Photometer System
MultiDirect

GB **Instruction manual**

Safety precautions

⚠ CAUTION ⚠

Reagents are formulated exclusively for chemical analysis and must not be used for any other purpose. Reagents must not get into the hands of children. Some of the reagents contain substances which are not entirely harmless environmentally. Be aware of the ingredients and take proper care when disposing of the test solution.

⚠ CAUTION ⚠

Please read this instruction manual before unpacking, setting up or using the photometer. Please read the method description completely before performing the test. Be aware of the risks of using the required reagents by reading the MSDS (Material Safety Data Sheets). Failure could result in serious injury to the operator or damage to the instrument.

MSDS: www.tintometer.de

⚠ CAUTION ⚠

Use the charger unit only with rechargeable batteries. Failure can result in serious injury to the operator or damage to the instrument.
Do not use charger with non rechargeables batteries.

⚠ CAUTION ⚠

The accuracy of the instrument is only valid if the instrument is used in an environment with controlled electromagnetic disturbances according to DIN 61326. Wireless devices, e.g. wireless phones, must not be used near the instrument.

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Part 1

Methods

1.1 Methods

1.1 Table of Methods

No.	Analysis	Reagent	Range	Displayed as	Method	λ [nm]	Page
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30	**Alkalinity, total T**	tablet	5-200	mg/l $CaCO_3$	Acid/Indicator [1,2,5]	610	14
35	**Alkalinity-p T**	tablet	5-300	mg/l $CaCO_3$	Acid/Indicator [1,2,5]	560	16
40	**Aluminium T**	tablet	0.01-0.3	mg/l Al	Eriochrome Cyanine R [2]	530	18
50	**Aluminium PP**	PP + liquid	0.01-0.25	mg/l Al	Eriochrome Cyanine R [2]	530	20
60	**Ammonia T**	tablet	0.02-1	mg/l N	Indophenol blue [2,3]	610	22
62	**Ammonia PP**	PP	0.01-0.8	mg/l N	Salicylate [2]	660	24
65	**Ammonia LR TT**	tube test	0.02-2.5	mg/l N	Salicylate [2]	660	26
66	**Ammonia HR TT**	tube test	1-50	mg/l N	Salicylate [2]	660	28
85	**Boron T**	tablet	0.1-2	mg/l B	Azomethine [3]	430	30
80	**Bromine T**	tablet	0.05-13	mg/l Br_2	DPD [5]	530	32
63	**Chloramine, mono PP**	PP + liquid	0.04-4.50	mg/l Cl_2	Indophenol	660	34
90	**Chloride T**	tablet	0.5 -25	mg/l Cl	Silver nitrate/ turbidity	530	38
100	**Chlorine T ***	tablet	0.01-6	mg/l Cl_2	DPD [1,2,3]	530	40, 42
101	**Chlorine L ***	liquid	0.02-4	mg/l Cl_2	DPD [1,2,3]	530	40, 46
110	**Chlorine PP ***	PP	0.02-2	mg/l Cl_2	DPD [1,2]	530	40, 50
120	**Chlorine dioxide T**	tablet	0.05-11	mg/l ClO_2	DPD, Glycine [1,2]	530	54
105	**Chlorine HR (KI) T**	tablet	5-200	mg/l Cl_2	KI/Acid [5]	530	60
130	**COD LR TT**	tube test	0 -150	mg/l O_2	Dichromate/H_2SO_4 [1,2]	430	62
131	**COD MR TT**	tube test	0 -1500	mg/l O_2	Dichromate/H_2SO_4 [1,2]	610	64
132	**COD HR TT**	tube test	0 -15	g/l O_2	Dichromate/H_2SO_4 [1,2]	610	66
204	**Colour**	direct reading	0-500	mg/l Pt	Pt-Co-Scale [1,2] (APHA)	430	68
150	**Copper T ***	tablet	0.05-5	mg/l Cu	Biquinoline [4]	560	70
153	**Copper PP**	PP	0.05-5	mg/l Cu	Bicinchoninate	560	74
157	**Cyanide**	Powder + liquid	0.01-0.5	mg/l CN	Pyridine-barbituric acid [1]	580	76
160	**Cyanuric acid T**	tablet	2-160	mg/l Cys	Melamine	530	78
165	**DEHA T**	tablet + liquid	20-500	µg/l DEHA	PPST [3]	560	80
167	**DEHA PP**	PP + liquid	20-500	µg/l DEHA	PPST [3]	560	82
170	**Fluoride L**	liquid	0.05-2	mg/l F	SPADNS [2]	580	84
190	**Hardness, Calcium T**	tablet	50-900	mg/l $CaCO_3$	Murexide [4]	560	86

* = free, combined, total; PP = powder pack; T = tablet;
L = liquid; TT = tube test; LR = low range; MR = middle range; HR = high range;
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1.1 Methods

1.1 Table of Methods

No.	Analysis	Reagent	Range	Displayed as	Method	λ [nm]	Page
191	**Hardness, Calcium 2T**	tablet	0-500	mg/l $CaCO_3$	Murexide [4]	560	88
200	**Hardness, total T**	tablet	2-50	mg/l $CaCO_3$	Metallphthalein [3]	560	90
201	**Hardness, total HR T**	tablet	20-500	mg/l $CaCO_3$	Metallphthalein [3]	560	92
205	**Hydrazine P**	powder	0.05-0.5	mg/l N_2H_4	4-(Dimethyl-amino)-benzaldehyde [3]	430	94
206	**Hydrazine L**	liquid	0.005-0.6	mg/l N_2H_4	4-(Dimethyl-amino)-benzaldehyde [3]	430	96
207	**Hydrazine C**	Vacu-vial	0.01-0.7	mg/l N_2H_4	PDMAB	430	98
210	**Hydrogen peroxide**	tablet	0.03-3	mg/l H_2O_2	DPD/catalyst [5]	530	100
215	**Iodine T**	tablet	0.05-3.6	mg/l I	DPD [5]	530	102
220	**Iron T**	tablet	0.02-1	mg/l Fe	PPST [3]	560	104,106
222	**Iron PP**	PP	0.02-3	mg/l Fe	1,10-Phenan-troline [3]	530	104,108
223	**Iron (TPTZ) PP**	PP	0.02-1.8	mg/l Fe	TPTZ	580	104,110
240	**Manganese T**	tablet	0.2-4	mg/l Mn	Formaldoxime	530	112
242	**Manganese LR PP**	PP + liquid	0.01-0.7	mg/l Mn	PAN	560	114
243	**Manganese HR PP**	PP + liquid	0,1-18	mg/l Mn	Periodate oxidation [2]	530	116
250	**Molybdate T**	tablet	1-50	mg/l MoO_4	Thioglycolate [4]	430	118
252	**Molybdate HR PP**	PP	0.5-66	mg/l MoO_4	Mercaptoacetic acid	430	120
265	**Nitrate TT**	tube test	1-30	mg/l N	Chromotropic acid	430	122
270	**Nitrite T**	tablet	0.01-0.5	mg/l N	N-(1-Naphthyl)-ethylendiamine [2,3]	560	124
272	**Nitrite LR PP**	PP	0.01-0.3	mg/l N	Diazotization	530	126
280	**Nitrogen, total LR TT**	tube test	0.5-25	mg/l N	Persulfate digestion method	430	128
281	**Nitrogen, total HR TT**	tube test	5-150	mg/l N	Persulfate digestion method	430	130
290	**Oxygen, active T**	tablet	0.1-10	mg/l O_2	DPD	530	132
292	**Oxygen, dissolved**	Vacu-vial	10-800	µg/l O_2	Rhodazine D™	530	134
300	**Ozone (DPD) T**	tablet	0.02-2	mg/l O_3	DPD/Glycine [5]	530	136
70	**PHMB T**	tablet	2-60	mg/l PHMB	Buffer/Indicator	560	142
320	**Phosphate, T ortho LR**	tablet	0.05-4	mg/l PO_4	Ammonium-molybdate [2,3]	660	144, 146

* = free, combined, total; PP = powder pack; T = tablet;
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1.1 Methods

1.1 Table of Methods

No.	Analysis	Reagent	Range	Displayed as	Method	λ [nm]	Page
321	**Phosphate, ortho HR T**	tablet	1-80	mg/l PO_4	Vanando-molybdate [2]	430	144, 148
323	**Phosphate, PP ortho**	PP	0.06-2.5	mg/l PO_4	Ascorbic acid [2]	660	144, 150
324	**Phosphate, ortho TT**	tube test	0.06-5	mg/l PO_4	Ascorbic acid [2]	660	144, 152
327	**Phosphate 1 C, ortho**	Vacu-vial	5-40	mg/l PO_4	Vanado-molybdate [2]	430	144, 154
328	**Phosphate 2 C, ortho**	Vacu-vial	0.05-5	mg/l PO_4	Stannous chloride [2]	660	144, 156
325	**Phosphate, hydr. TT**	tube test	0.02-1.6	mg/l P	Acid digestion, Ascorbic acid [2]	660	144, 158
326	**Phosphate, total TT**	tube test	0.02-1.1	mg/l P	Acid persulf digestion, Ascorbic acid [2]	660	144, 160
316	**Phosphonate PP**	PP	0-125	mg/l	Persulfate UV-Oxidation	660	162
329	**pH-Value LR T**	tablet	5.2-6.8	—	Bromocresolpurple [5]	560	166
330	**pH-Value T**	tablet	6.5-8.4	—	Phenolred [5]	560	168
331	**pH-Value L**	liquid	6.5-8.4	—	Phenolred [5]	560	170
332	**pH-Value HR T**	tablet	8.0-9.6	—	Thymolblue [5]	560	172
340	**Potassium T**	tablet	0.7-12	mg/l K	Tetraphenylborate-Turbidity [4]	430	174
350	**Silica T**	tablet	0.05-4	mg/l SiO_2	Silicomolybdate [2,3]	660	176
351	**Silica LR PP**	PP	0.1-1.6	mg/l SiO_2	Heteropolyblue [2]	660	178
352	**Silica HR PP**	PP	1-90	mg/l SiO_2	Silicomolybdate [2]	430	180
212	**Sodium hypochlorite T**	tablet	0.2-16	% NaOCl	Potassium iodide [5]	530	182
355	**Sulfate T**	tablet	5-100	mg/l SO_4	Bariumsulfate-Turbidity	610	184
360	**Sulfate PP**	PP	5-100	mg/l SO_4	Bariumsulfate-Turbidity [2]	530	186
365	**Sulfide**	tablet	0.04-0.5	mg/l S	DPD/Catalyst [3,4]	660	188
370	**Sulfite T**	tablet	0.1-5	mg/l SO_3	DTNB	430	190
384	**Suspended Solids**	direct reading	0-750	mg/l TSS	photometric	660	192
386	**Turbidity**	direct reading	0-1000	FAU	Attenuated Radiation Method	530	194
390	**Urea T**	tablet + liquid	0.1-2.5	mg/l Urea	Indophenol/ Urease	610	196
400	**Zinc T**	tablet	0.02 -1	mg/l Zn	Zincon [3]	610	198

* = free, combined, total; PP = powder pack; T = tablet;
L = liquid; TT = tube test; LR = low range; MR = middle range; HR = high range;
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1.1 Methods

The precision of Lovibond® Reagent Systems (tablets, powder packs and tube tests) is identical to the precision specified in standards literature such as American Standards (AWWA), ISO etc.

Most of the data referred to in these standard methods relates to Standard Solutions. Therefore they are not readily applicable to drinking-, boiler- or waste-water, since various interferences can have a major influence on the accuracy of the method.
For this reason we don't state such potentially misleading data.

Due to the fact that each sample is different, the only way to check the tolerances ('precision') is the Standard Additions Method.

According to this method, first the original sample is tested. Then further samples (2 to 4) are taken and small amounts of a Standard Solution are added, and further results are obtained. The amounts added range from approximately half, up to double the amount present in the sample itself.
These supplementary results make it possible to estimate the actual concentration of the original sample by comparison.

Literature

The reagent formulations are based on internationally recognised test methods. Some are described in national and/or international guidelines.

1) Deutsche Einheitsverfahren zur Wasser-, Abwasser- und Schlammuntersuchung
2) Standard Methods for the Examination of Water and Wastewater; 18th Edition, 1992
3) Photometrische Analysenverfahren, Schwedt,
   Wissenschaftliche Verlagsgesellschaft mbH, Stuttgart 1989
4) Photometrische Analyse, Lange / Vejdelek, Verlag Chemie 1980
5) Colorimetric Chemical Analytical Methods, 9th Edition, London

Notes for searching:

Active Oxygen	->	Oxygen, activ
Alkalinity-m	->	Alkalinity, total
Alkalinity, total	->	Alkalinity, total
Biguanide	->	PHMB
Calcium Hardness	->	Hardness, Calcium
Total Hardness	->	Hardness, total
Monochloramine	->	Chloramine, mono
m-Value	->	Alkalinity, total
p-Value	->	Alkalinity-p
Silicon dioxide	->	Silica
total Alkalinity	->	Alkalinity, total
total Hardness	->	Hardness, total
Langelier Saturation Index (Water Balance)	->	**Mode function 70**

1.1 Methods

2 0 Acid demand to pH 4.3 with Tablet

0.1 – 4 mmol/l

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber.

5. Add **one ALKA-M-PHOTOMETER tablet** straight from the foil to the water sample and crush the tablet using a clean stirring rod.

6. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.

7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.

The result is shown in the display as Acid demand to pH 4.3 in mmol/l.

1.1 Methods

Notes:

1. The terms total Alkalinity, Alkalinity-m, m-Value and Acid demand to pH 4.3 are identical.
2. For accurate results exactly 10 ml of water sample must be taken for the test.

1.1 Methods

3 0 Alkalinity, total = Alkalinity-m = m-Value with Tablet

5 – 200 mg/l $CaCO_3$

Ø 24 mm

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.
4. Remove the vial from the sample chamber.
5. Add **one ALKA-M-PHOTOMETER tablet** straight from the foil to the water sample and crush the tablet using a clean stirring rod.
6. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.
7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.

The result is shown in the display as total Alkalinity.

1.1 Methods

Notes:

1. The terms total Alkalinity, Alkalinity-m, m-Value and Alkalinity to pH 4.3 are identical.
2. For accurate results exactly 10 ml of water sample must be taken for the test.
3. Conversion table:

	Acid demand to pH 4.3 DIN 38 409 ($K_{S4.3}$)	German °dH*	English °eH*	French °fH*
1 mg/l $CaCO_3$	0.02	0.056	0.07	0.1

*Carbonate hardness (reference = Hydrogencarbonate-anions)

Example:
10 mg/l $CaCO_3$ = 10 mg/l x 0.056 = 0.56 °dH
10 mg/l $CaCO_3$ = 10 mg/l x 0.02 = 0.2 mmol/l

4. ▲ $CaCO_3$
   °dH
   °eH
   °fH
   ▼ °aH

1.1 Methods

3 5

Alkalinity-p = p-value with Tablet

5 – 300 mg/l $CaCO_3$

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber.

5. Add **one ALKA-P-PHOTOMETER tablet** straight from the foil to the water sample and crush the tablet using a clean stirring rod.

6. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.

7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.

The result is shown in the display as Alkalinity-p.

1.1 Methods

Notes

1. The terms Alkalinity-p, p-Value and Alkalinity to pH 8.2 are identical.
2. For accurate test results exactly 10 ml of water sample must be taken for the test.
3. This method was developed from a volumetric procedure for the determination of Alkalinity-p. Due to undefined conditions, the deviations from the standardised method may be greater.
4. Conversion table:

	mg/l $CaCO_3$	°dH	°fH	°eH
1 mg/l $CaCO_3$	----	0.056	0.10	0.07
1 °dH	17.8	----	1.78	1.25
1 °fH	10.0	0.56	----	0.70
1 °eH	14.3	0.80	1.43	----

▲ $CaCO_3$
°dH
°eH
°fH
▼ °aH

5. By determining Alkalinity-p and Alkalinity-m it is possible to classify the alkalinity as Hydroxide, Carbonate and Hydrogencarbonate.

   The following differentiation is only valid if:

   a) no other alkalis are present and

   b) Hydroxide und Hydrogen are not present in the same water sample.

   If condition b) is not fulfilled please get additional information from "Deutsche Einheitsverfahren zur Wasser-, Abwasser- und Schlammuntersuchung, D 8".

   Case 1: Alkalinity-p = 0
   Hydrogen carbonate = m
   Carbonate = 0
   Hydroxide = 0

   Case 2: Alkalinity-p > 0 and Alkalinity-m > 2p
   Hydrogen carbonate = m – 2p
   Carbonate = 2p
   Hydroxide = 0

   Case 3: Alkalinity-p > 0 and Alkalinity-m < 2p
   Hydrogen carbonate = 0
   Carbonate = 2m – 2p
   Hydroxide = 2p – m

1.1 Methods

4 0 Aluminium with Tablet

0.01 – 0.3 mg/l Al

Ø 24 mm

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber.

5. Add **one ALUMINIUM No. 1 tablet** straight from the foil to the water sample and crush the tablet using a clean stirring rod (dissolve the tablet).

6. Add **one ALUMINIUM No. 2 tablet** straight from the foil to the same water sample and crush the tablet using a clean stirring rod.

7. Close the vial tightly with the cap and swirl gently several times until the tablets are dissolved.

8. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

9. Press **TEST** key.
Wait for a **reaction period of 5 minutes.**

Countdown
5:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l Aluminium.

1.1 Methods

Notes:

1. Before use, clean the vials and the measuring beaker with Hydrochloric acid (approx. 20%). Rinse them thoroughly with deionised water.
2. To get accurate results the sample temperature must be between 20°C and 25°C.
3. A low test result may be given in the presence of Fluorides and Polyphosphates. The effect of this is generally insignificant unless the water has fluoride added artificially. In this case, the following table should be used:

Fluoride	Displayed value: Aluminium [mg/l Al]					
[mg/l F]	0.05	0.10	0.15	0.20	0.25	0.30
0.2	0.05	0.11	0.16	0.21	0.27	0.32
0.4	0.06	0.11	0.17	0.23	0.28	0.34
0.6	0.06	0.12	0.18	0.24	0.30	0.37
0.8	0.06	0.13	0.20	0.26	0.32	0.40
1.0	0.07	0.13	0.21	0.28	0.36	0.45
1.5	0.09	0.20	0.29	0.37	0.48	---

   Example: If the result of Aluminium determination is 0.15 mg/l Al and the Fluoride concentration is known to be 0.4 mg/l F, the true concentration of Aluminium is 0.17 mg/l Al.

4. ▲ Al
   ▼ Al_2O_3

1.1 Methods

5 0 Aluminium with Vario Powder Pack

0.01 – 0.25 mg/l Al

Ø 24 mm

Use two clean vials (24 mm Ø) and mark one as blank for zeroing.

1. Fill **20 ml of water sample** in a 100 ml beaker.

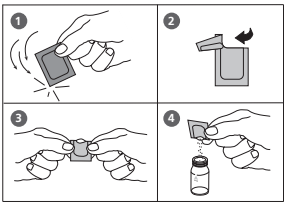
2. Add the contents of **one Vario Aluminum ECR F20 Powder Pack** straight from the foil to the water sample.

3. Dissolve the powder using a clean stirring rod.

Countdown 1
0:30
start: ↵

4. Press **[↵]** key.
   Wait for a **reaction period of 30 seconds.**

After the reaction period is finished proceed as follows:



5. Add the contents of **one Vario Hexamine F20 Powder Pack** straight from the foil to the same water sample.

6. Dissolve the powder using a clean stirring rod.

7. Add **1 drop of Vario Aluminum ECR Masking Reagent** in the vial marked as blank.

8. Add 10 ml of the prepared water sample to the vial (this is the blank).

9. Add the remaining 10 ml of the prepared water sample in the second clean vial (this is the sample).

10. Close the vials tightly with the caps and swirl several times to mix the contents.

Countdown 2
5:00
start: ↵

11. Press **[↵]** key.

    Wait for **a reaction period of 5 minutes**.

1.1 Methods

After the reaction period is finished proceed as follows:

12. Place the vial (the blank) in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

13. Press **ZERO** key.

14. Remove the vial from the sample chamber.

15. Place the vial (the sample) in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

16. Press **TEST** key.

The result is shown in the display in mg/l Aluminium.

Notes:

1. Before use, clean the vials and the measuring beaker with Hydrochloric acid (approx. 20%). Rinse them thoroughly with deionised water.
2. To get accurate results the sample temperature must be between 20°C and 25°C.
3. A low test result may be given in the presence of Fluorides and Polyphosphates. The effect of this is generally insignificant unless the water has fluoride added artificially. In this case, the following table should be used:

Fluoride	Displayed value: Aluminium [mg/l Al]					
[mg/l F]	0.05	0.10	0.15	0.20	0.25	0.30
0.2	0.05	0.11	0.16	0.21	0.27	0.32
0.4	0.06	0.11	0.17	0.23	0.28	0.34
0.6	0.06	0.12	0.18	0.24	0.30	0.37
0.8	0.06	0.13	0.20	0.26	0.32	0.40
1.0	0.07	0.13	0.21	0.28	0.36	0.45
1.5	0.09	0.20	0.29	0.37	0.48	---

Example: If the result of Aluminium determination is 0.15 mg/l Al and the Fluoride concentration is known to be 0.4 mg/l F, the true concentration of Aluminium is 0.17 mg/l Al.

4. ▲ Al
   ▼ Al_2O_3

1.1 Methods

60 Ammonia with Tablet

0.02 – 1 mg/l N

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample,** close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.
4. Remove the vial from the sample chamber.
5. Add **one AMMONIA No. 1 tablet** straight from the foil to the water sample and crush the tablet using a clean stirring rod.
6. Add **one AMMONIA No. 2 tablet** straight from the foil to the same water sample and crush the tablet using a clean stirring rod.
7. Close the vial tightly with the cap and swirl several times until the tablets are dissolved.
8. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

9. Press **TEST** key.
   Wait for a **reaction period of 10 minutes.**

Countdown
10:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l Ammonia as N.

1.1 Methods

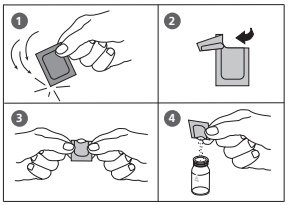
Notes:

1. The tablets must be added in the correct sequence.
2. The AMMONIA No. 1 tablet will only dissolve completely after the AMMONIA No. 2 tablet has been added.
3. The temperature of the sample is important for full colour development. At a temperature below 20°C the reaction period is 15 minutes.
4. Sea water samples:
   Ammonia conditioning reagent is required when testing sea water or brackish water samples to prevent precipitation of salts.
   Fill the test tube with the sample to the 10 ml mark and add one level spoonful of Conditioning Powder. Mix to dissolve, then continue as described in the test instructions.
5. Conversion:
   mg/l NH_4 = mg/l N x 1.29
   mg/l NH_3 = mg/l N x 1.22
6. ▲ N
   NH_4
   ▼ NH_3

1.1 Methods

6 2 Ammonia with Vario Powder Pack

0.01 – 0.8 mg/l N



Use two clean vials (24 mm Ø) and mark one as blank for zeroing.

1. Fill a clean vial (24 mm Ø) with **10 ml of deionised water** (this is the blank).
2. Fill the other clean vial (24 mm Ø) with **10 ml of water sample** (this is the sample).
3. Add the contents of **one Vario Ammonia Salicylate F10 Powder Pack** straight from the foil to each vial.
4. Close the vials tightly with the caps and swirl several times to mix the contents.

Countdown 1
3:00
start: ↵

5. Press **[↵]** key.
   Wait for a **reaction period of 3 minutes**.

   After the reaction period is finished proceed as follows:
6. Add the contents of **one Vario Ammonia Cyanurate F10 Powder Pack** straight from the foil to each sample.
7. Close the vials tightly with the caps and swirl several times to mix the contents.

Countdown 2
15:00
start: ↵

8. Press **[↵]** key.
   Wait for a **reaction period of 15 minutes**.

   After the reaction period is finished proceed as follows:
9. Place the vial (the blank) in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

10. Press **ZERO** key.
11. Remove the vial from the sample chamber.
12. Place the vial (the sample) in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

13. Press **TEST** key.

    The result is shown in the display in mg/l Ammonia as N.

1.1 Methods

Notes:

1. Extremely basic or acidic water samples should be adjusted with 0.5 mol/l (1 N) Sulfuric acid solution or 1 mol/l (1 N) Sodium hydroxide solution to pH 7.
2. Interferences:

Interfering substance	**Interference levels and treatments**
Calcium	greater than 1000 mg/l $CaCO_3$
Iron	Interferes at all levels. Correct as follows: a) determine the concentration of iron present in the sample by performing a total Iron test b) add the same iron concentration as determined to the deionised water (step 1). The interference will be blanked out successfully.
Magnesium	greater than 6000 mg/l $CaCO_3$
Nitrate	greater than 100 mg/l NO_3-N
Nitrite	greater than 12 mg/l NO_2-N
Phosphate	greater than 100 mg/l PO_4-P
Sulfate	greater than 300 mg/l SO_4
Sulfide	intensifies the colour
Glycine, Hydrazine, Colour, Turbidity	Less common interferences such as Hydrazine and Glycine will cause intensified colours in the prepared sample. Turbidity and colour will give erroneous high values. Samples with severe interferences require distillation.

3. ▲ N
   NH_4
   ▼ NH_3

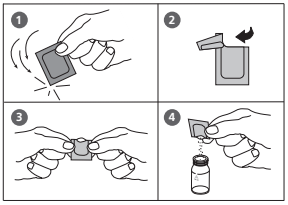
1.1 Methods

6 5 Ammonia LR with Vario Tube Test

0.02 – 2.5 mg/l N

Ø 16 mm

Insert the adapter for 16 mm Ø vials.



1. Open one white capped reaction vial and add **2 ml deionised water** (this is the blank).
2. Open another white capped reaction vial and add **2 ml of water sample** (this is the sample).
3. Add the contents of **one Vario Ammonia Salicylate F5 Powder Pack** straight from the foil into each vial.
4. Add the contents of **one Vario Ammonia Cyanurate F5 Powder Pack** straight from the foil into each vial.
5. Close the vials tightly with the caps and swirl several times to dissolve the powder.

Countdown 1
20:00
start: ↵

6. Press **[↵]** key.
   Wait for a **reaction period of 20 minutes**.

   After the reaction period is finished proceed as follows:

7. Place the vial (the blank) in the sample chamber making sure that the marks are aligned. Place the cover on the adapter.

prepare Zero
press ZERO

8. Press **ZERO** key.
9. Remove the vial from the sample chamber.
10. Place the vial (the sample) in the sample chamber making sure that the marks are aligned. Place the cover on the adapter.

Zero accepted
prepare Test
press TEST

11. Press **TEST** key.

    The result is shown in the display in mg/l Ammonia as N.

1.1 Methods

Notes:

1. Strong alkaline or acidic water samples must be adjusted to approx. pH 7 before analysis (use 1 mol/l Hydrochloric acid resp. 1 mol/l Sodium hydroxide).
2. If chlorine is known to be present, add one drop of 0.1 mol/l Sodium thiosulfate for each 0.3 mg/l Cl_2 in a one litre water sample.
3. Iron interferes with the test. The interferences will be eliminated as follows: Determine the amount of total iron present in the water sample. To produce the blank add an iron standard solution with the same iron concentration to the vial (point 1) instead of deionised water
4. Conversion:
   mg/l NH_4 = mg/l N x 1.29
   mg/l NH_3 = mg/l N x 1.22
5. ▲ N
   NH_4
   ▼ NH_3

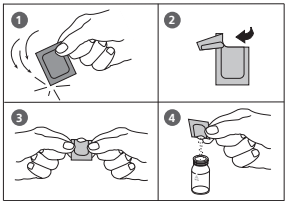
1.1 Methods

6 6 Ammonia HR with Vario Tube Test

1 – 50 mg/l N

Ø 16 mm

Insert the adapter for 16 mm Ø vials.



1. Open one white capped reaction vial and add **0.1 ml deionised water** (this is the blank).
2. Open another white capped reaction vial and add **0.1 ml of water sample** (this is the sample).
3. Add the contents of **one Vario Ammonia Salicylate F5 Powder Pack** straight from the foil into each vial.
4. Add the contents of **one Vario Ammonia Cyanurate F5 Powder Pack** straight from the foil into each vial.
5. Close the vials tightly with the caps and swirl several times to dissolve the powder.

Countdown 1
20:00
start: ↵

6. Press **[↵]** key.
   Wait for a **reaction period of 20 minutes**.

   After the reaction period is finished proceed as follows:

7. Place the vial (the blank) in the sample chamber making sure that the marks are aligned. Place the cover on the adapter.

prepare Zero
press ZERO

8. Press **ZERO** key.
9. Remove the vial from the sample chamber.
10. Place the vial (the sample) in the sample chamber making sure that the marks are aligned. Place the cover on the adapter.

Zero accepted
prepare Test
press TEST

11. Press **TEST** key.

    The result is shown in the display in mg/l Ammonia as N.

1.1 Methods

Notes:

1. Strong alkaline or acidic water samples must be adjusted to approx. pH 7 before analysis (use 1 mol/l Hydrochloric acid resp. 1 mol/l Sodium hydroxide).
2. If chlorine is known to be present, add one drop of 0.1 mol/l Sodium thiosulfate for each 0.3 mg/l Cl_2 in a one litre water sample.
3. Iron interferes with the test. The interferences will be eliminated as follows: Determine the amount of total iron present in the water sample. Add an iron standard solution with the same concentration to the vial (point 1) instead of deionised water to produce the blank.
4. Conversion:
   mg/l NH_4 = mg/l N x 1.29
   mg/l NH_3 = mg/l N x 1.22
5. ▲ N
   NH_4
   ▼ NH_3

1.1 Methods

8 5

Boron with Tablet

0.1 – 2 mg/l B

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.
4. Remove the vial from the sample chamber.
5. Add **one BORON No. 1 tablet** straight from the foil to the water sample and crush the tablet using a clean stirring rod and dissolve the tablet.
6. Add **one BORON No. 2 tablet** straight from the foil to the same water sample and crush the tablet using a clean stirring rod.
7. Close the vial tightly with the cap and swirl several times until the tablets are dissolved.
8. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

9. Press **TEST** key.

Countdown
20:00

Wait for a **reaction period of 20 minutes.**

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l Boron.

1.1 Methods

Notes:

1. The tablets must added in the correct sequence.
2. The sample solution should have a pH value between 6 and 7.
3. Interferences are prevented by the presence of EDTA in the tablets.
4. The rate of colour development depends on the temperature. The temperature of the sample must be 20 °C ± 1 °C.
5. ▲ B
   ▼ H_3BO_3

1.1 Methods

8 0

Bromine with Tablet

0.05 – 13 mg/l Br_2

Ø 24 mm

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber and **empty it, leaving a few drops remaining in the vial**.

5. Add **one DPD No. 1 tablet** straight from the foil and crush the tablet using a clean stirring rod.

6. Add water sample to the 10 ml mark.

7. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.

8. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

9. Press **TEST** key.

The result is shown in the display in mg/l Bromine.

1.1 Methods

Notes:

1. Vial cleaning:
   As many household cleaners (e.g. dishwasher detergent) contain reducing substances, the subsequent determination of Bromine may show lower results. To avoid any measurement errors, only use glassware free of Chlorine demand.
   Preparation: Put all applicable glassware into Sodium hypochlorite solution (0.1 g/l) for one hour, then rinse all glassware thoroughly with deionised water.
2. Preparing the sample:
   When preparing the sample, the lost of Bromine, e.g. by pipetting or shaking, must be avoided. The analysis must take place immediately after taking the sample.
3. The DPD colour development is carried out at a pH value of 6.2 to 6.5. The reagent tablet therefore contains a buffer for the pH adjustment. Strong alkaline or acidic water samples must be adjusted between pH 6 and pH 7 before the reagent is added (use 0.5 mol/l Sulfuric acid resp. 1 mol/l Sodium hydroxide).
4. Exceeding the measuring range:
   Concentrations above 22 mg/l Bromine can lead to results showing 0 mg/l.
   In this case, the water sample must be diluted with water free of Bromine.
   10 ml of the diluted sample should be mixed with the reagent and the measurement repeated.

   Oxidising agents such as Chlorine, Ozone etc. interfere as they react in the same way as Bromine.

1.1 Methods

6 3 Chloramine (Mono) and Free Ammonia with Vario Powder Pack and Liquid Reagent

0.04 – 4.50 mg/l Cl_2

Chloramine (Mono)
>> with NH4
without NH4

The following selection is shown in the display:

>> with NH4

for the determination of Monochloramine and free Ammonia

>> without NH4

for the determination of Monochloramine

Select the desired determination with the arrow keys [▲] and [▼]. Confirm with [↵] key.

Notes:

1. Full colour development – temperature
   The reaction periods indicated in the manual refer to a sample temperature between 18° and 20°C. Due to the fact that the reaction period is strongly influenced by sample temperature, you have to adjust both reaction periods according to the following table:

Sample temperature in °C	Reaction period in min
5	10
10	8
16	6
20	5
23	2.5
25	2

1.1 Methods

6 3 Chloramine (Mono) with Vario Powder Pack

0.04 – 4.50 mg/l Cl_2

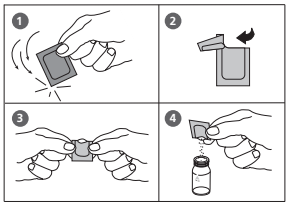
1. Fill a clean vial (24 mm Ø) with **10 ml of the water sample**, close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber.



5. Add the contents of **one Vario Monochlor FRGT Powder Pack** straight from the foil into the water sample.

6. Close the vial tightly with the cap and swirl several times to mix the contents (approx. 20 seconds).

7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.
   Wait for a **reaction period of 5 minutes**.

Countdown
5:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l Monochloramine.

▲ mg/l Cl_2
mg/l NH_2Cl
▼ mg/l N

Notes:
see previous page

6 3 Chloramine (Mono) and Free Ammonia with Vario Powder Pack and Liquid Reagent

0.04 – 4.50 mg/l Cl_2
0.01 – 0.50 mg/l NH_4-N

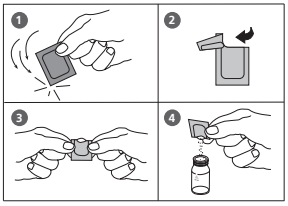
Ø 24 mm

Use two clean vials (24 mm Ø) and mark one as the chloramine vial, the other as the ammonia vial.

1. Fill both vials (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.
2. Place the chloramine vial in the sample chamber making sure that the ⨯ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.
4. Remove the vial from the sample chamber.



5. Add the contents of **one Vario Monochlor F RGT Powder Pack** straight from the foil into the chloramine vial.
6. Close the vial tightly with the cap and swirl several times to mix the contents (approx. 20 seconds).
7. Add **1 drop of Vario Free Ammonia Reagent Solution** into the ammonia vial (Note 1).
8. Close the vial tightly with the cap and invert several times to mix the contents.
9. Place the chloramine vial in the sample chamber making sure that the ⨯ marks are aligned.

Zero accepted
prepare T1
press TEST

10. Press **TEST** key.
    Wait for a **reaction period of 5 minutes**.

Countdown
5:00

    After the reaction period is finished the measurement starts automatically.

11. Remove the vial from the sample chamber.
12. Add the contents of **one Vario Monochlor F RGT Powder Pack** straight from the foil into the ammonia vial.

1.1 Methods

13. Close the vial tightly with the cap and swirl several times to mix the contents (approx. 20 seconds).

14. Place the ammonia vial in the sample chamber making sure that the ⧖ marks are aligned.

T1 accepted
prepare T2
press TEST

15. Press **TEST** key.
    Wait for a **reaction period of 5 minutes**.

Countdown
5:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l Monochloramine and mg/l free Ammonia.

***,** mg/l Cl2**

***,** mg/l NH2Cl**

***,** mg/l N [NH2Cl]**

The reading of Monochloramine is shown as:

▲ mg/l Cl_2
mg/l NH_2Cl
▼ mg/l N

The reading of Ammonia is shown as N.

Conversion:
mg/l NH_4 = mg/l N · 1.29
mg/l NH_3 = mg/l N · 1.22

***,** mg/l N [NH4]**

Notes:

1. Hold the bottle vertically and squeeze slowly.
2. To determine the ammonia concentration the difference between the chloramine (T1) and the sum of chloramine and ammonia (T2) is calculated. If T2 exceeds the range limit the following message is displayed:
   $NH_2Cl + NH_4 > 0.5$ mg/l
   In this case the sample has to be diluted and the measurement repeated.
3. see also page 34

1.1 Methods

9 0 Chloride with Tablet

0.5 – 25 mg/l Cl

Ø 24 mm

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.
4. Remove the vial from the sample chamber.
5. Add **one CHLORIDE T1 tablet** straight from the foil to the water sample, crush the tablet using a clean stirring rod and dissolve the tablet.
6. Add **one CHLORIDE T2 tablet** straight from the foil to the same water sample and crush the tablet using a clean stirring rod.
7. Close the vial tightly with the cap and swirl gently several times until the tablet is dissolved (Note 1).
8. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

9. Press **TEST** key.

   Wait for a **reaction period of 2 minutes**.

Countdown
2:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l Chloride.

1.1 Methods

Notes:

1. Ensure that all particles of the tablet are dissolved – Chloride causes an extremely fine distributed turbidity with a milky appearance.
   Heavy shaking leads to bigger sized particles which can cause false readings.
2. High concentrations of electrolytes and organic compounds have different effects on the precipitation reaction.
3. Ions which also form deposits with Silver nitrate in acidic media, such as Bromides, Iodides and Thiocyanates, interfere with the analysis.
4. Highly alkaline water should – if necessary – be neutralised using Nitric acid before analysis.

1.1 Methods

1 0 0 Chlorine with Tablet

0.01 – 6 mg/l Cl_2

1 0 1 Chlorine with Liquid Reagent

0.02 - 4 mg/l Cl_2

1 1 0 Chlorine with Vario Powder Pack

0.02 - 2 mg/l Cl_2

Chlorine
>> diff
free
total

The following selection is shown in the display:

>> diff

for the differentiated determination of free, combined and total Chlorine.

>> free

for the determination of free Chlorine.

>> total

for the determination of total Chlorine.

Select the desired determination with the arrow keys [▲] and [▼]. Confirm with [↵] key.

1.1 Methods

Notes:

1. Vial cleaning:
   As many household cleaners (e.g. dishwasher detergent) contain reducing substances, the subsequent determination of Chlorine may show lower results. To avoid any measurement errors, only use glassware free of Chlorine demand.
   Preparation: Put all applicable glassware into Sodium hypochlorite solution (0.1 g/l) for one hour, then rinse all glassware thoroughly with deionised water.
2. For individual testing of free and total Chlorine, the use of different sets of glassware is recommended (EN ISO 7393-2, 5.3)
3. Preparing the sample:
   When preparing the sample, the lost of Chlorine, e.g. by pipetting or shaking, must be avoided. The analysis must take place immediately after taking the sample.
4. The DPD colour development is carried out at a pH value of 6.2 to 6.5. The reagents therefore contain a buffer for the pH adjustment.
   Strong alkaline or acidic water samples must be adjusted between pH 6 and pH 7 before the reagent is added (use 0.5 mol/l Sulfuric acid resp. 1 mol/l Sodium hydroxide).
5. Exceeding the measuring range:
   Concentrations above:
   10 mg/l Chlorine using tablets
   4 mg/l Chlorine using liquid reagents
   2 mg/l using powder packs
   can lead to results showing 0 mg/l. In this case, the water sample must be diluted with water free of Chlorine. 10 ml of the diluted sample should be mixed with the reagent and the measurement repeated.
6. Turbidity (can lead to errors):
   The use of the DPD No. 1 tablet (method 100) in samples with high Calcium ion contents* and/or high conductivity* can lead to turbidity of the sample and therefore incorrect measurements. In this case, the reagent tablet DPD No. 1 High Calcium should be used as an alternative. Even if turbidity does occur after the DPD No. 3 tablet has been added, this can be prevented by using the DPD No. 1 HIGH CALCIUM tablet.
   ** it is not possible to give exact values, because the development of turbidity depends on the nature of the sample.*
7. If `???` is displayed at a differentiated test result see page 262.

Oxidising agents such as Bromine, Ozone etc. interfere as they react in the same way as Chlorine.

1.1 Methods

1 0 0 Chlorine, differentiated determination with Tablet

0.01 – 6 mg/l Cl_2

Ø 24 mm

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample,** close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber and **empty it, leaving a few drops remaining in the vial**.

5. Add **one DPD No. 1 tablet** straight from the foil and crush the tablet using a clean stirring rod.

6. Add water sample to the 10 ml mark.

7. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.

8. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare T1
press TEST

9. Press **TEST** key.

10. Remove the vial from the sample chamber.

11. Add **one DPD No. 3 tablet** straight from the foil to the same water sample and crush the tablet using a clean stirring rod.

12. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.

13. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

T1 accepted
prepare T2
press TEST

14. Press **TEST** key.
Wait for a **reaction period of 2 minutes**.

Countdown
2:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in:

***,** mg/l free Cl**
***,** mg/l comb Cl**
***,** mg/l total Cl**

mg/l free Chlorine
mg/l combined Chlorine
mg/l total Chlorine

Notes:

See page 41.

1.1 Methods

1 0 0 Chlorine, free with Tablet

0.01 – 6 mg/l Cl_2

Ø 24 mm

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.
4. Remove the vial from the sample chamber and **empty it, leaving a few drops remaining in the vial**.
5. Add **one DPD No. 1 tablet** straight from the foil and crush the tablet using a clean stirring rod.
6. Add water sample to the 10 ml mark.
7. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.
8. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

9. Press **TEST** key.

The result is shown in the display in mg/l free Chlorine.

Notes:

See page 41.

1.1 Methods

1 0 0

Chlorine, total with Tablet

0.01 – 6 mg/l Cl_2

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber and **empty it, leaving a few drops remaining in the vial**.

5. Add **one DPD No. 1 tablet** and **one DPD No. 3 tablet** straight from the foil and crush the tablets using a clean stirring rod.

6. Add water sample to the 10 ml mark.

7. Close the vial tightly with the cap and swirl several times until the tablets are dissolved.

8. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

9. Press **TEST** key.
Wait for a **reaction period of 2 minutes**.

Countdown
2:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l total Chlorine.

Notes:

See page 41.

1.1 Methods

1 0 1 Chlorine, differentiated determination with Liquid Reagent

0.02 – 4 mg/l Cl_2

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.
4. Remove the vial from the sample chamber and **empty the vial**.
5. Fill the vial with drops of the same size by holding the bottle vertically and squeeze slowly:

   6 drops of DPD 1 buffer solution

   2 drops of DPD 1 reagent solution
6. Add water sample to the 10 ml mark.
7. Close the vial tightly with the cap and swirl several times to mix the contents.
8. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare T1
press TEST

9. Press **TEST** key.
10. Remove the vial from the sample chamber.
11. **Add 3 drops of DPD 3 solution** to the same water sample.
12. Close the vial tightly with the cap and swirl several times to mix the contents.

1.1 Methods

13. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

T1 accepted **prepare T2** **press TEST**

14. Press **TEST** key.
Wait for a **reaction period of 2 minutes**.

Countdown **2:00**

After the reaction period is finished the reading starts automatically.

The result is shown in the display in:

***,** mg/l free Cl**	mg/l free Chlorine
***,** mg/l comb. Cl**	mg/l combined Chlorine
***,** mg/l total Cl**	mg/l total Chlorine

Notes:

1. After use replace the bottle caps securely noting the colour coding.
2. **Store the reagent bottles in a cool, dry place ideally between 6°C and 10°C**.
3. Also see page 41.

1.1 Methods

(1) (0) (1) Chlorine, free with Liquid Reagent

0.02 – 4 mg/l Cl_2

prepare Zero
press ZERO

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.
3. Press **ZERO** key.
4. Remove the vial from the sample chamber and **empty the vial**.
5. Fill the vial with drops of the same size by holding the bottle vertically and squeeze slowly:

   6 drops of DPD 1 buffer solution

   2 drops of DPD 1 reagent solution
6. Add water sample to the 10 ml mark.
7. Close the vial tightly with the cap and swirl several times to mix the contents.
8. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

9. Press **TEST** key.

   The result is shown in the display in mg/l free Chlorine.

Notes (free and total Chlorine):

1. After use replace the bottle caps securely noting the colour coding.
2. **Store the reagent bottles in a cool, dry place ideally between 6 °C and 10 °C.**
3. Also see page 41.

1.1 Methods

1 0 1 Chlorine, total with Liquid Reagent

0.02 – 4 mg/l Cl_2

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber and **empty the vial**.

5. Fill the vial with drops of the same size by holding the bottle vertically and squeeze slowly:

   6 drops of DPD 1 buffer solution

   2 drops of DPD 1 reagent solution

   3 drops of DPD 3 solution

6. Add water sample to the 10 ml mark.

7. Close the vial tightly with the cap and swirl several times to mix the contents.

8. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

9. Press **TEST** key.
   Wait for a **reaction period of 2 minutes**.

Countdown
2:00

After the reaction period is finished the reading starts automatically.

The result is shown in the display in mg/l total Chlorine.

1.1 Methods

1 1 0 Chlorine, differentiated determination with Vario Powder Pack

0.02 – 2 mg/l Cl_2

Ø 24 mm

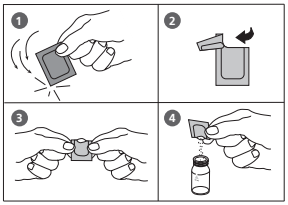
1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber.



5. Add the contents of **one Vario Chlorine FREE-DPD/ F10 Powder Pack** straight from the foil to the water sample.

6. Close the vial tightly with the cap and swirl several times to mix the contents (approx. 20 seconds).

7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare T1
press TEST

8. Press **TEST** key.

9. Remove the vial from the sample chamber, empty the vial, rinse vial and cap several times and then fill the vial with **10 ml of water sample**.

10. Add the contents of **one Vario Chlorine TOTAL-DPD / F10 Powder Pack** straight from the foil to the water sample.

11. Close the vial tightly with the cap and swirl several times to mix the contents (approx. 20 seconds).

12. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

T1 accepted
prepare T2
press TEST

13. Press **TEST** key.
Wait for a **reaction period of 3 minutes.**

Countdown
3:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in:

***,** mg/l free Cl**
***,** mg/l comb. Cl**
***,** mg/l total Cl**

mg/l free Chlorine
mg/l combined Chlorine
mg/l total Chlorine

Notes:

See page 41.

1.1 Methods

1 1 0 Chlorine, free with Vario Powder Pack

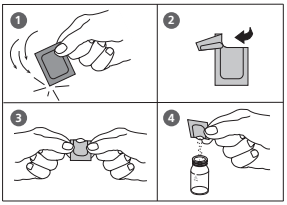
0.02 – 2 mg/l Cl_2

Ø 24 mm

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.
4. Remove the vial from the sample chamber.



5. Add the contents of **one Vario Chlorine FREE-DPD / F10 Powder Pack** straight from the foil to the water sample.
6. Close the vial tightly with the cap and swirl several times to mix the contents (approx. 20 seconds).
7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.

The result is shown in the display in mg/l free Chlorine.

Notes:

See page 41.

1.1 Methods

1 1 0 Chlorine, total with Vario Powder Pack

0.02 – 2 mg/l Cl_2

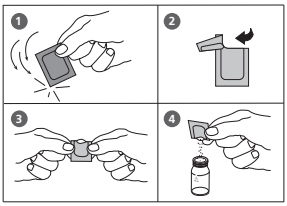
1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber.



5. Add the contents of **one Vario Chlorine TOTAL-DPD / F10 Powder Pack** straight from the foil to the water sample.

6. Close the vial tightly with the cap and swirl several times to mix the contents (approx. 20 seconds).

7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.
Wait for a **reaction period of 3 minutes**.

Countdown
3:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l total Chlorine.

Notes:

See page 41.

1.1 Methods

120 Chlorine dioxide with Tablet

0.05 – 11 mg/l ClO_2

Chlorine dioxide **>> with Cl** **without Cl**

The following selection is shown in the display:

>> with Cl

for the determination of Chlorine dioxide in the presence of Chlorine.

>> without Cl

for the determination of Chlorine dioxide in the absence of Chlorine.

Select the desired determination with the arrow keys [▲] and [▼]. Confirm with [↵] key.

1.1 Methods

Notes:

1. Vial cleaning:

   As many household cleaners (e.g. dishwasher detergent) contain reducing substances, the subsequent determination of Chlorine dioxide may show lower results. To avoid any measurement errors, only use glassware free of Chlorine demand.

   Preparation: Put all applicable glassware into Sodium hypochlorite solution (0.1 g/l) for one hour, then rinse all glassware thoroughly with deionised water.

2. Preparing the sample:

   When preparing the sample, the lost of Chlorine dioxide, e.g. by pipetting or shaking, must be avoided. The analysis must take place immediately after taking the sample.

3. The DPD colour development is carried out at a pH value of 6.2 to 6.5. The reagent tablet therefore contains a buffer for the pH adjustment.

   Strong alkaline or acidic water samples must be adjusted between pH 6 and pH 7 before the tablet is added (use 0.5 mol/l Sulfuric acid resp. 1 mol/l Sodium hydroxide).

4. Exceeding the measuring range:

   Concentrations above 19 mg/l Chlorine dioxide can lead to results showing 0 mg/l. In this case, the water sample must be diluted with water free of Chlorine dioxide. 10 ml of the diluted sample should be mixed with the reagent and the measurement repeated.

5. If `???` is displayed at a differentiated test result see page 262.

Oxidising agents such as Chlorine, Ozone etc. interfere as they react in the same way as Chlorine dioxide.

1.1 Methods

120 Chlorine dioxide in the presence of Chlorine with Tablet

0.05 – 11 mg/l ClO_2

Ø 24 mm

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.
4. Remove the vial from the sample chamber and **empty it, leaving a few drops remaining in the vial**.
5. Add **one DPD No. 1 tablet** straight from the foil and crush the tablet using a clean stirring rod.
6. **Fill a second clean vial with 10 ml of water sample.**
7. Add **one GLYCINE tablet** straight from the foil and crush the tablet using a clean stirring rod.
8. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.
9. **Transfer the contents of the second vial into the prepared vial.**
10. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.
11. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare T1
press TEST

12. Press **TEST** key.

1.1 Methods

13. Remove the vial from the sample chamber, empty the vial, rinse vial and cap several times. Fill with **a few drops of the water sample**.

14. Add **one DPD No. 1 tablet** straight from the foil and crush the tablet using a clean stirring rod.

15. Add water sample to the 10 ml mark.

16. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.

17. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

T1 accepted
prepare T2
press TEST

18. Press **TEST** key.

19. Remove the vial from the sample chamber.

20. Add **one DPD No. 3 tablet** straight from the foil to the same water sample and crush the tablet using a clean stirring rod.

21. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.

22. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

T2 accepted
prepare T3
press TEST

23. Press **TEST** key.
Wait for a **reaction period of 2 minutes**.

Countdown
2:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in:

***,** mg/l ClO_2 [Cl]**

as Chlorine dioxide in mg/l Chlorine,
or

***,** mg/l ClO_2**

as Chlorine dioxide in mg/l ClO_2.

***,** mg/l free Cl**
***,** mg/l comb. Cl**
***,** mg/l total Cl**

mg/l free Chlorine
mg/l combined Chlorine
mg/l total Chlorine

Notes:

See next page.

1.1 Methods

Notes: (Chlorine dioxide in the presence of Chlorine)

1. The conversion factor to convert Chlorine dioxide as Chlorine to Chlorine dioxide as ClO_2 is approximately 0.4 (more exactly 0.38).
   mg/l ClO_2 = mg/l ClO_2 [Cl] x 0.38
   ▲ ClO_2[Cl]
   ▼ ClO_2
   (Chlorine dioxide displayed as Chlorine units ClO_2 [Cl] has its origin in swimming poolwater treatment according to DIN 19643.)
2. The total Chlorine result given includes the contribution of the Chlorine dioxide (as Chlorine) reading. For true total Chlorine value subtract the Chlorine dioxide (as Chlorine) reading from the quoted total Chlorine reading.
3. Also see page 55.

1.1 Methods

1 2 0 Chlorine dioxide in absence of Chlorine with Tablet

0.05 – 11 mg/l ClO_2

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.
4. Remove the vial from the sample chamber and **empty it, leaving a few drops remaining in the vial**.
5. Add **one DPD No. 1 tablet** straight from the foil and crush the tablet using a clean stirring rod.
6. Add water sample to the 10 ml mark.
7. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.
8. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

9. Press **TEST** key.

The result is shown in the display

***,** mg/l ClO_2 [Cl]**

as Chlorine dioxide in mg/l Chlorine,

***,** mg/l ClO_2**

or
as Chlorine dioxide in mg/l ClO_2.

Notes:

See page 55.

1.1 Methods

1 0 5 Chlorine HR (KI) with Tablet

5 – 200 mg/l Cl_2

Insert the adapter for 16 mm Ø vials.

1. Fill a clean vial (16 mm Ø) with **8 ml of water sample**, close tightly with the cap.

2. Place the vial in the sample chamber making sure that the marks are aligned. Place the cover on the adapter.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber.

5. Add **one CHLORINE HR (KI) tablet** straight from the foil to the water sample and crush the tablet using a clean stirring rod.

6. Add **one ACIDIFYING GP tablet** straight from the foil to the same water sample and crush the tablet using a clean stirring rod.

7. Close the vial tightly with the cap and swirl several times until the tablets are dissolved.

8. Place the vial in the sample chamber making sure that the marks are aligned. Place the cover on the adapter.

Zero accepted
prepare Test
press TEST

9. Press **TEST** key.

   The result is shown in the display in mg/l Chlorine.

1.1 Methods

Notes:

1. Oxidising agents interfere as they react in the same way as Chlorine.

1.1 Methods

1 3 0 COD LR with Vario Tube Test

0 – 150 mg/l O_2


Ø 16 mm

Insert the adapter for 16 mm Ø vials.

1. Open one white capped reaction vial and add **2 ml deionised water** (this is the blank (Note 1)).

2. Open another white capped reaction vial and add **2 ml of water sample** (this is the sample).

3. Close the vials tightly with the cap. Invert the vial gently several times to mix the contents.
   (CAUTION: The vial will become hot during mixing!)

4. Heat the vials for **120 minutes** in the preheated reactor at a temperature of **150°C**.

5. **(CAUTION: The vials are hot!)**
   Remove the tubes from the heating block and allow them to cool to 60°C or less. Mix the contents by carefully inverting each tube several times while still warm. Then allow the tubes to cool to ambient temperature before measuring. (Note 2).

6. Place the vial (the blank (Note 3, 4)) in the sample chamber making sure that the marks are aligned. Place the cover on the adapter.

prepare Zero
press ZERO

7. Press **ZERO** key.

8. Remove the vial from the sample chamber.

9. Place the vial (the sample (Note 3, 4)) in the sample chamber making sure that the marks are aligned. Place the cover on the adapter.

Zero accepted
prepare Test
press TEST

10. Press **TEST** key.
    The result is shown in the display in mg/l COD.

1.1 Methods

Notes:

1. Run samples and blanks with the same batch of vials. The blank is stable when stored in the dark and can be used for further measurements with vials of the same batch.
2. Do not place the hot vials in the sample chamber. Cool the vials to room temperature for final measurements.
3. Suspended solids in the vial lead to incorrect measurements. For this reason it is important to place the vials carefully in the sample chamber. The precipitate at the bottom of the sample should be not suspended.
4. Clean the outside of the vials with a towel. Finger prints or other marks will be removed.
5. Samples can be measured when the Chloride content does not exceed 1000 mg/l.
6. In exceptional cases, compounds contained in the water cannot be oxidized adequately, so results may be lower than reference methods.

1.1 Methods

1 3 1 COD MR with Vario Tube Test

0 – 1500 mg/l O_2


Ø 16 mm

Insert the adapter for 16 mm Ø vials.

1. Open one white capped reaction vial and add **2 ml deionised water** (this is the blank (Note 1)).

2. Open another white capped reaction vial and add **2 ml of water sample** (this is the sample).

3. Close the vials tightly with the cap. Invert the vial gently several times to mix the contents.
   (CAUTION: The vial will become hot during mixing!)

4. Heat the vials for **120 minutes** in the preheated reactor at a temperature of **150°C**.

5. **(CAUTION: The vials are hot!)**
   Remove the tubes from the heating block and allow them to cool to 60°C or less. Mix the contents by carefully inverting each tube several times while still warm. Then allow the tubes to cool to ambient temperature before measuring. (Note 2).

6. Place the vial (the blank (Note 3, 4)) in the sample chamber making sure that the marks are aligned. Place the cover on the adapter.

prepare Zero
press ZERO

7. Press **ZERO** key.

8. Remove the vial from the sample chamber.

9. Place the vial (the sample (Note 3, 4)) in the sample chamber making sure that the marks are aligned. Place the cover on the adapter.

Zero accepted
prepare Test
press TEST

10. Press **TEST** key.
    The result is shown in the display in mg/l COD.

1.1 Methods

Notes:

1. Run samples and blanks with the same batch of vials. The blank is stable when stored in the dark and can be used for further measurements with vials of the same batch.
2. Do not place the hot vials in the sample chamber. Cool the vials to room temperature for final measurements.
3. Suspended solids in the vial lead to incorrect measurements. For this reason it is important to place the vials carefully in the sample chamber. The precipitate at the bottom of the sample should be not suspended.
4. Clean the outside of the vials with a towel. Finger prints or other marks will be removed.
5. Samples can be measured when the Chloride content does not exceed 1000 mg/l.
6. In exceptional cases, compounds contained in the water cannot be oxidized adequate, so results may be lower than reference methods.
7. For samples under 100 mg/l COD it is recommended to repeat the test with the tube test for COD LR.

1.1 Methods

1 3 2 COD HR with Vario Tube Test

0 – 15 g/l O_2 (≙ 0 – 15 000 mg/l O_2)

Ø 16 mm

Insert the adapter for 16 mm Ø vials.

1. Open one white capped reaction vial and add **0.2 ml deionised water** (this is the blank (Note 1)).

2. Open another white capped reaction vial and add **0.2 ml of water sample** (this is the sample).

3. Close the vials tightly with the cap. Invert the vial gently several times to mix the contents.
   (CAUTION: The vial will become hot during mixing!)

4. Heat the vials for **120 minutes** in the preheated reactor at a temperature of **150°C**.

5. **(CAUTION: The vials are hot!)**
   Remove the tubes from the heating block and allow them to cool to 60°C or less. Mix the contents by carefully inverting each tube several times while still warm. Then allow the tubes to cool to ambient temperature before measuring. (Note 2).

6. Place the vial (the blank (Note 3, 4)) in the sample chamber making sure that the marks are aligned. Place the cover on the adapter.

prepare Zero
press ZERO

7. Press **ZERO** key.

8. Remove the vial from the sample chamber.

9. Place the vial (the sample (Note 3, 4)) in the sample chamber making sure that the marks are aligned. Place the cover on the adapter.

Zero accepted
prepare Test
press TEST

10. Press **TEST** key.
    The result is shown in the display in **g/l** COD.

1.1 Methods

Notes:

1. Run samples and blanks with the same batch of vials. The blank is stable when stored in the dark and can be used for further measurements with vials of the same batch.
2. Do not place the hot vials in the sample chamber. Cool the vials to room temperature for final measurements.
3. Suspended solids in the vial lead to incorrect measurements. For this reason it is important to place the vials carefully in the sample chamber. The precipitate at the bottom of the sample should be not suspended.
4. Clean the outside of the vials with a towel. Finger prints or other marks will be removed.
5. Samples can be measured when the Chloride content does not exceed 10 000 mg/l.
6. In exceptional cases, compounds contained in the water cannot be oxidized adequate, so results may be lower than reference methods.
7. For samples under 1 g/l COD it is recommended to repeat the test with the test kit for COD MR or for samples under 0,1 g/l COD with the tube test COD LR.

1.1 Methods

2 0 4 Colour, true and apparent (APHA Platinum-Cobalt Standard Method)

0 – 500 Pt-Co units (mg/l Pt)

Sample preparation (Note 4):

Step A

Filter approx. **50 ml deionised water** through a membrane filter with a pore width of 0.45 µm.
Discard the filtrate. Filter another **50 ml deionised water** and keep it for zeroing.

Step B

Filter approx. **50 ml water sample** using the same filter. Keep this filtrate for sample measurement.


Ø 24 mm

1. Fill a clean vial (24 mm Ø) with **10 ml of the filtrated deionised water** (from Step A), close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber and empty it completely.

5. Rinse the vial with the filtrated water sample and fill with **10 ml filtrated water sample** (from Step B).

6. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

7. Press **TEST** key.

The result is shown in the display in Pt-Co units (mg/l Pt).

1.1 Methods

Notes:

1. This colour scale was originally developed by A. Hazen as a visual comparison scale. It is therefore necessary to ascertain whether the extinction maximum of the water sample is in the range from 420 to 470 nm, as this method is only suitable for water samples with yellowish to yellowish-brown coloration. Where applicable, a decision should be made based on visual inspection of the water sample.
2. This method 204 – Colour (Hazen) – is calibrated on the basis of the standards specified by "Standard Methods for the Examination of Water and Wastewater" (also see EN ISO 7887:1994).
   1 Pt-Co colour unit = 1 mg/L of platinum as chloroplatinate ion
3. The estimated detection limit is 15 mg/L Pt.
4. Colour may be expressed as "apparent" or "true" colour. The apparent colour is defined as the colour of a solution due to dissolved substances and suspended particles in the sample. This manual describes the determination of true colour by filtration of the water sample. To determine the apparent colour, non-filtrated deionised water and sample are measured.
5. Sample collection, preservation and storage:
   Pour the water sample into clean glass or plastic containers and analyse as soon as possible after the sample is taken. If this is not possible, fill the container right up to the top and seal tightly. Do not stir the sample; avoid lengthy contact with the air.
   The sample may be stored in a dark place at a temperature of 4°C for 24 hours. Before performing measurements, the water sample must be brought up to room temperature.

1.1 Methods

(1) (5) (0) **Copper with Tablet**

0.05 – 5 mg/l Cu

Copper
>> diff
free
total

The following selection is shown in the display:

>> diff

for the differentiated determination of free, combined and total Copper.

>> free

for the determination of free Copper.

>> total

for the determination of total Copper.

Select the desired determination with the arrow keys [▲] and [▼]. Confirm with [↵] key.

Note:

1. If ??? is displayed at the diffentiated test result see page 262.

1.1 Methods

(1) (5) (0)

Copper, differentiated determination with Tablet

0.05 – 5 mg/l Cu

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.
4. Remove the vial from the sample chamber.
5. Add **one COPPER No. 1 tablet** straight from the foil to the water sample and crush the tablet using a clean stirring rod.
6. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.
7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare T1
press TEST

8. Press **TEST** key.
9. Remove the vial from the sample chamber.
10. Add **one COPPER No. 2 tablet** straight from the foil to the same water sample and crush the tablet using a clean stirring rod.
11. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.
12. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

T1 accepted
prepare T2
press TEST

13. Press **TEST** key.

The result is shown in the display in:
mg/l free Copper
mg/l combined Copper
mg/l total Copper

***,** mg/l free Cu**
***,** mg/l comb Cu**
***,** mg/l total Cu**

1.1 Methods

1 5 0 Copper, free with Tablet

0.05 – 5 mg/l Cu

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.
4. Remove the vial from the sample chamber.
5. Add **one COPPER No. 1 tablet** straight from the foil to the water sample and crush the tablet using a clean stirring rod.
6. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.
7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.

The result is shown in the display in mg/l free Copper.

1.1 Methods

1 5 0 Copper, total with Tablet

0.05 – 5 mg/l Cu

Ø 24 mm

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.
4. Remove the vial from the sample chamber.
5. Add **one COPPER No. 1 tablet and one COPPER No. 2 tablet** straight from the foil to the water sample and crush the tablets using a clean stirring rod.
6. Close the vial tightly with the cap and swirl several times until the tablets are dissolved.
7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.

The result is shown in the display in mg/l total Copper.

1.1 Methods

1 5 3 Copper, free (Note 1) with Vario Powder Pack

0.05 – 5 mg/l Cu

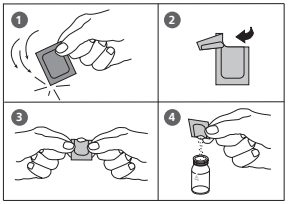
Ø 24 mm

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample,** close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.



4. Remove the vial from the sample chamber.

5. Add the contents of **one Vario Cu 1 F10 Powder Pack** straight from the foil to the water sample.

6. Close the vial tightly with the cap and swirl several times to mix the contents (Note 3).

7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.

   Wait for a **reaction period of 2 minutes**.

Countdown
2:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l Copper.

1.1 Methods

Notes:

1. For determination of total Copper digestion is required.
2. Extremely acid water samples (pH 2 or less) must be adjusted between pH 4 and pH 6 before the reagent is added (with 8 mol/l Potassium hydroxide solution KOH). Caution: pH values above 6 can lead to Copper precipitation.
3. Accuracy is not affected by undissolved powder.
4. Interferences:

Cyanide, CN^-	Cyanide prevents full colour development. Add 0.2 ml Formaldehyde to 10 ml water sample and wait for a reaction time of 4 minutes (Cyanide is masked). After this perform test as described. Multiply the result by 1.02 to correct the sample dilution by Formaldehyde.
Silver, Ag^+	If a turbidity remains and turns black, silver interference is likely. Add 10 drops of saturated Potassium chloride solution to 75 ml of water sample. Filtrate through a fine filter. Use 10 ml of the filtered water sample to perform test.

1.1 Methods

1 5 7 Cyanide with Reagent Test

0.01 – 0.5 mg/l CN

Ø 24 mm

1. Fill a clean vial (24 mm Ø) with **2 ml of water sample and 8 ml of deionised water**, close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber.

5. Add **2 level spoons No. 4 (grey) Cyanide-11** into the prepared water sample, close the vial tightly with the cap and invert several times to mix the contents.

6. Add **2 level spoons No. 4 (grey) Cyanide-12**, close the vial tightly with the cap and invert several times to mix the contents.

7. Fill the vial with drops of the same size by holding the bottle vertically and squeeze slowly:

   3 drops of Cyanide-13

8. Close the vial tightly with the cap and invert several times to mix the contents.

9. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

10. Press **TEST** key.

    Wait for a reaction **period of 10 minutes.**

Countdown
10:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l Cyanide.

1.1 Methods

Notes:

1. Only free Cyanide and Cyanides that can be destroyed by Chlorine are determined by this test.
2. In the present of Thiocyanate, heavy metal complexes, colorants or aromatic amines, the cyanide must be separated out by distillation before analysis is performed.
3. **Store the reagents in closed containers at a temperature of + 15°C to + 25°C.**

1.1 Methods

1 6 0 Cyanuric acid with Tablet

2 – 160 mg/l Cys

Ø 24 mm

1. Fill a clean vial (24 mm Ø) with **5 ml of the water sample** and **5 ml deionised water** (Note 1), close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber.

5. Add **one CYANURIC ACID tablet** straight from the foil to the prepared water sample and crush the tablet using a clean stirring rod.

6. Close the vial tightly with the cap and swirl several times until the tablet is dissolved (Note 2, 3).

7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.

The result is shown in the display in mg/l Cyanuric acid.

1.1 Methods

Notes:

1. Use deionised water or tap water free of Cyanuric acid.
2. If Cyanuric acid is present a cloudy solution will occur.
   Small single particles are not necessarily caused by Cyanuric acid.
3. Dissolve the tablet completely (therefore swirl the vial approx. 1 minute).
   Un-dissolved particles of the tablet can cause results that are too high.

1.1 Methods

(1) (6) (5) DEHA (N,N-Diethylhydroxylamine) with Tablet and Liquid Reagent

20 – 500 µg/l DEHA / 0.02 – 0.5 mg/l DEHA

Ø 24 mm

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample,** close tightly with the cap (Note 2).
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.
4. Remove the vial from the sample chamber.
5. Fill the vial with drops of the same size by holding the bottle vertically and squeeze slowly:

   6 drops (0.25ml) of DEHA solution

6. Close the vial tightly with the cap and swirl several times to mix the contents.
7. Add **one DEHA tablet** straight from the foil to the same water sample and crush the tablet using a clean stirring rod.
8. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.
9. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

10. Press TEST key.

    Wait for a **reaction period of 10 minutes**.

Countdown
10:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display as DEHA.

1.1 Methods

Notes:

1. Application: Testing of residual corrosion inhibitors (Oxygen scavengers) in boiler feed water or condensate.
2. Before using clean the vials with Hydrochloric acid (approx. 20%). Rinse thoroughly with deionised water.
3. Keep the sample dark during colour development time. UV-light (sunlight) causes high measurement results.
4. Ideal temperature for full colour development is 20 °C ± 2 °C.
5. Interferences:
   - Iron (II) interferes at all concentrations:
     Repeat the test procedure but without adding the DEHA solution. If the displayed result is above 20 µg/l subtract this value from the DEHA test result.
   - Substances which reduce Iron (III) interfere. Substances which complex iron strongly may interfere also.
   - Substances which may interfere when present in concentrations at:

Borate (as $Na_2B_4O_7$)	500 mg/l
Cobalt	0.025 mg/l
Copper	8.0 mg/l
Hardness (as $CaCO_3$)	1000 mg/l
Lignosulfonates	0.05 mg/l
Manganese	0.8 mg/l
Molybdenum	80 mg/l
Nickel	0.8 mg/l
Phosphate	10 mg/l
Phosphonates	10 mg/l
Sulfate	1000 mg/l
Zinc	50 mg/l

6. There is an option to change the unit from mg/l to µg/l.
   ▲ mg/l
   ▼ µg/l

1.1 Methods

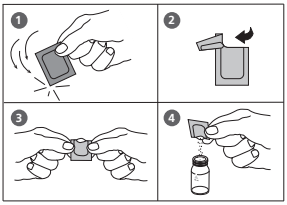
1 6 7 DEHA (N,N-Diethylhydroxylamin) with Vario Powder Pack and Liquid Reagent

20 – 500 µg/l DEHA / 0.02 – 0.5 mg/l DEHA

Use two clean vials (24 mm Ø) and mark one as blank for zeroing (Note 2).

1. Fill a clean vial with **10 ml deionised water** (this is the blank).

2. Fill the second clean vial with **10 ml of the water sample** (this is the sample).



3. Add the contents of **one Vario OXYSCAV 1 Rgt Powder Pack** straight from the foil into each vial.

4. Close the vials tightly with the caps and swirl several times to mix the contents.

5. Add **0.20 ml VARIO DEHA 2 Rgt Solution** to each vial (Note 4).

6. Close the vials tightly with the caps and swirl several times to mix the contents.

Countdown 1
10:00
start: ↵

7. Press **[↵]** key.

   Wait for a reaction **period of 10 minutes** (Note 5).

   After the reaction period is finished proceed as follows:

8. Place the vial (the blank) in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

9. Press **ZERO** key.

10. Remove the vial from the sample chamber.

11. Place the vial (the sample) in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

12. Press **TEST** key.

    The result is shown in the display as DEHA.

1.1 Methods

Notes:

1. Application: Testing of residual corrosion inhibitors (Oxygen scavengers) in boiler feed water or condensate.
2. Before using clean the vials with Hydrochloric acid (approx. 20%). Rinse thoroughly with deionised water.
3. Ideally temperature for full colour development is 25°C ± 3 °C.
4. Volume should always be metered by using suitable pipette (class A).
5. Keep blank and sample dark during colour development time. UV-light (sunlight) causes high measurement results.
6. Interferences:
   - Iron (II) interferes at all concentrations:
     Repeat the test procedure but without adding the VARIO DEHA Rgt 2 solution. If the displayed result is above 20 µg/l subtract this value from the DEHA test result.
   - Substances which reduce Iron (III) interfere. Substances which complex iron strongly may interfere also.
   - Substances who may interfere when present in concentrations at:

Borate (as $Na_2B_4O_7$)	500 mg/l
Cobalt	0.025 mg/l
Copper	8.0 mg/l
Hardness (as $CaCO_3$)	1000 mg/l
Lignosulfonates	0.05 mg/l
Manganese	0.8 mg/l
Molybdenum	80 mg/l
Nickel	0.8 mg/l
Phosphate	10 mg/l
Phosphonates	10 mg/l
Sulfate	1000 mg/l
Zinc	50 mg/l

7. There is an option to change the unit from mg/l to µg/l.
   ▲ mg/l
   ▼ µg/l

1.1 Methods

1 7 0 Fluoride with Liquid Reagent

0.05 – 2 mg/l F

Caution: See notes!

1. Fill a clean vial (24 mm Ø, Note 8) with **exactly 10 ml of water sample** (Note 4), close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber.

5. Add **exactly 2 ml SPADNS reagent solution** (Note 4) to the water sample.
Caution: Vial is filled up to the top! (Note 8)

6. Close the vial tightly with the cap and swirl several times to mix the contents.

7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.

The result is shown in the display in mg/l Fluoride.

1.1 Methods

Notes:

1. The same batch of SPADNS reagent solution must be used for adjustment and test. The adjustment process needs to be performed for each new batch of SPADNS reagent solution (see Standard Methods 20th, 1998, APHA, AWWA, WEF 4500 F D., S. 4-82). The procedure is described in chapter 2.4.5 "Calibration – Fluoride Method 170" on page 234.
2. During adjustment and test the same vial should be used for zeroing and test, as different vials may exhibit minor tolerances.
3. The calibration solution and the water samples to be tested should have the same temperature (± 1°C).
4. As the test result is highly dependent on exact sample and reagent volumes, the sample and reagent volumes should always be metered by using a 10 ml resp. 2 ml volumetric pipette (class A).
5. The accuracy of the test methods decreases above a level of 1.2 mg/l Fluoride. Although the results are sufficiently accurate for most applications, even more exact results can be achieved by 1:1 dilution of the sample prior to use and subsequent multiplication of the result by 2.
6. SPADNS reagent solution contains Arsenite. Chlorine concentrations up to 5 mg/l do not interfere.
7. Seawater and wastewater samples must be distilled.
8. It is convenient to use special vials with larger volume.

1.1 Methods

(1) (9) (0) Hardness, Calcium with Tablet

50 – 900 mg/l $CaCO_3$

1. Fill a clean vial (24 mm Ø) with **10 ml deionised water**.
2. Add **one CALCHECK tablet** straight from the foil to the deionised water and crush the tablet using a clean stirring rod.
3. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.
4. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

5. Press **ZERO** key.

Countdown
2:00

Wait for a **reaction period of 2 minutes**.

After the reaction period is finished the measurement starts automatically.

6. Remove the vial from the sample chamber.
7. Add **2 ml of water sample** to the prepared vial. **Caution: Vial is filled up to the top! (Note 4)**
8. Close the vial tightly with the cap and swirl several times (5x) to mix the contents.
9. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

10. Press **TEST** key.

The result is shown in the display as Calcium Hardness.

1.1 Methods

Notes:

1. Strong alkaline or acidic water samples must be adjusted between pH 4 and pH 10 before the tablet is added (use 1 mol/l Hydrochloric acid resp. 1 mol/l Sodium hydroxide).
2. The tolerance of the method is increasing with higher concentrations. When diluting samples, this should be take into account, always measuring in the first third of the range.
3. This method was developed from a volumetric procedure for the determination of calcium. Due to undefined conditions, the deviations from the standardised method may be greater.
4. It is convenient to use special vials with larger volume.
5. ▲ $CaCO_3$
   °dH
   °eH
   °fH
   ▼ °aH

1.1 Methods

1 9 1 Hardness, Calcium 2T with Tablet

0 – 500 mg/l $CaCO_3$

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.
4. Remove the vial from the sample chamber.
5. Add **one CALCIO H No. 1 tablet** straight from the foil to the 10 ml water sample, crush the tablet using a clean stirring rod and dissolve the tablet completely.
6. Add **one CALCIO H No. 2 tablet** straight from the foil to the same water sample and crush the tablet using a clean stirring rod.
7. Close the vial tightly with the cap and swirl gently several times until the tablet is completely dissolved.
8. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

9. Press **TEST** key.

   Wait for a **reaction period of 2 minutes**.

Countdown
2:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display as Calcium Hardness.

1.1 Methods

Notes:

1. To optimise the readings an optional batch related calibration can be performed using Mode 40, see page 232.
2. Strong alkaline or acidic water samples must be adjusted to a pH-value between pH 4 and 10 before the tablets are added (use 1 mol/l Hydrochloride acid resp. 1 mol/l Sodium hydroxide).
3. For accurate test results exactly 10 ml of water sample must be taken for the test.
4. This method was developed from a volumetric procedure for the determination of Calcium Hardness. Due to undefined conditions, the deviations from the standardised method may be greater.
5. The tolerance of the method is increasing with higher concentrations. When diluting samples, this should be taken in account, always measuring in the first third of the range.
6. Interferences:
   - Magnesium hardness up to 200 mg/l $CaCO_3$ does not interfere.
   - Iron concentration above 10 mg/l may cause low results.
   - Zinc concentration above 5 mg/l may cause high results.
7. ▲ $CaCO_3$
   °dH
   °eH
   °fH
   ▼ °aH

1.1 Methods

(2) (0) (0)

Hardness, total with Tablet

2 – 50 mg/l $CaCO_3$

Ø 24 mm

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber.

5. Add **one HARDCHECK P tablet** straight from the foil to the water sample and crush the tablet using a clean stirring rod.

6. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.

7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.

   Wait for a **reaction period of 5 minutes**.

Countdown
5:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display as total Hardness.

1.1 Methods

Notes:

1. Strong alkaline or acidic water samples must be adjusted between pH 4 and pH 10 before the tablet is added (use 1 mol/l Hydrochloric acid resp. 1mol/l Sodium hydroxide).
2. Conversion table:

	mg/l $CaCO_3$	°dH	°fH	°eH
1 mg/l $CaCO_3$	----	0,056	0,10	0,07
1 °dH	17,8	----	1,78	1,25
1 °fH	10,0	0,56	----	0,70
1 °eH	14,3	0,80	1,43	----

3. ▲ $CaCO_3$
   °dH
   °eH
   °fH
   ▼ °aH

1.1 Methods

(2) (0) (1)

Hardness, total HR with Tablet

20 – 500 mg/l $CaCO_3$

1. Fill a clean vial (24 mm Ø) with **1 ml of water sample** and **9 ml of deionised water**, close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber.

5. Add **one HARDCHECK P tablet** straight from the foil to the water sample and crush the tablet using a clean stirring rod.

6. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.

7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.

   Wait for a **reaction period of 5 minutes**.

Countdown
5:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display as total Hardness.

1.1 Methods

Notes:

1. Strong alkaline or acidic water samples must be adjusted between pH 4 and pH 10 before the tablet is added (use 1 mol/l Hydrochloric acid resp. 1mol/l Sodium hydroxide).
2. Conversion table:

	mg/l $CaCO_3$	°dH	°fH	°eH
1 mg/l $CaCO_3$	----	0,056	0,10	0,07
1 °dH	17,8	----	1,78	1,25
1 °fH	10,0	0,56	----	0,70
1 °eH	14,3	0,80	1,43	----

3. ▲ $CaCO_3$
   °dH
   °eH
   °fH
   ▼ °aH

1.1 Methods

2 0 5 Hydrazine with Powder Reagent

0.05 – 0.5 mg/l N_2H_4 / 50 – 500 µg/l N_2H_4

Ø 24 mm

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample** (Note 1, 2), close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber.

5. Add **1 g HYDRAZINE test powder** (Note 3) to the water sample.

6. Close the vial tightly with the cap and swirl several times to mix the contents.

Countdown
10:00
start: ↵

7. Press [↵] key.
   Wait for a **reaction period of 10 minutes.**

   After the reaction period is finished proceed as follows:

8. Any slight turbidity that occurs when the reagent is added must be removed by filtration (Note 4).

9. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

10. Press **TEST** key.

    The result is shown in the display as Hydrazine.

1.1 Methods

Notes:

1. If the water sample is cloudy, you must filter it before performing the zero calibration.
2. The temperature of the water sample should not exceed 21 °C.
3. Using the Hydrazine spoon: 1 g is equivalent to one level spoon.
4. Qualitative folded filter papers for medium precipitates are recommended.
5. In order to check whether the reagent has aged (if it has been stored for a lengthy period), perform the test as described above using tap water. If the result is above the detection limit of 0.05 mg/l, you should only use the reagent with reservations as there may be a major deviation in results.
6. There is an option to change the unit from mg/l to µg/l.
   ▲ mg/l
   ▼ µg/l

1.1 Methods

2 0 6 Hydrazine with Vario Liquid Reagent

0.005 – 0.6 mg/l N_2H_4 / 5 – 600 µg/l N_2H_4


Ø 24 mm

Use two clean vials (24 mm Ø) and mark one as blank for zeroing.

1. Fill a clean vial with **10 ml deionised water** (this is the blank).
2. Add **1 ml VARIO Hydra 2 Rgt Solution** into the vial (Note 3).
3. Close the vial tightly with the cap and swirl several times to mix the contents.
4. Place the vial (the blank) in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

5. Press **ZERO** key.
6. Remove the vial from the sample chamber.
7. Fill the second clean vial with **10 ml of the water sample** (this is the sample).
8. Add **1 ml VARIO Hydra 2 Rgt Solution** into the vial.
9. Close the vial tightly with the cap and swirl several times to mix the contents.
10. Place the vial (the blank) in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

11. Press **TEST** key.
    Wait for a **reaction period of 12 minutes**.

Countdown
12:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display as Hydrazine.

1.1 Methods

Notes:

1. Samples cannot be preserved and must be analysed immediately.
2. Sample temperature should be 21 °C ± 4 °C.
3. The blank may develop a faint yellow colour due to the reagent.
4. Interferences:
   - Ammonia causes no interferences up to 10 mg/l.
     At a concentration of 20 mg/l it is possible that the test result increases by 20 %.
   - Morpholine does not interfere up to 10 mg/l.
   - Highly coloured or turbid samples:
     Mix 1 part deionised water with 1 part household bleach. Add 1 drop of this mixture into 25 ml water sample and mix. Use 10 ml prepared sample in place of deionised water in point 1.
     Note: at point 7 use the unprepared water sample.
     Principle: Hydrazine is oxidised by household bleach. Colour interference will be eliminated by zeroing.
5. There is an option to change the unit from mg/L to µg/L.
   ▲ mg/l
   ▼ µg/l

1.1 Methods

(2) (0) (7) **Hydrazine with Vacu-vials® K-5003** (see Notes)

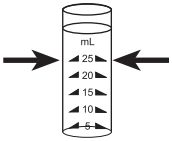
0.01 – 0.7 mg/l N_2H_4 / 10 – 700 µg/l N_2H_4

Insert the adapter for 13 mm Ø vials.

1. Place the blank in the sample chamber. The blank is part of the test kit.

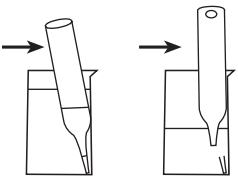
prepare Zero
press ZERO

2. Press **ZERO** key.



3. Remove the blank from the sample chamber.

4. Fill the sample container to the 25 ml mark with the water sample.



5. Place one Vacu-vial® in the sample container. Snap the tip by pressing the vial against the side of the sample container. The Vacu-vial® breaks at the neck and the vial fills automatically. A small volume of inert gas remains in the Vacu-vial®.

6. Mix the contents of the Vacu-vial® by inverting it several times, allowing the bubble to move from one end to the other. Dry the outside of the vial.

7. Place the Vacu-vial® in the sample chamber.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.
Wait for a **reaction period of 10 minutes**.

Countdown
10:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display as Hydrazine.

1.1 Methods

Notes:

1. This method is adapted from CHEMetrics.
2. Read the original test instruction and the MSDS (delivered with the test) before performing the test. MSDS also available at www.chemetrics.com.
3. Vacu-vials® is a registered trade mark of the company CHEMetrics, Inc. / Calverton, U.S.A.
4. There is an option to change the unit from mg/l to µg/l.
   ▲ mg/l
   ▼ µg/l

1.1 Methods

2 1 0 Hydrogen peroxide with Tablet

0.03 – 3 mg/l H_2O_2

Ø 24 mm

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber and **empty it, leaving a few drops remaining in the vial**.

5. Add **one HYDROGENPEROXIDE LR tablet** straight from the foil and crush the tablet using a clean stirring rod.

6. Add water sample to the 10 ml mark.

7. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.

8. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

9. Press **TEST** key.

   Wait for a **reaction period of 2 minutes**.

Countdown
2:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l Hydrogen peroxide.

1.1 Methods

Notes:

1. Vial cleaning:
   As many household cleaners (e.g. dishwasher detergent) contain reducing substances, the subsequent determination of Hydrogen peroxide may show lower results. To avoid any measurement errors, only use glassware free of Chlorine demand.
   Preparation: Put all applicable glassware into Sodium hypochlorite solution (0.1 g/l) for one hour, then rinse all glassware thoroughly with deionised water.
2. Preparing the sample:
   When preparing the sample, the lost of Hydrogen peroxide, e.g. by pipetting or shaking, must be avoided. The analysis must take place immediately after taking the sample.
3. The DPD colour development is carried out at a pH value of 6.2 to 6.5. The reagent tablet therefore contains a buffer for the pH adjustment.
   Strong alkaline or acidic water samples must be adjusted between pH 6 and pH 7 before the tablet is added (use 0.5 mol/l Sulfuric acid resp. 1 mol/l Sodium hydroxide).
4. Exceeding the measuring range:
   Concentrations above 5 mg/l Hydrogen peroxide can lead to results showing 0 mg/l. In this case, the water sample must be diluted with water free of Hydrogen peroxide. 10 ml of the diluted sample should be mixed with the reagent and the measurement repeated.

   Oxidising agents such as Chlorine, Ozone etc. interfere as they react in the same way as Hydrogen peroxide.

1.1 Methods

2 1 5 Iodine with Tablet

0.05 – 3.6 mg/l I

Ø 24 mm

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.
4. Remove the vial from the sample chamber and **empty it, leaving a view drops remaining in the vial.**
5. Add **one DPD No. 1 tablet** straight from the foil to the water sample and crush the tablet using a clean stirring rod.
6. Add water sample to the 10 ml mark.
7. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.
8. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

9. Press **TEST** key.

The result is shown in the display in mg/l Iodine.

1.1 Methods

Notes:

1. Oxidising reagents, such as Chlorine, Bromine, etc. interfere as they react in the same way as Iodine.

1.1 Methods

220 Iron with Tablet

0.02 – 1 mg/l Fe

*Determination of total dissolved Iron Fe^{2+} and Fe^{3+}

222 Iron with Vario Powder Pack

0.02 – 3 mg/l Fe

*Determination of all dissolved iron and most undissolved forms of iron.

223 Iron, total with Vario Powder Pack

0.02 – 1.8 mg/l Fe

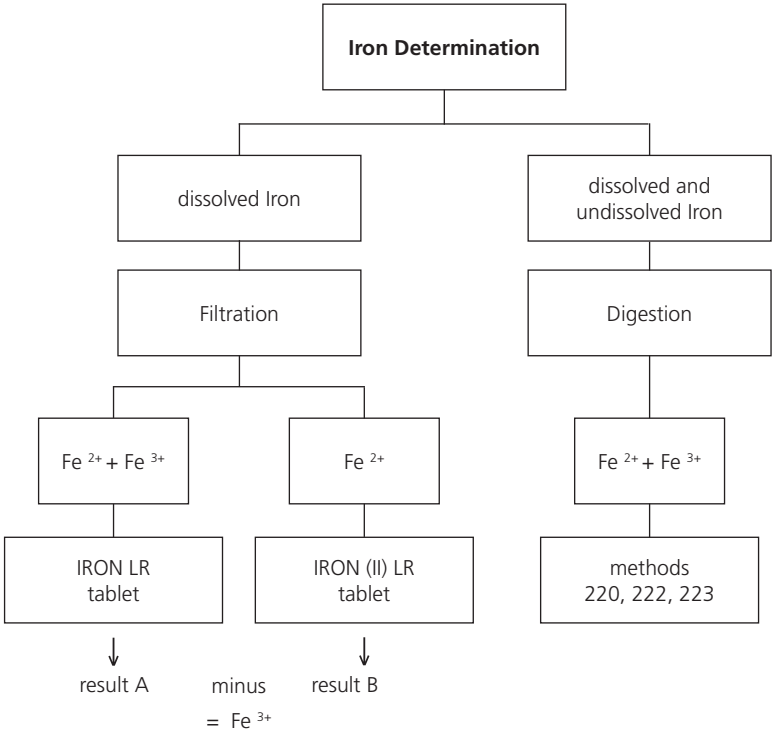
*Determination of all dissolved iron and most undissolved forms of iron; most undissolved iron oxides are recovered by the reagent.

*This information refers to analysis of the water sample without digestion.

Further information can be found in the method notes.

1.1 Methods

Notes:



Digestion procedure for the determination of total dissolved and undissolved iron.

1. Add 1 ml of concentrated sulfuric acid to 100 ml water sample. Heat and boil for 10 minutes or until all particles are dissolved. After cooling down, the sample is set to a pH-value of 3 to 6 by using ammonia solution. Refill with deionised water to the previous volume of 100 ml and mix well. 10 ml of this pre-treated solution is used for the following analysis. Perform as described by the selected test method.
2. Water which has been treated with organic compounds like corrosion inhibitors must be oxidised where necessary to break down the iron. Therefore add 1 ml concentrated sulfuric acid and 1 ml concentrated nitric acid to 100 ml water sample and boil to approx. half volume. After cooling down, proceed as described above.

1.1 Methods

2 2 0 Iron (Note 1) with Tablet

0.02 – 1 mg/l Fe

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber.

5. Add **one IRON LR tablet** straight from the foil to the water sample and crush the tablet using a clean stirring rod.

6. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.

7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.
Wait for a **reaction period of 5 minutes**.

Countdown
5:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l Iron.

1.1 Methods

Notes:

1. This method determines the total dissolved Iron as Fe^{2+} and Fe^{3+}.
2. The IRON (II) LR tablet is used for differentiation – as described above – instead of the IRON LR tablet.
3. For the determination of total dissolved and undissolved iron digestion is required. An example is described on page 105.

1.1 Methods

2 2 2 Iron (Note 1) with Vario Powder Pack

0.02 – 3 mg/l Fe

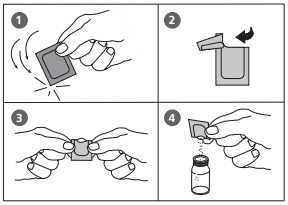
1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.

2. Place the vial in the sample chamber making sure that the marks ⧖ are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber.



5. Add the contents of **one Vario Ferro F10 Powder Pack straight** from the foil to the water sample.

6. Close the vial tightly with the cap and swirl several times to mix the contents (Note 4).

7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.

   Wait for a **reaction period of 3 minutes (Note 5)**.

Countdown
3:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l Iron.

1.1 Methods

Notes:

1. The reagent reacts with all dissolved iron and most undissolved forms of iron in the water sample.
2. Iron oxide requires prior digestion: use mild, vigorous or Digesdahl digestion (e.g. for digestion with acid see page 105).
3. Very strong alkaline or acidic water samples must be adjusted to a pH value between 3 and 5 before analysis.
4. Accuracy is not affected by undissolved powder.
5. Water samples containing visible rust should be allowed to react for at least five minutes.

1.1 Methods

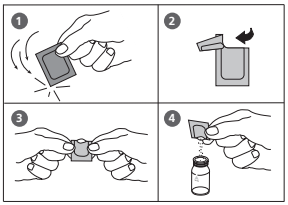
(2) (2) (3)

Iron, total (TPTZ, Note 1) with Vario Powder Pack

0.02 – 1.8 mg/l Fe

Use two clean vials (24 mm Ø) and mark one as blank for zeroing.

1. Fill a clean vial with **10 ml deionised water** (this is the blank).



2. Fill the second clean vial with **10 ml of the water sample** (this is the sample).

3. Add the contents of **one Vario IRON TPTZ F10 Powder Pack** straight from the foil into each vial.

4. Close the vials tightly with the caps and swirl several times to mix the contents.

Countdown
3:00
start: ↵

5. Press **[↵]** key.

   Wait for a reaction **period of 3 minutes.**

   After the reaction period is finished proceed as follows:

6. Place the vial (the blank) in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

7. Press **ZERO** key.

8. Remove the vial from the sample chamber.

9. Place the vial (the sample) in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

10. Press **TEST** key.

    The result is shown in the display in mg/l Iron.

1.1 Methods

Notes:

1. For determination of total Iron digestion is required.
   TPTZ reagent recovers most insoluble iron oxides without digestion.
2. Rinse all glassware with 1:1 Hydrochloric acid solution first and then rinse with deionised water to remove iron deposits that can cause slightly high results.
3. Strong alkaline or acidic water samples must be adjusted between pH 3 and pH 8 before the reagent is added (use 0.5 mol/l Sulfuric acid resp. 1 mol/l Sodium hydroxide).
4. Interferences:
   When interferences occur, colour development is inhibited or a precipitate is formed.
   The values below refer to a standard with an iron concentration of 0.5 mg/l.
   The following substances do not interfere when present up to the levels given:

Substance	**no inerference to**
Cadmium	4.0 mg/l
Chromium$^{(3+)}$	0.25 mg/l
Chromium$^{(6+)}$	1.2 mg/l
Cobalt	0.05 mg/l
Copper	0.6 mg/l
Cyanide	2.8 mg/l
Manganese	50 mg/l
Mercury	0.4 mg/l
Molybdenum	4.0 mg/l
Nickel	1.0 mg/l
Nitrite Ion	0.8 mg/l

1.1 Methods

2 4 0 Manganese with Tablet

0.2 – 4 mg/l Mn

Ø 24 mm

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber.

5. Add **one MANGANESE LR 1 tablet** straight from the foil to the water sample and crush the tablet using a clean stirring rod.

6. Add **one MANGANESE LR 2 tablet** straight from the foil to the same water sample and crush the tablet using a clean stirring rod.

7. Close the vial tightly with the cap and swirl several times until the tablets are dissolved.

8. Place the vial in the sample chamber making sure that the marks ⧖ are aligned.

Zero accepted
prepare Test
press TEST

9. Press **TEST** key.

   Wait for a **reaction period of 5 minutes.**

Countdown
5:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l Manganese.

1.1 Methods

Note:

1. ▲ Mn
   MnO_4
   ▼ $KMnO_4$

1.1 Methods

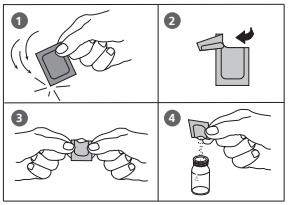
2 4 2 Manganese LR with Vario Powder Pack

0.01 – 0.7 mg/l Mn

Ø 24 mm

Use two clean vials (24 mm Ø) and mark one as blank for zeroing (Note 1).

1. Fill a clean vial with **10 ml of deionised water** (this is the blank).
2. Fill the second clean vial with **10 ml of water sample** (this is the sample).



3. Add the contents of **one Vario Ascorbic Acid Powder Pack** straight from the foil into each vial. (Note 2)
4. Close the vials tightly with the caps and swirl several times to mix the contents.
5. Fill each vial with drops of the same size by holding the bottle vertically and squeeze slowly (Note 3):
   15 drops of Alkaline Cyanide reagent solution
6. Close the vials tightly with the caps and swirl several times to mix the contents.
7. Fill each vial with drops of the same size by holding the bottle vertically and squeeze slowly:
   21 drops of PAN Indicator solution
8. Close the vials tightly with the caps and swirl several times to mix the contents.

Countdown 1
2:00
start: ↵

9. Press **[↵]** key.
   Wait for a **reaction period of 2 minutes** (Note 4).

   After the reaction period is finished proceed as follows:

10. Place the vial (the blank) in the sample chamber making sure that the marks are ⧖ aligned.

prepare Zero
press ZERO

11. Press **ZERO** key.
12. Remove the vial from the sample chamber.
13. Place the vial (the sample) in the sample chamber making sure that the marks are ⧖ aligned.

Zero accepted
prepare Test
press TEST

14. Press **TEST** key.

    The result is shown in the display in mg/l Manganese.

1.1 Methods

Notes:

1. Rinse all glassware with 1:1 Nitric acid solution first and then rinse with deionised water.
2. Water samples that contain more than 300 mg/l $CaCO_3$ hardness: after adding the Vario Ascorbic Acid Powder Pack add additionally 10 drops of Rochelle Salt Solution.
3. After addition of the reagent solution "Alkaline-Cyanide" a cloudy or turbid solution may form in some water samples. The turbidity should disappear after point 7.
4. Water samples containing more than 5 mg/l iron should be allowed to react for at least 10 minutes.
5. Conversion:
   mg/l MnO_4 = mg/l Mn x 2.17
6. ▲ Mn
   MnO_4
   ▼ $KMnO_4$

1.1 Methods

2 4 3 Manganese HR with Vario Powder Pack

0.1 – 18 mg/l Mn

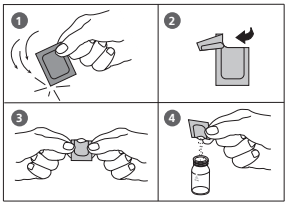
1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber.



5. Add the contents of **one Vario Manganese Citrate Buffer F10 Powder Pack** straight from the foil to the water sample.

6. Close the vial tightly with the cap and swirl several times to mix the contents.

7. Add the contents of **one Vario Sodium periodate F10 Powder Pack** straight from the foil to the same water sample.

8. Close the vial tightly with the cap and swirl several times to mix the contents.

9. Place the vial in the sample chamber making sure that the ⧖ marks are aligned

Zero accepted
prepare Test
press TEST

10. Press **TEST** key.

    Wait for a **reaction period of 2 minutes**.

Countdown
2:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l Manganese.

1.1 Methods

Notes:

1. This test is applicable for the determination of soluble Manganese in water and wastewater.
2. Highly buffered water samples or extreme pH values may exceed the buffering capacity of the reagents and requires sample pre-treatment.
   If samples were acidified for storing, adjust the pH between 4 and 5 with 5 mol/l (5 N) Sodium hydroxide before test. Do not exceed pH 5, as manganese may precipitate.
3. Interferences:

Interfering substance	**Interference level**
Calcium	greater than 700 mg/l
Chloride	greater than 70 000 mg/l
Iron	greater than 5 mg/l
Magnesium	greater than 100 000 mg/l

4. ▲ Mn
   MnO_4
   ▼ $KMnO_4$

1.1 Methods

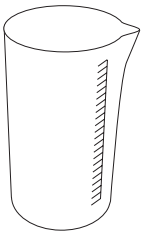
250 Molybdate with Tablet

1 – 50 mg/l MoO_4 / 0.6 – 30 mg/l Mo

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample,** close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.



4. Remove the vial from the sample chamber and **empty the vial**.
5. Fill **20 ml of water sample** in a 100 ml beaker.
6. Add **one MOLYBDATE HR No. 1 tablet** straight from the foil to the water sample and crush the tablet using a clean stirring rod.
7. Add **one MOLYBDATE HR No. 2 tablet** straight from the foil to the same water sample and crush the tablet using a clean stirring rod.
8. Dissolve the tablets using a clean stirring rod.
9. Rinse out the vial with the prepared water sample and then fill to the 10 ml mark.
10. Close the vial tightly with the cap.
11. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

12. Press **TEST** key.

    The result is shown in the display in mg/l Molybdate / Molybdenum.

1.1 Methods

Notes:

1. The tablets must be added in the correct sequence.
2. Under test conditions (pH 3.8 – 3.9) iron does not interfere nor do other metals at levels likely to be found in industrial water systems.
3. Conversions:
   mg/l Mo = mg/l MoO_4 x 0.6
   mg/l Na_2MoO_6 = mg/l MoO_4 x 1.3
4. ▲ MoO_4
   Mo
   ▼ Na_2MoO_4

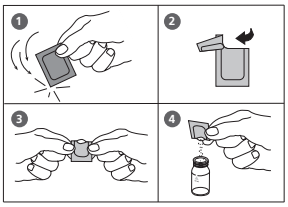
2 5 2 Molybdate / Molybdenum HR with Vario Powder Pack

0.5 – 66 mg/l MoO_4 / 0.3 – 40 mg/l Mo

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.
4. Remove the vial from the sample chamber.



5. Add the contents of **one Vario Molybdenum HR 1 F10 Powder Pack** straight from the foil to the water sample.
6. Close the vial tightly with the cap and swirl several times to mix the contents.
7. Add the contents of **one Vario Molybdenum HR 2 F10 Powder Pack** straight from the foil to the same water sample.
8. Close the vial tightly with the cap and swirl several times to mix the contents.
9. Add the contents of **one Vario Molybdenum HR 3 F10 Powder Pack** straight from the foil to the same water sample.
10. Close the vial tightly with the cap and swirl several times to mix the contents.
11. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

12. Press **TEST** key.
    Wait for a **reaction period of 5 minutes**.

Countdown
5:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l Molybdate / Molybdenum.

1.1 Methods

Notes:

1. Filter turbid water samples using filter paper and funnel before analysis.
2. Highly buffered water samples or extreme pH values should be adjusted to a pH of nearly 7 with 1 mol/l Nitric acid or 1 mol/l Sodium hydroxide.
3. Concentrations above 10 mg/l Cu causes too high test values if the reaction time of 5 minutes is increased. So it is very important to perform the test procedure as described.
4. Substances which may interfere when present in concentrations at:

Aluminium	50 mg/l
Chromium	1000 mg/l
Iron	50 mg/l
Nickel	50 mg/l
Nitrite	all levels

5. ▲ MoO_4
   Mo
   ▼ Na_2MoO_4

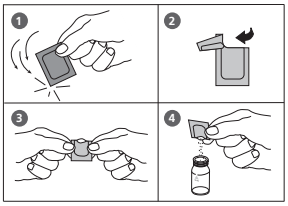
1.1 Methods

2 6 5 Nitrate with Tube Test

1 – 30 mg/l N

Ø 16 mm

Insert the adapter for 16 mm Ø vials.



1. Open one white capped reaction vial (Reagent A) and add **1 ml deionised water** (this is the blank).

2. Open another white capped reaction vial (Reagent A) and add **1 ml of water sample** (this is the sample).

3. Add the contents of **one Vario Nitrate Chromotropic Powder Pack straight** from the foil into each vial.

4. Close the vials tightly with the caps and invert gently several times (10 x) to mix the contents (Note 1).

Countdown
5:00
start: ↵

5. Press **[↵]** key.

   Wait for a **reaction period of 5 minutes**.

6. After the reaction period is finished proceed as follows:

7. Place the vial (the blank) in the sample chamber making sure that the marks are ▽ aligned. Place the cover on the adapter.

prepare Zero
press ZERO

8. Press **ZERO** key.

9. Remove the vial from the sample chamber.

10. Place the vial (the sample) in the sample chamber making sure that the marks are ▽ aligned. Place the cover on the adapter.

Zero accepted
prepare Test
press TEST

11. Press **TEST** key.

    The result is shown in the display in mg/l Nitrate.

1.1 Methods

Notes:

1. Some solids may not dissolve.
2. Conversion:
   mg/l NO_3 = mg/l N x 4.43
3. ▲ N
   ▼ NO_3

1.1 Methods

2 7 0 Nitrite with Tablet

0.01 – 0.5 mg/l N

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.

2. Place the vial in the sample chamber making sure that the marks ⧖ are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber.

5. Add **one NITRITE LR tablet** straight from the foil to the water sample and crush the tablet using a clean stirring rod.

6. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.

7. Place the vial in the sample chamber making sure that the marks ⧖ are aligned.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.
Wait for a **reaction period of 10 minutes**.

Countdown
10:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l Nitrite.

1.1 Methods

Notes:

1. The following ions can produce interferences since under the reaction conditions they cause precipitation:
   Antimony (III), Iron (III), Lead, Mercury (I), Silver, Chloroplatinate, Metavanadate and Bismuth.

   Copper (II)-ions may cause lower test results as they accelerate the decomposition of the Diazonium salt.

   It is unlikely in practice that these interfering ions will occur in such high concentrations that they cause significant reading errors.
2. Conversion:
   mg/l NO_2 = mg/l N x 3.29
3. ▲ N
   ▼ NO_2

1.1 Methods

2 7 2 Nitrite LR with Vario Powder Pack

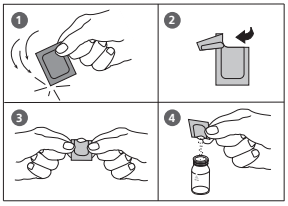
0.01 – 0.3 mg/l N

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.



4. Remove the vial from the sample chamber.

5. Add the contents of **one Vario Nitri 3 Powder Pack** straight from the foil to the water sample.

6. Close the vial tightly with the cap and swirl several times to mix the contents.

7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.

Countdown
20:00

Wait for a **reaction period of 20 minutes**.

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l Nitrite.

1.1 Methods

Notes:

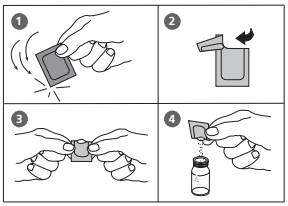
1. Interferences:
   - Strong oxidizing and reducing substances interfere.
   - Cupric and ferrous ions cause low results.
   - Antimonous, Auric, Bismuth, Chloroplatinate, Ferric, Lead, Mercurous, Metavanadate, Silver ions interfere by causing precipitation.
   - In samples with very high concentrations of Nitrate (> 100 mg/L N) a small amount of Nitrite will be found. Such high levels of Nitrate appear to undergo a slight amount of reduction to Nitrite, either spontaneously or during the reaction time of the test.
2. ▲ N
   ▼ NO_2

1.1 Methods

2 8 0 Nitrogen, total LR with Vario Tube Test

0.5 – 25 mg/l N

Ø 16 mm



Insert the adapter for 16 mm Ø vials.

1. **Open two TN Hydroxide LR digestion vials** and add **one Vario TN Persulfate Rgt. Powder Pack** (Note 2, 3).
2. Add **2 ml deionised water** to the prepared vial (this is the blank, Note 4, 5).
3. Add **2 ml of water sample** to the other prepared vial (this is the sample).
4. Close the vials with the caps and shake to mix the contents (at least 30 seconds, Note 6).
5. Heat the vials for **30 minutes** in the preheated reactor at a temperature of **100°C** (Note 7).
6. After 30 minutes remove the vials from the reactor. **(CAUTION: The vials are hot!)**
   Allow the vials to cool to room temperature.
7. Open the cooled digestion vials and add the contents of **one Vario TN Reagent A Powder Pack** to each vial (Note 2).
8. Close the vials with the caps and shake to mix the contents (at least 15 seconds).

Countdown
3:00
start: ↵

9. Press **[↵]** key.
   Wait for a **reaction period of 3 minutes.**

   After the reaction period is finished proceed as follows:

10. Open the digestion vials and add the contents of **one Vario TN Reagent B Powder Pack** to each vial (Note 2).
11. Close the vials with the caps and shake to mix the contents (at least 15 seconds, Note 8).

Countdown
2:00
start: ↵

12. Press **[↵]** key.
    Wait for a **reaction period of 2 minutes**.

    After the reaction period is finished proceed as follows:

13. Open **two TN Acid LR/HR (Reagent C) vials** and add **2 ml of the digested, treated blank** to one vial (this is the blank).
14. Add **2 ml of the digested, treated water sample** to the other TN Acid LR/HR vial (this is the sample).
15. Close the vials with the caps and swirl the vials gently several times to mix the contents (10 x, Note 9). **(CAUTION: Vials warm up).**

1.1 Methods

prepare Zero
press ZERO

Countdown
5:00

Zero accepted
prepare Test
press TEST

16. Place the vial (the blank) in the sample chamber making sure that the marks ⍋ are aligned. Place the cover on the adapter.
17. Press **ZERO** key.
    Wait for a **reaction period of 5 minutes**.

    After the reaction period is finished the measurement starts automatically.
18. Remove the vial from the sample chamber.
19. Place the vial (the sample, Note 10) in the sample chamber making sure that the marks ⍋ are aligned. Place the cover on the adapter.
20. Press **TEST** key.

    The result is shown in the display in mg/l Nitrogen.

Notes:

1. Appropriate safety precautions and a good lab technique should be used during the whole procedure.
2. Use a funnel to add the reagent.
3. Wipe off any Persulfate reagent that may get on the lid or the tube threads.
4. Volumes for samples and blank should always be metered by using 2 ml volumetric pipettes (class A).
5. One blank is sufficient for each set of samples.
6. The reagent may not dissolve completely.
7. It is very important to remove the vials from the reactor after exactly 30 minutes.
8. The reagent will not completely dissolve.
9. Hold the vial in a vertical position with the cap pointing up. Turn the vial upside-down. Wait for all of the solution to flow down to the cap. Return the vial to the upright position. Wait for all the solution to flow to the bottom of the vial. This process is one inversion; 10 inversions = approx. 30 seconds.
10. After zero calibration with the blank it is possible to measure several samples.
11. Large quantities of nitrogen free, organic compounds which are included in some water samples may reduce the effectiveness of the digestion by reacting with the Persulfate reagent. Samples which are well known to contents large quantities of organic compounds must be diluted and digestion and measurement must be repeated for checking the effectiveness of the digestion.
12. Application: for water, wastewater and seawater
13. Interferences:
    Interfering substances that resulted in a concentration change of 10%:
    Bromide more than 60 mg/l and Chloride more than 1000 mg/l produce positive interferences.

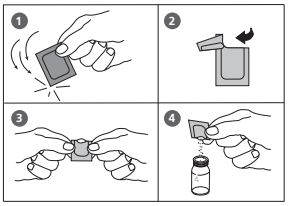
    TN = Total Nitrogen
14. ▲ N
    NH_4
    ▼ NH_3

1.1 Methods

2 8 1 Nitrogen, total HR with Vario Tube Test

5 – 150 mg/l N

Ø 16 mm



Insert the adapter for 16 mm Ø vials.

1. **Open two TN Hydroxide HR digestion vials** and add **one Vario TN Persulfate Rgt. Powder Pack** (Note 2, 3).
2. Add **0.5 ml deionised water** to the prepared vial (this is the blank, Note 4, 5).
3. Add **0.5 ml of water sample** to the other prepared vial (this is the sample).
4. Close the vials with the caps and shake to mix the contents (at least 30 seconds, Note 6).
5. Heat the vials for **30 minutes** in the preheated reactor at a temperature of **100°C** (Note 7).
6. After 30 Minutes remove the vials from the reactor. **(CAUTION: The vials are hot!)**
   Allow the vials to cool to room temperature.
7. Open the cooled digestion vials and add the contents of **one Vario TN Reagent A Powder Pack** to each vial (Note 2).
8. Close the vials with the caps and shake to mix the contents (at least 15 seconds).

Countdown
3:00
start: ↵

9. Press **[↵]** key.
   Wait for a **reaction period of 3 minutes.**
   After the reaction period is finished proceed as follows:
10. Open the digestion vials and add the contents of **one Vario TN Reagent B Powder Pack** to each vial (Note 2).
11. Close the vials with the caps and shake to mix the contents (at least 15 seconds, Note 8).

Countdown
2:00
start: ↵

12. Press **[↵]** key.
    Wait for a **reaction period of 2 minutes**.
    After the reaction period is finished proceed as follows:
13. Open **two TN Acid LR/HR (Reagent C) vials** and add **2 ml of the digested, treated blank** to one vial (this is the blank).
14. Add **2 ml of the digested, treated water sample** to the other TN Acid LR/HR vial (this is the sample).
15. Close the vials with the caps and swirl the vials gently several times to mix the contents (10 x, Note 9).
    (CAUTION: Vials warm up).

1.1 Methods

prepare Zero
press ZERO

Countdown
5:00

Zero accepted
prepare Test
press TEST

16. Place the vial (the blank) in the sample chamber making sure that the ⍋ marks are aligned. Place the cover on the adapter.
17. Press **ZERO** key.
    Wait for a **reaction period of 5 minutes**.

    After the reaction period is finished the measurement starts automatically.
18. Remove the vial from the sample chamber.
19. Place the vial (the sample, Note 10) in the sample chamber making sure that the ⍋ marks are aligned. Place the cover on the adapter.
20. Press **TEST** key.

    The result is shown in the display in mg/l Nitrogen.

Notes:

1. Appropriate safety precautions and a good lab technique should be used during the whole procedure.
2. Use a funnel to add the reagent.
3. Wipe off any Persulfate reagent that may get on the lid or the tube threads.
4. Volumes for samples and blank should always be metered by using suitable pipettes (class A).
5. One blank is sufficient for each set of samples.
6. The reagent may not dissolve completely.
7. It is very important to remove the vials from the reactor after exactly 30 minutes.
8. The reagent will not completely dissolve.
9. Hold the vial in a vertical position with the cap pointing up. Turn the vial upside-down. Wait for all of the solution to flow down to the cap. Return the vial to the upright position. Wait for all the solution to flow to the bottom of the vial. This process is one inversion; 10 inversions = approx. 30 seconds.
10. After zero calibration with the blank it is possible to measure several samples.
11. Large quantities of nitrogen free, organic compounds which are included in some water samples may reduce the effectiveness of the digestion by reacting with the Persulfate reagent. Samples which are well known to contents large quantities of organic compounds must be diluted and digestion and measurement must be repeated for checking the effectiveness of the digestion.
12. Application: for water, wastewater and seawater
13. Interferences:
    Interfering substances that resulted in a concentration change of 10%:
    Bromide more than 60 mg/l and Chloride more than 1000 mg/l produce positive interferences.
    TN = Total Nitrogen
14. ▲ N
    NH_4
    ▼ NH_3

1.1 Methods

2 9 0 Oxygen, active * with Tablet

0.1 – 10 mg/l O_2

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber.

5. Add **one DPD No. 4 tablet** straight from the foil to the water sample and crush the tablet using a clean stirring rod.

6. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.

7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.

   Wait for a **reaction period of 2 minutes**.

Countdown
2:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l active Oxygen.

1.1 Methods

Notes:

*** Active Oxygen is a synonym for a common disinfectant (based on "Oxygen") in Swimming Pool Treatment.**

1. When preparing the sample, the lost of Oxygen, e.g. by pipetting or shaking, must be avoided.
2. The analysis must take place immediately after taking the sample.

2 9 2 Oxygen, dissolved with Vacu-vials® K-7553 (see Notes)

10 – 800 µg/l O_2

Insert the adapter for 13 mm Ø round vials.

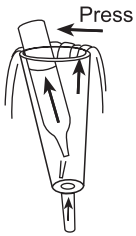
1. Place the blank in the sample chamber. The blank is part of the test kit.

prepare Zero
press ZERO

2. Press **ZERO** key.

3. Remove the blank from the sample chamber.

4. Water should flow through the special sample container for several minutes to remove any air bubbles sticking at the surface.



   The water must flow from the bottom to the top.

5. When the sample container is bubble-free press one Vacu-vial® into the lower edge of the sample container. The Vacu-vial® breaks at the neck and the vial fills automatically.

   A small volume of inert gas remains in the Vacu-vial®.

6. Remove the Vacu-vial® point downwards from the sample container immediately.

   As the contents of the vial has a higher density than water, it is important to remove the vial from the sample container within 5 seconds to prevent any loss of reagent.

7. The Vacu-vial® is closed with one finger (covered with a glove) to prevent entry of air. Invert the vial several times. Dry the outside of the vial.

8. Place the Vacu-vial® in the sample chamber.

Zero accepted
prepare Test
press TEST

9. Press **TEST** key.

   The result is shown in the display in µg/l Oxygen.

1.1 Methods

Notes:

1. This method is adapted from CHEMetrics.
2. Read the original test instruction and the MSDS (delivered with the test) before performing the test. MSDS also available at www.chemetrics.com.
3. Vacu-vials® is a registered trade mark of the company CHEMetrics, Inc. / Calverton, U.S.A.

1.1 Methods

3 0 0 Ozone with Tablet

0.02 – 2 mg/l O_3

Ozon	
>>	**with Cl**
	without Cl

The following selection is shown in the display:

>>	**with Cl**

for the determination of Ozone in the presence of Chlorine.

>>	**without Cl**

for the determination of Ozone in the absence of Chlorine.

Select the desired method with the arrow keys [▲] and [▼]. Confirm with [↵] key.

1.1 Methods

Notes:

1. Vial cleaning:
   As many household cleaners (e.g. dishwasher detergent) contain reducing substances, the subsequent determination of Ozone may show lower results. To avoid any measurement errors, only use glassware free of Chlorine demand.
   Preparation: Put all applicable glassware into Sodium hypochlorite solution (0.1 g/l) for one hour, then rinse all glassware thoroughly with deionised water.
2. Preparing the sample:
   When preparing the sample, the lost of Ozone, e.g. by pipetting or shaking, must be avoided. The analysis must take place immediately after taking the sample.
3. The DPD colour development is carried out at a pH value of 6.2 to 6.5. The reagent tablet therefore contains a buffer for the pH adjustment.
   Strong alkaline or acidic water samples must be adjusted between pH 6 and pH 7 before the tablet is added (use 0.5 mol/l Sulfuric acid resp. 1 mol/l Sodium hydroxide).
4. Turbidity (can lead to errors):
   The use of the DPD No. 1 tablet in samples with high Calcium ion contents* and/or high conductivity* can lead to turbidity of the sample and therefore incorrect measurements.
   ** it is not possible to give exact values, because the development of turbidity depends on the nature of the sample.*
5. Exceeding the measuring range:
   Concentrations above 6 mg/l Ozone can lead to results showing 0 mg/l. In this case, the water sample must be diluted with water free of Ozone. 10 ml of the diluted sample should be mixed with the reagent and the measurement repeated.
6. If ??? is displayed at the diffentiated test result see page 262.

   Oxidising agents such as Bromine, Chlorine etc. interfere as they react in the same way as Ozone.

1.1 Methods

3 0 0 Ozone, in the presence of Chlorine with Tablet

0.02 – 2 mg/l O_3

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.
4. Remove the vial from the sample chamber and **empty it, leaving a few drops remaining in the vial**.
5. Add **one DPD No. 1 tablet** and **one DPD No. 3 tablet** straight from the foil and crush the tablets using a clean stirring rod.
6. Add water sample to the 10 ml mark.
7. Close the vial tightly with the cap and swirl several times until the tablets are dissolved.
8. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare T1
press TEST

9. Press **TEST** key.
   Wait for a **reaction period of 2 minutes**.

Countdown
2:00

After the reaction period is finished the measurement starts automatically.

10. Remove the vial from the sample chamber, empty the vial, rinse vial and cap several times. Fill the vial with **a few drops of water sample**.
11. Add **one DPD No. 1 tablet** and **one DPD No. 3 tablet** straight from the foil and crush the tablets using a clean stirring rod.

12. **Fill a second clean vial with 10 ml of water sample.**

13. Add **one GLYCINE tablet** straight from the foil and crush the tablet using a clean stirring rod.

14. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.

15. **Transfer the contents of the second vial into the prepared vial.**

16. Close the vial tightly with the cap and swirl several times until the tablets are dissolved.

17. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

T1 accepted
prepare T2
press TEST

18. Press **TEST** key.
Wait for a **reaction period of 2 minutes**.

Countdown
2:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in:

***,** mg/l O_3**
***,** mg/l total Cl**

mg/l Ozone
mg/l total Chlorine

Notes:

See page 137.

3 0 0 Ozone, in absence of Chlorine with Tablet

0.02 – 2 mg/l O_3

prepare Zero
press ZERO

1. Fill a clean vial (24 mm Ø) **with 10 ml of water sample**, close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.
3. Press **ZERO** key.
4. Remove the vial from the sample chamber and **empty it, leaving a few drops remaining in the vial**.
5. Add **one DPD No. 1 tablet** and **one DPD No. 3 tablet** straight from the foil and crush the tablets using a clean stirring rod.
6. Add water sample to the 10 ml mark.
7. Close the vial tightly with the cap and swirl several times until the tablets are dissolved.
8. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

9. Press **TEST** key.
   Wait for a **reaction period of 2 minutes**.

Countdown
2:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in

mg/l Ozone.

Notes:
See page 137.

1.1 Methods

7 0 PHMB (Biguanide) with Tablet

2 – 60 mg/l PHMB

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber.

5. Add **one PHMB PHOTOMETER tablet** straight from the foil to the water sample and crush the tablet using a clean stirring rod.

6. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.

7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.

   The result is shown in the display in mg/l PHMB.

1.1 Methods

Notes:

1. Clean vials with the brush immediately after analysis.
2. Vials and stirring rods may turn blue after prolonged use. In this case clean vials and stirring rods with a laboratory detergent (see chapter 1.2.2 Cleaning of vials and accessories for analysis). Rinse vials and caps thoroughly with tap water and then with deionised water.
3. The test result is influenced by Hardness and Total Alkalinity.
   The calibration of this method was done using water with the following concentration:
   Ca-Hardness: 200 mg/l $CaCO_3$
   Total Alkalinity: 120 mg/l $CaCO_3$

1.1 Methods

3 2 0

Phosphate, ortho LR with Tablet

0.05 – 4 mg/l PO_4
Determination of ortho-Phosphate ions

3 2 1

Phosphate, ortho HR with Tablet

1 – 80 mg/l PO_4
Determination of ortho-Phosphate ions

3 2 3

Phosphate, ortho with Vario Powder Pack

0.06 – 2.5 mg/l PO_4
Determination of ortho-Phosphate ions

3 2 4

Phosphate, ortho with Vario Tube Test

0.06 – 5 mg/l PO_4
Determination of ortho-Phosphate ions

3 2 7

Phosphat 1, ortho with Vacu-vials®

5 – 40 mg/l PO_4
Determination of ortho-Phosphate ions

3 2 8

Phosphat 2, ortho with Vacu-vials®

0.05 – 5 mg/l PO_4
Determination of ortho-Phosphate ions

3 2 5

Phosphate, acid hydrolizable with Vario Tube Test

0.02 – 1.6 mg/l P
Determination of ortho-Phosphate ions + condensed, inorganic Phosphates

3 2 6

Phosphate, total with Vario Tube Test

0.02 – 1.1 mg/l P
Determination of ortho-Phosphate ions + condensed, inorganic Phosphates + organically combined Phosphates

1.1 Methods

General:
Ortho-Phosphate ions react with the reagent to form an intense blue colour (methods **320**, **323**, **324**, **325** and **326**).
Phosphate in organic and condensed inorganic forms (meta-, pyro- and polyphosphates) must be converted to ortho-Phosphate ions before analysis.
Pretreatment of the sample with acid and heat provides the conditions for hydrolysis of the condensed inorganic forms. Organically combined phosphates are converted to ortho-Phosphate ions by heating with acid and persulfate.
The amount of organically combined phosphates can be calculated:
mg/l Phosphate, organic = mg/l Phosphate, total – mg/l Phosphate, acid hydrolysable

In methods **321** and **327** the ortho-Phosphate ions react with the Vanadate-molybdate-reagent under acid conditions to form a yellow coloured product.

Notes – only for tube tests and tests with powder packs:
323, 324, 325, 326

1. Application: for water, wastewater and seawater.
2. Highly buffered samples or samples with extreme pH values should be adjusted between pH 2 and pH 10 before analysis (with 1 mol/l Hydrochloric acid or 1 mol/l Sodium hydroxide).
3. Interferences:
   Large amounts of turbidity may cause inconsistent results.

Interfering substance	**Interference level:**
Aluminium	greater than 200 mg/l
Arsenate	at any level
Chromium	greater than 100 mg/l
Copper	greater than 10 mg/l
Iron	greater than 100 mg/l
Nickel	greater than 300 mg/l
Silica (Silicium dioxide)	greater than 50 mg/l
Silicate	greater than 10 mg/l
Sulfide	at any level
Zinc	greater than 80 mg/l

Phosphate, ortho ≙ Phosphorus, reactive

1.1 Methods

3 2 0 Phosphate, ortho LR with Tablet

0.05 – 4 mg/l PO_4

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.

2. Place the vial in the sample chamber making sure that the marks ⧖ are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber.

5. Add **one PHOSPHATE No. 1 LR tablet** straight from the foil to the water sample and crush the tablet using a clean stirring rod.

6. Add **one PHOSPHATE No. 2 LR tablet** straight from the foil to the same water sample and crush the tablet using a clean stirring rod.

7. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.

8. Place the vial in the sample chamber making sure that the marks ⧖ are aligned.

Zero accepted
prepare Test
press TEST

9. Press **TEST** key.
Wait for a **reaction period of 10 minutes**.

Countdown
10:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l ortho-Phosphate.

1.1 Methods

Notes:

1. Only ortho-Phosphate ions react.
2. The tablets must be added in the correct sequence.
3. The test sample should have a pH-Value between 6 and 7.
4. Interferences:
   Higher concentrations of Cu, Ni, Cr (III), V (V) and W (VI) interfere due to their colour.
   Silicates do not interfere (masked by Citric acid in the tablets).
5. see also page 145
6. Conversion:
   mg/l P = mg/l PO_4 x 0.33
   mg/l P_2O_5= mg/l PO_4 x 0.75
7. ▲ PO_4
   P
   ▼ P_2O_5

1.1 Methods

3 2 1 Phosphate HR, ortho with Tablet

1 – 80 mg/l PO_4

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.
4. Remove the vial from the sample chamber.
5. Add **one PHOSPHATE HR P1 tablet** straight from the foil to the water sample and crush the tablet using a clean stirring rod.
6. Add **one PHOSPHATE HR P2 tablet** straight from the foil to the same water sample and crush the tablet using a clean stirring rod.
7. Close the vial tightly with the cap and swirl several times until the tablets are dissolved.
8. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

9. Press **TEST** key.

   Wait for a **reaction period of 10 minutes**.

Countdown
10:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l ortho-Phosphate.

1.1 Methods

Notes:

1. For samples under 5 mg/l PO_4 it is reccommended to analyse the water sample with method 320 "Posphate LR, ortho with Tablet".
2. Only ortho-Phosphate ions react.
3. see also page 145
4. Conversions:
   mg/l P = mg/l PO_4 x 0.33
   mg/l P_2O_5 = mg/l PO_4 x 0.75
5. ▲ PO_4
   P
   ▼ P_2O_5

1.1 Methods

3 2 3

Phosphate, ortho with Vario Powder Pack

0.06 – 2.5 mg/l PO_4

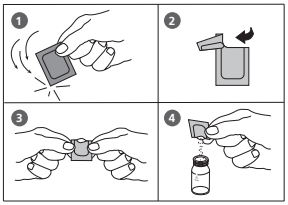
1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber.



5. Add the contents of **one VARIO Phosphate Rgt. F10 Powder Pack** straight from the foil to the water sample.

6. Close the vial tightly with the cap and swirl several times to mix the contents (approx. 10–15 sec., Note 1).

7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.

   Wait for a **reaction period of 2 minutes**.

Countdown
2:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l ortho-Phosphate.

1.1 Methods

Notes:

1. The reagent does not dissolve completely.
2. see also page 145
3. Conversions:
   mg/l P = mg/l PO_4 x 0.33
   mg/l P_2O_5 = mg/l PO_4 x 0.75
4. ▲ PO_4
   P
   ▼ P_2O_5

1.1 Methods

3 2 4

Phosphate, ortho with Vario Tube Test

0.06 – 5 mg/l PO_4

Ø 16 mm

Insert the adapter for 16 mm Ø vials.

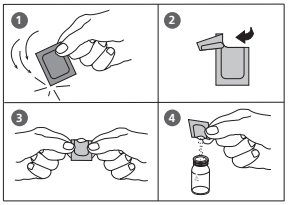
1. Open one white capped reaction **tube PO_4-P Dilution** and add **5 ml of water sample.**

2. Place the vial in the sample chamber making sure that the ⍙ marks are aligned.
Place the cover on the adapter.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber.



5. Add the contents of **one Vario Phosphate Rgt. F10 Powder Pack** straight from the foil to the water sample (Note 1).

6. Close the vial tightly with the cap and swirl several times to mix the contents (approx. 10–15 sec., Note 2).

7. Place the vial in the sample chamber making sure that the ⍙ marks are aligned.
Place the cover on the adapter.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.

Wait for a **reaction period of 2 minutes**.

Countdown
2:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l ortho-Phosphate.

1.1 Methods

Notes:

1. Use a funnel to add the reagent.
2. The reagent does not dissolve completely.
3. see also page 145
4. Conversions:
   mg/l P = mg/l PO_4 x 0.33
   mg/l P_2O_5 = mg/l PO_4 x 0.75
5. ▲ PO_4
   P
   ▼ P_2O_5

1.1 Methods

3 2 7

Phosphate 1, ortho with Vacu-vials® K-8503 (see Notes)

5 – 40 mg/l PO_4

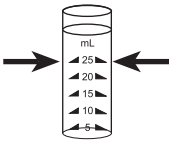
Insert the adapter for 13 mm Ø vials.

1. Place the blank in the sample chamber. The blank is part of the test kit.

prepare Zero
press ZERO

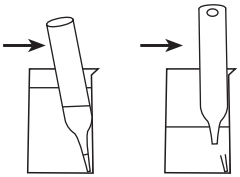
2. Press **ZERO** key.

3. Remove the blank from the sample chamber.



4. Fill the sample container to the 25 ml mark with the water sample.

5. Place one Vacu-vial® in the sample container. Snap the tip by pressing the vial against the side of the sample container.



The Vacu-vial® breaks at the neck and the vial fills automatically.

A small volume of inert gas remains in the Vacu-vial®.

6. Mix the contents of the Vacu-vial® by inverting it several times, allowing the bubble to move from one end to the other. Dry the outside of the vial.

7. Place the Vacu-vial® in the sample chamber.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.
Wait for a **reaction period of 5 minutes**.

Countdown
5:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l ortho-Phosphate.

1.1 Methods

Notes:

1. This method is adapted from CHEMetrics.
2. Read the original test instruction and the MSDS (delivered with the test) before performing the test. MSDS also available at www.chemetrics.com.
3. Vacu-vials® is a registered trade mark of the company CHEMetrics, Inc. / Calverton, U.S.A.
4. Only ortho-Phosphate ions react.
5. Sulfide, Thiosulfate and Thiocyanate cause low test results.
6. ▲ PO_4
   P
   ▼ P_2O_5

3 2 8 Phosphate 2, ortho with Vacu-vials® K-8513 (see Notes)

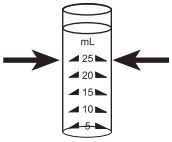
0.05 – 5 mg/l PO_4

Insert the adapter for 13 mm Ø vials.

1. Place the blank in the sample chamber. The blank is part of the test kit.

prepare Zero
press ZERO

2. Press **ZERO** key.



3. Remove the blank from the sample chamber.

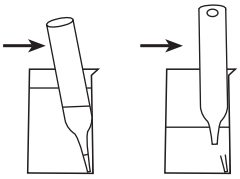
4. Fill the sample container to the 25 ml mark with the water sample.

5. Fill the sample container with drops of the same size by holding the bottle vertically and squeeze slowly:

   2 drops A-8500 Activator Solution

6. Close the sample container with the cap tightly and swirl several times to mix the contents.

7. Place one Vacu-vial® in the sample container. Snap the tip by pressing the vial against the side of the sample container. The Vacu-vial® breaks at the neck and the vial fills automatically. A small volume of inert gas remains in the Vacu-vial®.



8. Mix the contents of the Vacu-vial® by inverting it several times, allowing the bubble to move from one end to the other. Dry the outside of the vial.

9. Place the Vacu-vial® in the sample chamber.

Zero accepted
prepare Test
press TEST

10. Press **TEST** key.

    Wait for a **reaction period of 3 minutes.**

Countdown
3:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l ortho-Phosphate.

1.1 Methods

Notes:

1. This method is adapted from CHEMetrics.
2. Read the original test instruction and the MSDS (delivered with the test) before performing the test. MSDS also available at www.chemetrics.com.
3. Vacu-vials® is a registered trade mark of the company CHEMetrics, Inc. / Calverton, U.S.A.
4. Only ortho-Phosphate ions react.
5. Sulfide, Thiosulfate and Thiocyanate cause low test results.
6. ▲ PO_4
   P
   ▼ P_2O_5

1.1 Methods

3 2 5 Phosphate, acid hydrolyzable with Vario Tube Test

0.02 – 1.6 mg/l P (≙ 0.06 – 5 mg/l PO_4)

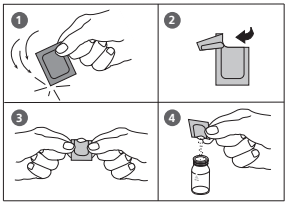
Ø 16 mm

Insert the adapter for 16 mm Ø vials.

1. Open one white capped **digestion tube PO4-P Acid reagent** and add **5 ml of water sample**.
2. Close the vial tightly with the cap. Invert the vial gently several times to mix the contents.
3. Heat the vials for **30 minutes** in the preheated reactor at a temperature **of 100°C**.
4. After 30 minutes remove the vial from the reactor. **(CAUTION: The vials are hot!)**
   Allow the vials to cool to room temperature.
5. Open the cooled digestion vial and add **2 ml 1.00 N Sodium hydroxide solution** to the vial.
6. Close the vial with the cap and invert the vial gently several times to mix the contents.
7. Place the vial in the sample chamber making sure that the ⍋ marks are aligned. Place the cover on the adapter.

prepare Zero
press ZERO

8. Press **ZERO** key.
9. Remove the vial from the sample chamber.



10. Add the contents of **one Vario Phosphate Rgt. F10 Powder Pack** straight from the foil to the vial (Note 2).
11. Close the vial tightly with the cap and swirl several times to mix the contents (approx. 10–15 sec., Note 3).
12. Place the vial in the sample chamber making sure that the ⍋ marks are aligned. Place the cover on the adapter.

Zero accepted
prepare Test
press TEST

13. Press **TEST** key.
    Wait for a **reaction period of 2 minutes**.

Countdown
2:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l acid hydrolyzable Phosphate.

1.1 Methods

Notes:

1. Appropriate safety precautions and a good lab technique should be used during the whole procedure.
2. Use a funnel to add the reagent.
3. The reagent does not dissolve completely.
4. see also page 145
5. Conversions:
   mg/l PO_4 = mg/l P x 3.07
   mg/l P_2O_5 = mg/l P x 2.29
6. ▲ PO_4
   P
   ▼ P_2O_5

3 2 6 Phosphate, total with Vario Tube Test

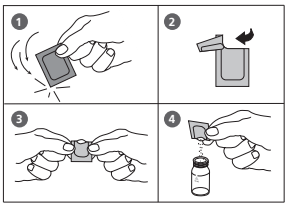
0.02 – 1.1 mg/l P (≙ 0.06 – 3.5 mg/l PO_4)

Ø 16 mm

Insert the adapter for 16 mm Ø vials.

1. Open one white capped **digestion tube PO4-P Acid reagent** and add **5 ml of water sample**.
2. Add the contents of **one Vario Potassium Persulfate F10 Powder Pack** straight from the foil to the vial (Note 2).
3. Close the vial tightly with the cap. Invert the vial several times to mix the contents.
4. Heat the vials for **30 minutes** in the preheated reactor at a temperature **of 100°C**.
5. After 30 minutes remove the vial from the reactor. **(CAUTION: The vials are hot!)**
   Allow the vials to cool to room temperature.
6. Open the cooled digestion vial and add **2 ml 1.54 N Sodium hydroxide solution** to the vial.
7. Close the vial with the cap and invert the vial gently several times to mix the contents.
8. Place the vial in the sample chamber making sure that the ⍋ marks are aligned. Place the cover on the adapter.

prepare Zero
press ZERO



9. Press **ZERO** key.
10. Remove the vial from the sample chamber.
11. Add the contents of **one Vario Phosphate Rgt. F10 Powder Pack** straight from the foil to the vial (Note 2).
12. Close the vial tightly with the cap and swirl several times to mix the contents (approx. 10–15 sec., Note 3).
13. Place the vial in the sample chamber making sure that the ⍋ marks are aligned. Place the cover on the adapter.

Zero accepted
prepare Test
press TEST

14. Press **TEST** key.
    Wait for a **reaction period of 2 minutes**.

Countdown
2:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l total Phosphate.

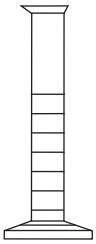
1.1 Methods

Notes:

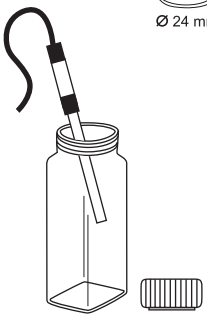
1. Appropriate safety precautions and a good lab technique should be used during the whole procedure.
2. Use a funnel to add the reagent.
3. The reagent does not dissolve completely.
4. see also page 145
5. Conversions:
   mg/l PO_4 = mg/l P x 3.07
   mg/l P_2O_5 = mg/l P x 2.29
6. ▲ P
   PO_4
   ▼ P_2O_5

3 1 6 Phosphonates Persulfate UV oxidation method with Vario Powder Pack

0 – 125 mg/l (see Table 1)



Ø 24 mm



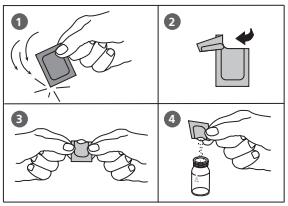
1. Choose the appropriate sample volume from table 1 (see following pages).
2. Pipette the chosen sample volume into a clean 50 ml graduated cylinder. If necessary fill up with deionised water to the 50 ml mark and mix well.
3. Fill a clean vial (24 mm Ø) with **10 ml of the prepared water sample** (this is the blank).
4. Transfer **25 ml of the prepared water sample** into the digestion vial.
5. Add the contents of **one Vario Potassium Persulfate F10 Powder Pack** straight from the foil to the digestion vial.
6. Close the digestion vial tightly with the cap and swirl until the reagent is dissolved completely.
7. Insert the UV lamp into the digestion vial (Note 3, 4, 5). **CAUTION: Wear UV safety goggles!**

Countdown 1
10:00
start: ↲

8. Switch the UV lamp on and wait for a **reaction period of 10 minutes**.
9. After the reaction period is finished switch the UV lamp off and remove the lamp from the vial.

Ø 24 mm

10. Fill a second vial (24 mm Ø) with **10 ml of the digested sample** (this is the sample).



11. Add the contents of **one Vario Phosphate Rgt. F10 Powder Pack** straight from the foil into each vial (blank and sample).
12. Close the vials tightly with the cap and swirl gently several times (30 sec.). (Note 6)

1.1 Methods

13. Place the vial (the blank) in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

14. Press **ZERO** key.

Countdown
2:00

Wait for a **reaction period of 2 minutes** (Note 7).

After the reaction period is finished the measurement starts automatically.

15. Remove the vial from the sample chamber.

16. Place the vial (the sample) in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

17. Press **TEST** key.

The result is shown in the display in mg/L PO_4^{3-}.

To calculate the actual phosphonate concentration multiply the reading with the corresponding dilution factor from table 1.

To calculate the active phosphonate concentration multiply the actual phosphonate concentration using the appropriate factor from table 2.

Notes:

1. Rinse all glassware with 1:1 Hydrochloric acid first and then rinse with deionised water. Do not use detergents with phosphates.
2. During UV digestion Phosphonates are converted to Ortho-Phosphates. This step is normally completed in 10 minutes. High organic loaded samples or a weak lamp can cause incomplete phosphate conversion.
3. UV lamp available on request.
4. While the UV lamp is on UV safety goggles must be worn.
5. For handling of the UV lamp see manufacturer's manual. Do not touch the surface of the UV lamp. Fingerprints will etch the glass. Wipe the UV lamp with a soft and clean tissue between measurements.
6. The reagent does not dissolve completely.
7. The given reaction time of 2 minutes refers to a water sample temperature of more than 15°C. At a sample temperature lower than 15 °C a reaction time of 4 minutes is required.

Tables:

see next page

1.1 Methods

Table 1:

Expected range (mg/L Phosphonate)	Sample volume in ml	Factor
0 – 2.5	50	0.1
0 – 5.0	25	0.2
0 – 12.5	10	0.5
0 – 25	5	1.0
0 – 125	1	5.0

Table 2:

Phosphonate type	Conversion factor for active phosphonate
PBTC	2.840
NTP	1.050
HEDPA	1.085
EDTMPA	1.148
HMDTMPA	1.295
DETPMPA	1.207
HPA	1.490

1.1 Methods

Interference levels decrease with increasing sample volume.
Example: Iron interferes above 200 mg/L if a sample volume of 5 ml is used.
At a sample volume of 10 ml the interference level decreases to 100 mg/L.

Table 3:

Interfering substances	Interference level using 5 ml of sample
Aluminium	100 mg/l
Arsenate	interferes at all concentrations
Benzotriazole	10 mg/l
Bicarbonate	1000 mg/l
Bromide	100 mg/l
Calcium	5000 mg/l
CDTA	100 mg/l
Chloride	5000 mg/l
Chromate	100 mg/l
Copper	100 mg/l
Cyanide	100 mg/l; increase the UV digestion to 30 minutes
Diethanoldithiocarbamate	50 mg/l
EDTA	100 mg/l
Iron	200 mg/l
Nitrate	200 mg/l
NTA	250 mg/l
Orthophosphate	15 mg/l
Phosphite and organophosphorus compounds	reacts quantitatively; Meta- and Polyphosphates do not interfere
Silica	500 mg/l
Silicate	100 mg/l
Sulfate	2000 mg/l
Sulfide	interferes at all concentrations
Sulfite	100 mg/l
Thiourea	10 mg/l
highly buffered samples or extreme sample pH	may exceed the buffering capacity of the reagents and require sample pretreatment

1.1 Methods

3 2 9 pH value LR 5.2 – 6.8 with Tablet

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber.

5. Add **one BROMOCRESOLPURPLE PHOTOMETER tablet** straight from the foil to the water sample and crush the tablet using a clean stirring rod.

6. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.

7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.

   The result is shown in the display as pH-value.

1.1 Methods

Notes:

1. For photometric determination of pH values only use BROMOCRESOLPURPLE tablets in black printed foil pack and marked with PHOTOMETER.
2. pH values below 5.2 and above 6.8 can produce results inside the measuring range. A plausibility test (pH-meter) is recommended.
3. The accuracy of the colorimetric determination of pH values depends on various boundary conditions (buffer capacity of the sample, salt contents etc.).
4. Salt error
   Correction of test results (average values) for samples with salt contents of:

Indicator	Salt contents		
Bromcresolpurple	1 molar – 0.26	2 molar – 0.33	3 molar – 0.31

The values of Parson and Douglas (1926) are based on the use of Clark and Lubs buffers.

1 Mol NaCl = 58.4 g/l = 5.8 %

1.1 Methods

3 3 0 pH value 6.5 – 8.4 with Tablet

Ø 24 mm

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.
4. Remove the vial from the sample chamber.
5. Add **one PHENOL RED PHOTOMETER tablet** straight from the foil to the water sample and crush the tablet using a clean stirring rod.
6. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.
7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.

   The result is shown in the display as pH-value.

1.1 Methods

Notes:

1. For photometric determination of pH values only use PHENOL RED tablets in black printed foil pack and marked with PHOTOMETER.
2. Water samples with low values of Alkalinity-m (below 35 mg/l $CaCO_3$) may give wrong pH readings.
3. pH values below 6.5 and above 8.4 can produce results inside the measuring range. A plausibility test (pH-meter) is recommended.
4. The accuracy of the colorimetric determination of pH values depends on various boundary conditions (buffer capacity of the sample, salt contents etc.).
5. Salt error
   Correction of test results (average values) for samples with salt contents of:

Indicator	Salt contents		
Phenol red	1 molar – 0.21	2 molar – 0.26	3 molar – 0.29

The values of Parson and Douglas (1926) are based on the use of Clark and Lubs buffers. 1 Mol NaCl = 58.4 g/l = 5.8 %

1.1 Methods

(3) (3) (1) pH value 6.5 – 8.4 with Liquid Reagent

Ø 24 mm

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.
4. Remove the vial from the sample chamber.
5. Fill the vial with drops of the same size by holding the bottle vertically and squeeze slowly:

   6 drops of PHENOL RED solution

6. Close the vial tightly with the cap and swirl several times to mix the contents.
7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare TEST
press Test

8. Press **TEST** key.

   The result is shown in the display as pH-value.

1.1 Methods

Notes:

1. When testing chlorinated water the residual chlorine contents can influence the colour reaction of the liquid reagent. This can be avoided (without interfering with the pH measurement) by adding a small crystal of Sodiumthiosulfate ($Na_2S_2O_3 \cdot 5\ H_2O$) to the sample before adding the PHENOL RED solution. PHENOL RED tablets already contain Thiosulfate.
2. Due to differing drop sizes results can show a discrepancy in accuracy by comparison with tablets. This can be minimised by using a pipette (0.18 ml PHENOL RED solution is equivalent to 6 drops).
3. After use replace the bottle cap securely.
4. **Store the reagent in a cool, dry place ideally between 6°C and 10°C.**

1.1 Methods

3 3 2 pH value HR 8.0 – 9.6 with Tablet

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.
4. Remove the vial from the sample chamber.
5. Add **one THYMOLBLUE PHOTOMETER tablet** straight from the foil to the water sample and crush the tablet using a clean stirring rod.
6. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.
7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare TEST
press Test

8. Press **TEST** key.

The result is shown in the display as pH-value.

1.1 Methods

Notes:

1. For photometric determination of pH values only use THYMOLBLUE tablets in black printed foil pack and marked with PHOTOMETER.
2. pH values below 8.0 and above 9.6 can produce results inside the measuring range. A plausibility test (pH-meter) is recommended.
3. The accuracy of the colorimetric determination of pH values depends on various boundary conditions (buffer capacity of the sample, salt contents etc.).
4. Salt error

   Correction of test results (average values) for samples with salt contents of:

Indicator	Salt contents		
Thymolblue	1 molar – 0.22	2 molar – 0.29	3 molar – 0.34

   The values of Parson and Douglas (1926) are based on the use of Clark and Lubs buffers. 1 Mol NaCl = 58.4 g/l = 5.8 %

1.1 Methods

3 4 0 Potassium with Tablet

0.7 – 12 mg/l K

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber.

5. Add **one Potassium T tablet** straight from the foil to the water sample and crush the tablet using a clean stirring rod.

6. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.

7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.

The result is shown in the display in mg/l Potassium.

1.1 Methods

Notes:

1. If Potassium is present a cloudy solution will appear.
   Single particles are not necessarily caused by Potassium.

1.1 Methods

3 5 0 Silica/Silicon dioxide with Tablet

0.05 – 4 mg/l SiO_2

prepare Zero
press ZERO

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.
3. Press **ZERO** key.
4. Remove the vial from the sample chamber.
5. Add **one SILICA No. 1 tablet** straight from the foil to the water sample and crush the tablet using a clean stirring rod.
6. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.

Countdown
5:00
start: ↵

7. Press [↵] key.
   Wait for a **reaction period of 5 minutes**.

   After the reaction period is finished proceed as follows:
8. Add **one SILICA PR tablet** straight from the foil to the same water sample and crush the tablet using a clean stirring rod.
9. Add **one SILICA No. 2 tablet** straight from the foil to the same water sample and crush the tablet using a clean stirring rod.
10. Close the vial tightly with the cap and swirl several times until the tablets are dissolved.
11. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

12. Press **TEST** key.
    Wait for a **reaction period of 1 minute**.

Countdown
1:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l Silica.

1.1 Methods

Notes:

1. The tablets must be added in the correct sequence.
2. Phosphate ions do not interfere under the given reaction conditions.
3. If Phosphate is known to be absent, the addition of the SILICA PR tablet may be omitted.
4. Conversion:
   mg/l Si = mg/l SiO_2 x 0.47
5. ▲ SiO_2
   ▼ Si

1.1 Methods

3 5 1

Silica LR / Silicon dioxide LR with Vario Powder Pack and Liquid Reagent

0.1 – 1.6 mg/l SiO_2

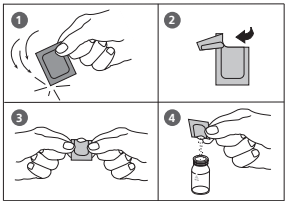
Use two clean vials (24 mm Ø) and mark one as blank for zeroing.

1. Fill each vial with **10 ml of water sample**.
2. Add **0.5 ml Vario Molybdate 3 reagent solution** into each vial.
3. Close the vials tightly with the caps and swirl the vials several times to mix the contents (Note 1).

Countdown
4:00
start: ↵

4. Press **[↵]** key.

   Wait for a **reaction period of 4 minutes** (Note 2).



After the reaction period is finished proceed as follows:

5. Add the contents of **one Vario Silica Citric Acid F10 Powder Pack** straight from the foil into each vial.
6. Close the vials tightly with the caps and swirl the vials several times to mix the contents.

Countdown
1:00
start: ↵

7. Press **[↵]** key.
   Wait for a **reaction period of 1 minute** (Note 3).

After the reaction period is finished proceed as follows:

8. Place the vial (the blank) in the sample chamber making sure that the ⧖ marks are aligned.
9. Add the contents of **one Vario Silica LR Amino Acid F F10 Powder Pack** straight from the foil into the vial (the sample).
10. Close the vial tightly with the cap and swirl several times to mix the contents.

1.1 Methods

prepare Zero
press ZERO

11. Press **ZERO** key (blank is already placed in the sample chamber - see point 8).

Countdown
2:00

Wait for a **reaction period of 2 minutes**.

After the reaction period is finished the zero-reading starts automatically.

12. Remove the vial from the sample chamber.

13. Place the vial (the sample) in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

14. Press **TEST** key.

The result is shown in the display in mg/l Silica.

Notes:

1. Close the vials with the cap immediately after adding the Vario Molybdate 3 reagent solution, otherwise low readings may result.
2. The given reaction time of 4 minutes refers to a water sample temperature of 20°C.
   At 30°C a reaction time of 2 minutes, at 10°C a reaction time of 8 minutes are required.
3. The given reaction time of 1 minute refers to a water sample temperature of 20°C.
   At 30°C a reaction time of 30 seconds, at 10°C a reaction time of 2 minutes are required.
4. Interferences:

Substance	Interference
Iron	large amounts interfere
Phosphate	does not interfere at concentrations less than 50 mg/l PO_4 at 60 mg/l PO_4 the interference is approx. – 2% at 75 mg/l PO_4 the interference is approx. – 11%
Sulfide	interferes at all levels

   Occasionally water samples contain forms of silica which reacts very slowly with Molybdate. The nature of these forms is not known.
   A pre-treatment with Sodium hydrogencarbonate and then with Sulfuric Acid will make these forms reactive to Molybdate (pre-treatment is given in "Standard Methods for the Examination of Water and Wastewater" under " Silica Digestion with Sodium Bicarbonate").

5. ▲ SiO_2
   ▼ Si

1.1 Methods

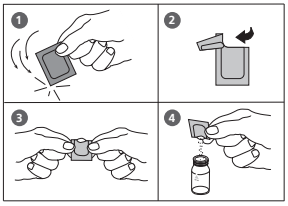
(3) (5) (2)

Silica HR / Silicon dioxide HR with Vario Powder Pack

1 – 90 mg/l SiO_2

Ø 24 mm

prepare Zero
press ZERO



1. Fill a clean vial (24 mm Ø) with **10 ml of water sample** (Note 1), close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.
3. Press **ZERO** key.
4. Remove the vial from the sample chamber.
5. Add the contents of **one Vario Silica HR Molybdate F10 Powder Pack** straight from the foil to the water sample.
6. Close the vial tightly with the cap and swirl several times to mix the contents.
7. Add the contents of **one Vario Silica HR Acid Rgt. F10 Powder Pack** straight from the foil to the same water sample (Note 2).
8. Close the vial with the caps tightly and swirl the vials several times to mix the contents.

Countdown
10:00
start: ↵

9. Press **[↵]** key.
   Wait for a **reaction period of 10 minutes**.

   After the reaction period is finished proceed as follows:
10. Add the contents of **one Vario Silica Citric Acid F10 Powder Pack** straight from the foil to the water sample (Note 3).
11. Close the vial tightly with the cap and swirl several times to mix the contents.
12. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

13. Press **TEST** key.
    Wait for a **reaction period of 2 minutes**.

Countdown
2:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l Silica.

1.1 Methods

Notes:

1. Temperature of the sample should be 15 °C – 25 °C.
2. If Silica or Phosphate is present a yellow colour is developed.
3. In this step any yellow colour due to Phosphate is removed.
4. Interferences:

Substance	**Interference**
Iron	large amounts interfere
Phosphate	does not interfere at concentrations less than 50 mg/l PO_4 at 60 mg/l PO_4 the interference is approx. – 2 % at 75 mg/l PO_4 the interference is approx. – 11 %
Sulfide	interferes at all levels

Occasionally water samples contain forms of silica which reacts very slowly with Molybdate. The nature of these forms is not known.

A pre-treatment with Sodium hydrogencarbonate and then with Sulfuric Acid will make these forms reactive to Molybdate (pre-treatment is given in "Standard Methods for the Examination of Water and Wastewater" under "Silica Digestion with Sodium Bicarbonate").

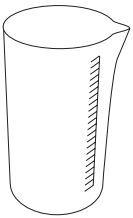
5. ▲ SiO_2
   ▼ Si

1.1 Methods

2 1 2

Sodium hypochlorite (Soda bleaching lye) with Tablet

0.2 – 16 % w/w NaOCl



Preparation:

1. Fill a 5 ml plastic syringe with the test solution, ensuring that all air bubbles are expelled. Transfer the 5 ml test solution slowly into a 100 ml beaker and dilute to the 100 ml mark with chlorine-free water. Mix thoroughly.
2. Fill a 5 ml plastic syringe with the diluted test solution (step 1) to the 1 ml mark, ensuring that all air bubbles are expelled. Transfer the 1 ml test solution slowly into a 100 ml beaker and dilute to the 100 ml mark with chlorine-free water. Mix thoroughly.

Performing test procedure:

Ø 24 mm

1. Fill a clean vial (24 mm Ø) with **10 ml of the prepared water sample**, close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.
4. Remove the vial from the sample chamber.
5. Add **one CHLORINE HR (KI) tablet** straight from the foil to the water sample and crush the tablet using a clean stirring rod.
6. Add **one ACIDIFYING GP tablet** straight from the foil to the same water sample and crush the tablet using a clean stirring rod.
7. Close the vial tightly with the cap and swirl several times until the tablets are dissolved.

8. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

9. Press **TEST** key.

The result is shown in the display in % w/w as available chlorine present in the original sample of Sodium hypochlorite.

Notes:

1. Please pay attention when handling sodium hypochlorite. The material has a very strong alkalinity and can cause corrosion. Contact with eyes, skin and clothes etc. has to be avoided. Refer to the detailed information the producer has supplied with the product.
2. The tablets must be added in the correct sequence.
3. This method provides a fast and simple test. The test can be performed on site but the result will not be as precise as a laboratory method.
4. By strictly following the test procedure, an accuracy of +/- 1 weight % can be achieved.

1.1 Methods

3 5 5 Sulfate with Tablet

5 – 100 mg/l SO_4

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample,** close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.
4. Remove the vial from the sample chamber.
5. Add **one SULFATE T tablet** straight from the foil to the water sample and crush the tablet using a clean stirring rod.
6. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.
7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.

The result is shown in the display in mg/l Sulfate.

1.1 Methods

Note:

1. If Sulfate is present a cloudy solution will appear.

1.1 Methods

3 6 0 Sulfate with Vario Powder Pack

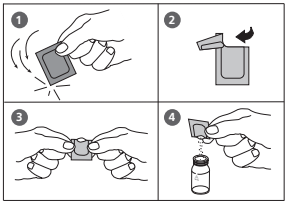
5 – 100 mg/l SO_4

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.



4. Remove the vial from the sample chamber.

5. Add the contents of **one Vario Sulpha 4 / F10 Powder Pack** straight from the foil to the water sample.

6. Close the vial tightly with the cap and swirl several times to mix the contents.

7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.
Wait for a **reaction period of 5 minutes.**

Countdown
5:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l Sulfate.

1.1 Methods

Note:

1. If Sulfate ions are present a cloudy solution will appear.

1.1 Methods

3 6 5 Sulfide with Tablet

0.04 – 0.5 mg/l S

Ø 24 mm

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample,** close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber.

5. Add **one SULFIDE No. 1 tablet** to the water sample and crush the tablet using a clean stirring rod and dissolve the tablet.

6. Add **one SULFIDE No. 2 tablet** to the same water sample and crush the tablet using a clean stirring rod.

7. Close the vial tightly with the cap and swirl several times until the tablets are dissolved.

8. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

9. Press **TEST** key.

   Wait for a **reaction period of 10 minutes.**

Countdown
10:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l Sulfide.

1.1 Methods

Notes:

1. The tablets must be added in the correct sequence.
2. Chlorine and other oxidizing agents which react with DPD do not interfere with the test.
3. To avoid loss of Sulfide collect the sample carefully with a minimum of aeration. It is essential to test the sample immediately after collection.
4. The sample temperature should be 20°C. A different temperature can lead to higher or lower results.
5. Conversion:
   H_2S = mg/l S x 1.06
6. ▲ S
   ▼ H_2S

1.1 Methods

3 7 0 Sulfite with Tablet

0.1 – 5 mg/l SO_3


Ø 24 mm

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.
4. Remove the vial from the sample chamber.
5. Add **one SULFITE LR tablet** straight from the foil to the water sample and crush the tablet using a clean stirring rod.
6. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.
7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.

   Wait for a **reaction period of 5 minutes**.

Countdown
5:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l Sulfite.

1.1 Methods

Notes:

1. ▲ SO_3
   ▼ Na_2SO_3

1.1 Methods

3 8 4 Suspended Solids

0 – 750 mg/l TSS


Ø 24 mm

Sample preparation:

Blend approx. 500 ml of the water sample in a blender at high speed for 2 minutes.

1. Fill a clean vial (24 mm Ø) with **10 ml of deionised water,** close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber and empty the vial completely.

5. Stir the blended water sample. Immediately rinse the vial with the water sample and fill with **10 ml water sample**.

6. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

7. Press **TEST** key.

   The result is shown in the display in mg/l TSS (Total Suspended Solids).

1.1 Methods

Note:

1. The photometric determination of Suspended Solids is based on a gravimetric method. In a lab this is usually done by evaporation of the filter residue of a filtrated water sample in an oven at 103°C – 105°C and weighing of the dried residue.
2. When higher accuracy is required perform a gravimetric determination of a water sample. The result can be used to calibrate the photometer with the same water sample.
3. The estimated detection limit is 20 mg/L TSS.
4. Collect water samples in clean plastic or glass bottles and analyse the water sample as soon as possible. It is possible to store the sample at 4°C for 7 days. Before measurement warm up the sample to the temperature at collection time.
5. Interferences:
   - Air bubbles interfere and can be removed by swirling the vial gently.
   - Colour interferes if light is absorbed at 660 nm.

1.1 Methods

3 8 6 Turbidity

0 – 1000 FAU

Ø 24 mm

1. Fill a clean vial (24 mm Ø) with **10 ml of deionised water,** close tightly with the cap.

2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.

4. Remove the vial from the sample chamber and empty the vial completely.

5. Stir the water sample. Immediately rinse the vial with the water sample and fill with **10 ml water sample**.

6. Close the vial tightly with the cap and swirl gently several times.

7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

8. Press **TEST** key.

The result is shown in the display in FAU.

1.1 Methods

Note:

1. This test uses an attenuated radiation method for the reading of FAU (Formazin Attenuation Units). The results can not be used for USEPA reporting purposes, but may be used for routine measurements. The attenuated radiation method is different from the Nephelometric method.
2. The estimated detection limit is 20 FAU.
3. Collect water samples in clean plastic or glass bottles and analyse the water sample as soon as possible. It is possible to store the sample at 4°C for 48 hours. Before measurement warm up the sample to the temperature at collection time. Temperature differences between measurement and sample collection can effect the turbidity of the sample.
4. Colour interferes if light is absorbed at 530 nm. For strong coloured water samples a filtrated portion of the sample can be used for zeroing instead of the deionised water.
5. Air bubbles interfere and can be removed using an ultrasonic bath.

1.1 Methods

3 9 0 Urea with Tablet and Liquid Reagent

0.1 – 2.5 mg/l $(NH_2)_2CO$ / mg/l Urea

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**, close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

3. Press **ZERO** key.
4. Remove the vial from the sample chamber.
5. Add **2 drops of Urea Reagent 1** to the water sample (Note 9).
6. Close the vial tightly with the cap and swirl several times to mix the contents.
7. Add **1 drop of Urea Reagent 2** (Urease) to the same water sample (Note 9).
8. Close the vial tightly with the cap and swirl several times to mix the contents.

Countdown
5:00
start: ↵

9. Press **[↵]** key.
   Wait for a **reaction period of 5 minutes**.

   After the reaction period is finished proceed as follows:

10. Add **one AMMONIA No. 1 tablet** straight from the foil to the prepared water sample and mix to dissolve with a clean stirring rod.
11. Add **one AMMONIA No. 2 tablet** straight from the foil to the same water sample and mix to dissolve with a clean stirring rod.

12. Close the vial tightly with the cap and swirl several times until the tablets are dissolved.

13. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
prepare Test
press TEST

14. Press **TEST** key.
Wait for a **reaction period of 10 minutes**.

Countdown
10:00

After the reaction period is finished the measurement starts automatically.

The result is shown in the display in mg/l Urea.

Notes:

1. The sample temperature should be between 20 °C and 30 °C.
2. Carry out the test at the latest one hour after sample taking.
3. Concentrations above 2 mg/l Urea can produce results inside the measuring range. In this case the water sample should be diluted with Urea free water and remeasured.
4. The tablets must be added in the correct sequence.
5. The AMMONIA No. 1 tablet will only dissolve completely after the AMMONIA No. 2 tablet has been added.
6. **Do not store reagent 1 (Urease) below 10°C; granulation is possible. Store reagent 2 (Urease) in the refrigerator at a temperature of 4°C to 8°C.**
7. Ammonia and chloramines are also measured during urea measurement.
8. Before analysing seawater samples, a measuring spoon of Ammonia Conditioning Powder must be added to the sample and swirled to dissolve before AMMONIA No. 1 tablet is added.
9. Fill the vial with drops of the same size by holding the bottle vertically and squeeze slowly.

1.1 Methods

4 0 0 Zinc with Tablet

0.02 – 1 mg/l Zn

1. Fill a clean vial (24 mm Ø) with **10 ml of water sample**.
2. Add **one COPPER / ZINC LR tablet** straight from the foil to the water sample, crush the tablet using a clean stirring rod.
3. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.
4. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

prepare Zero
press ZERO

5. Press **ZERO** key.

   Wait for a **reaction period of 5 minutes**.

   After the reaction period is finished the measurement starts automatically.

Countdown
5:00

6. Remove the vial from the sample chamber.
7. Add **one EDTA tablet** straight from the foil to the prepared vial and crush the tablet using a clean stirring rod.
8. Close the vial tightly with the cap and swirl several times until the tablet is dissolved.
9. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
press ZERO
press TEST

10. Press **TEST** key.

    The result is shown in the display in mg/l Zinc.

1.1 Methods

Notes:

1. The tablets must be added in the correct sequence.
2. In the case of high levels of residual chlorine, perform the analysis with a dechlorinated water sample. To dechlorinate add one DECHLOR tablet to the water sample (point 1). Crush and mix to dissolve the tablet. Then add the COPPER / ZINC LR tablet (point 2) and continue with the test procedure as described above.

1.2 Important notes

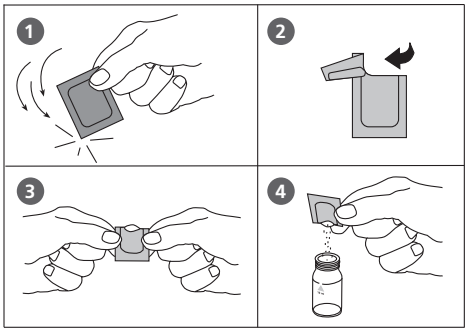
1.2.1 Correct use of reagents

The reagents must be added in the correct sequence.

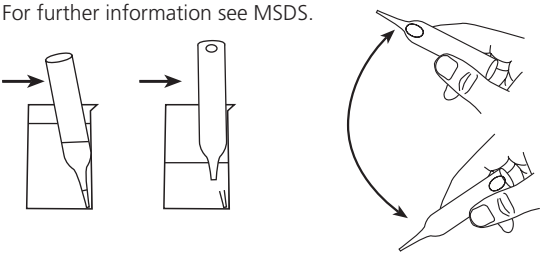
Tablet reagents:
The tablet reagents should be added to the water sample straight from the foil without touching them with the fingers.

Liquid reagents:
Add drops of the same size to the water sample by holding the bottle vertically and squeezing slowly.
After use replace the bottle caps securely noting the colour coding.
Note recommendation for storage (e.g. cool and dry).

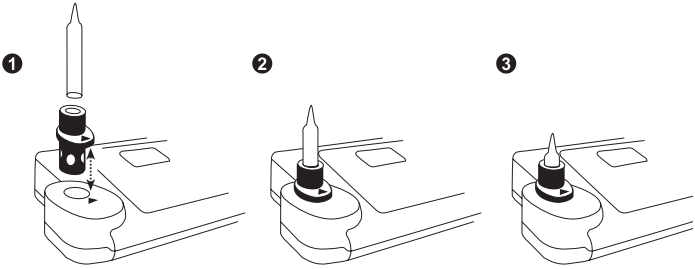
Powder Packs:



Vacu-vials® from CHEMetrics:
Vacu-vials® should be stored in the dark and at room temperature.
For further information see MSDS.



Correct position of the adapter and the vial (Ø 13 mm):



1.2.2 Cleaning of vials and accessories for analysis

Vials, caps and stirring rods should be cleaned thoroughly **after each analysis** to prevent interferences.

Procedure:

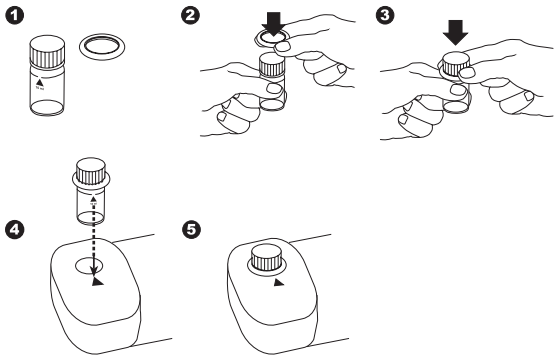
Clean vials and accessories after each analysis as soon as possible.

a. Clean vials and accessories with laboratory detergent
(e.g. Extran® MA 02 (neutral, phosphatic), Extran® MA 03 (alkaline, phosphate-free) from Merck KGaA).
b. Rinse thoroughly with tap water.
c. On demand (see Notes) perform special cleaning as required, e.g.: rinse with diluted Hydrochloric acid solution.
d. Rinse thoroughly with deionised water.

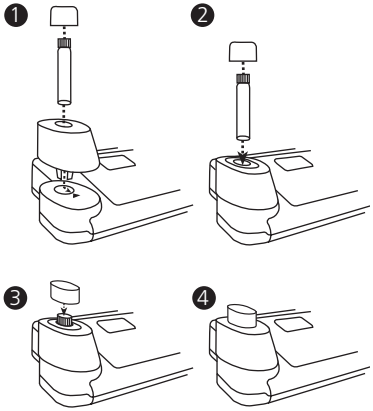
1.2.3 Guidelines for photometric measurements

1. Vials, caps and stirring rods should be cleaned thoroughly after each analysis to prevent interferences. Even minor reagent residues can cause errors in the test result.
2. The outside of the vial must be clean and dry before starting the analysis. Clean the outside of the vials with a towel. Fingerprints or other marks will be removed.
3. If there is no defined vial for the blank, the zeroing and the test must be carried out with the same vial as there may be slight differences in optical performance between vials.
4. The vials must be positioned in the sample chamber for zeroing and test with the Δ mark on the vial aligned with the ∇ mark on the instrument.

Correct position of the vial (Ø 24 mm):

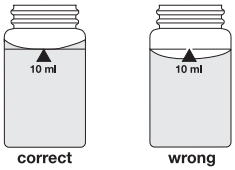


Correct position of the vial (Ø 16 mm):



5. Always perform zeroing and test with closed vial cap. Only use cap with sealing ring.
6. Bubbles on the inside wall of the vial lead to incorrect measurements. To prevent this, remove the bubbles by swirling the vial before performing the test.
7. Avoid spillage of water in the sample chamber. If water should leak into the instrument housing, it can destroy electronic components and cause corrosion.
8. Contamination of the lens in the sample chamber can result in errors. Check at regular intervals and – if necessary – clean the light entry surfaces of the sample chamber using a moist cloth or cotton buds.
9. Large temperature differences between the instrument and the environment can lead to errors – e.g. due to the formation of condensation in the area of the lens or on the vial.
10. To avoid errors caused by stray light do not use the instrument in bright sunlight.

Correct filling of the vial:



1.2.4 Sample dilution techniques

Proceed as follows for accurate dilutions:

Pipette the water sample (see table) into a 100 ml volumetric flask and fill up to 100 ml mark with deionised water. Swirl to mix the contents.

Water sample [ml]	Multiplication factor
1	100
2	50
5	20
10	10
25	4
50	2

Pipette the required volume of the diluted sample into the vial and proceed as described in the test methods.

Caution:

1. Dilution decreases accuracy.
2. Do not dilute water samples for measurement of pH values. This will lead to incorrect test results. If "Overrange" is displayed use another instrument (e.g. pH-meter).

1.2.5 Correcting for volume additions

If a larger volume of acid or base is used to pre-adjust the pH-value, a volume correction of the displayed result is necessary.

Example:

For adjusting the pH-value of a 100 ml water sample 5 ml of acid had to be added. The corresponding displayed result is 10 mg/l.

Total volume = 100 ml + 5 ml = 105 ml

Correction factor = 105 ml / 100 ml = 1.05

Corrected result = 10 mg/l x 1.05 = 10.5 mg/l

Part 2

Operating manual

2.1 Operation

2.1.1 Commissioning

Before working with the photometer insert the rechargeable batteries and the Lithium battery (delivery contents). The rechargeable batteries are not charged. See chapter 2.1.2 Saving data – Important Notes, 2.1.3 Replacement of rechargeable batteries resp. Lithium battery. and 2.1.4 Charging the rechargeable batteries.

Before using the photometer perform the following settings in the Mode-Menu:

- MODE 10: select language
- MODE 12: set date and time
- MODE 34: perform „Delete data"
- MODE 69: perform "User m. init" to initialise the userpolynomial system

See chapter 2.4 Photometer settings.

2.1.2 Saving data – Important Notes

The Lithium battery saves data (stored results and photometer setting) if there is no power from the power supply from the rechargeable batteries or the mains adapter.
Recommendation: Exchange of the lithium battery every 5 years.
Note: When neither mains adapter nor batteries supply energy to the instrument, all stored data and settings will be lost, if the lithium battery is taken out.
Recommendation: Keep the instrument connected to mains adapter supply while changing the lithium battery.

2.1.3 Replacement of rechargeable batteries resp. Lithium-battery

1. Switch the instrument off.
2. If necessary remove vial from the sample chamber.
3. Place the instrument upside down on a clean and even surface.
4. Unscrew the two screws (A) of the battery compartment cover (B).
5. Lift off battery compartment cover.
6. If necessary remove old rechargeable batteries (C) and/or the Lithium-battery (D) (See 2.1.4).
7. Place 7 new rechargeable batteries and/or the Lithium-battery.
   Ensuring the correct polarity!
8. Replace the battery compartment cover.
9. Tighten the screws carefully.

CAUTION
Dispose of used rechargeable batteries and Lithium-batteries in accordance with all federal, state and local regulations.

2.1.4 Charging the rechargeable batteries

The rechargeable batteries are uncharged in the instrument. As soon as the photometer is connected with the mains adapter to the mains the rechargeable batteries are charged.

Empty rechargeable batteries should be charged in the instrument for at least 5 days. 10 charging and discharging cycles are necessary before the rechargeable batteries obtain their full capacity.

it is possible to operate the instrument with the adapter with or without inserted rechargeable batteries.

2.1.5 Fuse

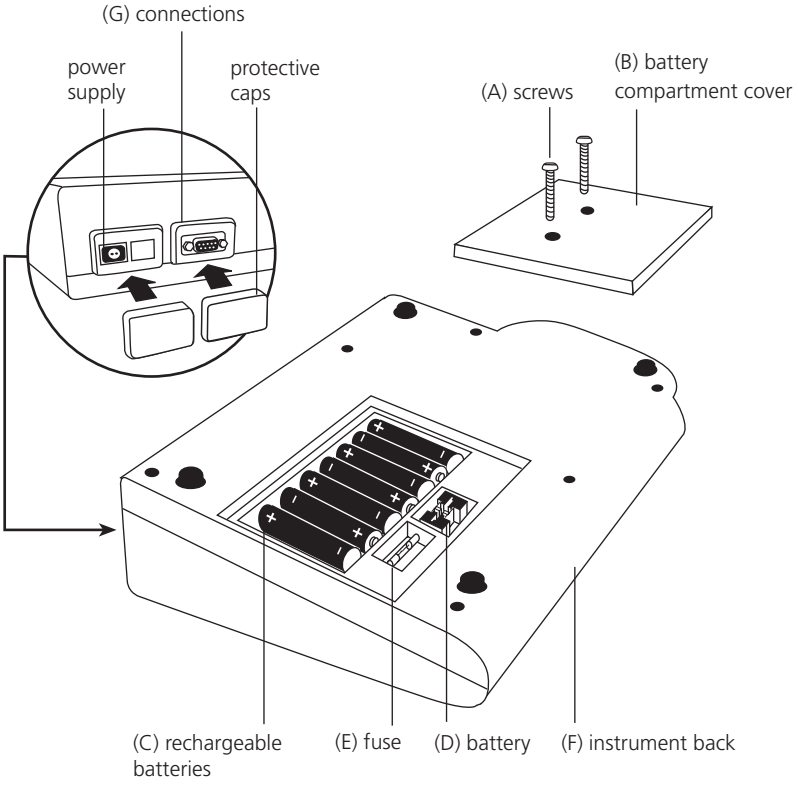
The instrument contains a fuse (E) (type: 1 A, inert, 20 mm).
If a replacement is necessary proceed as described in "Replacement of rechargeable batteries resp. Lithium-battery)". If the instrument can be operated with the mains adapter but not with the rechargeable batteries, the fuse could be defect (try new rechargeable batteries first).

2.1.6 Protective caps

If not used protect the two connections against damage (e.g. corrosion) caused by environmental influences (e.g. dust or splashing) keep the protective caps in place (G).

2.1.7 Instrument view

(A) screws
(B) battery compartment cover
(C) rechargeable batteries — 7 Ni-MH-rechargeable batteries (Type AA, 1100 mAh)
(D) battery — Lithium-battery (Type CR 2032, 3V)
(E) fuse — 1 A, inert, 20 mm
(F) instrument



2.2 Overview of function keys

Attention:
With the software-update V012.002.3.003.001 an "ESC-function" is implemented. If your keypad doesn't show an [Esc]-key please note that the grey key without a print (lowest key on the left) has the "ESC-function".

2.2.1 Overview

Switching the photometer on or off

Returning to selection of methods or previous menu

Function key: description in the text if key available

Function key: description in the text if key available

Function key: description in the text if key available


Confirming

Menu of photometer settings and further functions


Moving the cursor up or down

Storing of displayed test result

Performing Zero

Performing Test


Displaying date and time / user-countdown

2.2.2 Displaying time and date:


Press ["clock"] key.

19:30:22 2009-06-15

The display shows:

After 15 seconds the photometer reverts to the previous display automatically


or press **[↵]** key or [ESC].

2.2.3 User countdown

With this function the operator is able to define his own countdown.

Press [“clock”] key.

19.30.20 2009-06-15

The display shows time and date:

Press [“clock”] key.

Countdown
mm : ss
99 : 99

The display shows:

Either press [↲] key to accept the last used user countdown.

or

press any number key to start entering a new value

The entry comprises two digits each.
Enter minutes and seconds,
e.g.: 2 minutes, 0 seconds = [0][2][0][0].
Confirm with **[↲]** key.

Countdown
02:00
start: ↲

The display shows:

Start countdown with **[↲]** key.

After countdown has finished the photometer reverts to the previous display automatically.

2.3 Operation mode

Switch the photometer on by pressing the [ON/OFF] key.

selftest ...

The photometer performs an electronic self-test.

2.3.1 Automatic switch off

The instrument switches off automatically after 20 minutes. This is indicated 30 seconds before by a beeper. Press any key to avoid the instrument switching off.
As long as the instrument is working (for example countdown or printing) the automatic switch off is inactive.

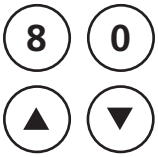
2.3.2 Selecting a method

>>	**30 Alkalinity-m**
	35 Alkalinity-p
	40 Aluminium

The display shows a selection:

There are two possibilities to select the required method:

a) enter method-number directly



e.g.: [8] [0] to select Bromine

b) press arrow key [▼] or [▲] to select the required method from the displayed list.

Confirm with [↵] key.

2.3.2.1 Method Information (F1)

Use [F1] key to switch between the compact and the detailed list for method selection.

100 Chlorine
0.02-6 mg/l Cl_2
Tablet
24 mm
DPD No 1
DPD No 3

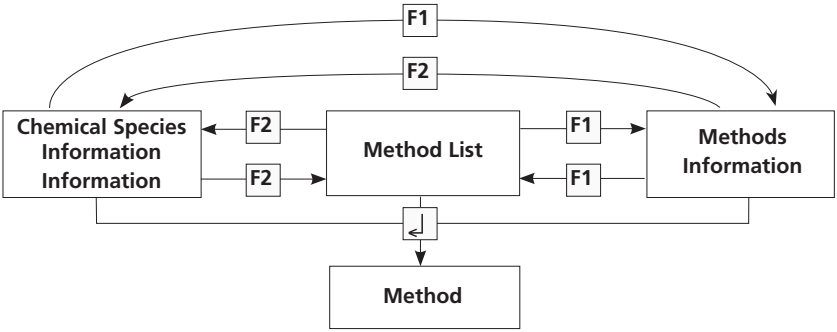
Example:

Line 1:	Method number, Method name
Line 2:	Range
Line 3:	Kind of reagent
Line 4:	Vial
Line 5-7:	Used reagent
tube =	reagent vial contained in tube test

2.3.2.2 Chemical Species Information

Pressing the F2 key the display shows a list with available chemical species and corresponding ranges. Changing chemical species see chapter 2.3.7 page 214.

320 Phosphate LR T **0.05-4 mg/l PO_4** **0.02-1.3 mg/l P** **0.04-3 mg/l P_2O_5**	Line 1: Method number, Method name Line 2: Range with chemical species 1 Line 3: Range with chemical species 2 Line 4: Range with chemical species 3



2.3.3 Differentiation

Chlorine **>> diff** **free** **total**	Differentiation is possible in some methods (e.g. Chlorine). The photometer then requires the type of determination.

Press arrow key [▼] or [▲] to select the required determination.

Confirm with **[↵]** key.

2.3.4 Performing Zero

prepare Zero **press ZERO**	The display shows:

Prepare a clean vial as described in "Method" and place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Press **ZERO** key.

Zero accepted **prepare Test** **press TEST**	The display shows:

2.3.5 Performing Tests

When zero calibration is complete, remove the vial from the sample chamber and perform the tests as described under "Method".

When the results have been displayed:

– with some methods you can change between different chemical species
– you can store and/or print out the results
– perform further analysis with the same zero
– select a new method

2.3.6 Ensuring reaction periods (countdown)

To ensure compliance with reaction periods a time delay is incorporated: the countdown. There are two kinds of countdowns:

Countdown
2:00

start: ↵

- Press **[↵]** key.
  Prepare water sample, start countdown with **[↵]** key and proceed as described in the mode description.
  The vial must not be placed in the sample chamber.

- Press **TEST** key.
  Prepare the water sample as described in the method description and place the vial in the sample chamber. The display shows the countdown by pressing the **TEST** key and the countdown is started automatically. After the reaction period is finished the measurement starts automatically.

Countdown
1:59

Notes:

1. It is possible to finish the working countdown by pressing the **[↵]** key. Reading starts immediately. In this case the operator is responsible for ensuring the necessary reaction period.
   Non-compliance with reaction periods leads to incorrect test results.
2. The time remaining is displayed continuously.
   The beeper indicates the last 10 seconds.

2.3.7 Changing chemical species

For some methods there is a possibility to change the chemical species of the test result. If the test result is displayed press arrow key [▲] or [▼].

Example:

320 Phosphate LR T -----[▼]----> 0.05-4 mg/l PO_4
<---- [▲] -----
1.00 mg/l PO_4

320 Phosphate LR T 0.02-1.3 mg/l P
0.33 mg/l P

<---- [▼] -----
------ [▲] ---->
320 Phosphate LR T 0.04-3 mg/l P_2O_5
0.75 mg/l P_2O_5

If the species of a test result is changed the displayed range is adjusted automatically.For an already stored result it is not possible to change the chemical species. The last displayed chemical species is kept by the instrument and will be displayed if this method is used the next time. If there is the possibility to change the chemical species for a method it is described in the manual. The arrows indicate the possible chemical species and are printed below the notes of the method:

▲ PO_4
P
▼ P_2O_5

2.3.8 Storing results

Store

Press **STORE** key while the test result is displayed.

Code No.:
_ _ _ _ _ _

The display shows:

① ⓪ ⓪ ⓪ ⓪ ⑥

- We advise you to enter a numeric code (up to 6 places). (A Code No. can contain references to the operator or the samplling location.)

  After entering confirm with **[↵]** key.

- If a code number is not necessary confirm by pressing **[↵]** directly. (The assignment for the Code No. is then 0 automatically.)

The entire data set is stored with date, time, Code No., method and test result.

Stored!

The display shows:

The test result is then shown again.

Note:

Storage: 900
free records left

The display shows the number of free data sets.

Storage: only 29
free records left

If there are less than 30 data sets free the display shows:

Clear the memory as soon as possible (see "Deleting stored results"). If memory capacity is used up it is impossible to save additional test results.

2.3.9 Printing results

If a printer is installed and switched on, it is possible to print out the test results (without saving it beforehand).

Press **F3** key.

The entire data set is printed with date, time, code no., method and test result. Printing example:

100 Chlorine T
0.02-6 mg/l Cl_2
Profi-Mode: no
2009-07-01 14:53:09
Test No.: 1
Code No.: 007
4.80 mg/l Cl_2

The test no. is an internal number that is set automatically if a test result is stored. It appears only on the print out.

2.3.10 Perform additional measurements

To perform additional tests using the same method:

Zero accepted
prepare Test
press TEST

- Press **TEST** key

The display shows:

Confirm with **TEST** key

or

- Press **ZERO** key to perform a new zero calibration.

prepare Zero
press ZERO

The display shows:

2.3.11 Selecting a new method

Press [ESC] key to return to method selection.

Or enter the required method number directly, e.g. [1] [6] [0] for Cyanuric acid.

Confirm with [↲] key.

2.3.12 Measure absorbance

Range: –2600 mAbs to +2600 mAbs

Method-No.	Title
900	mAbs 430 nm
910	mAbs 530 nm
920	mAbs 560 nm
930	mAbs 580 nm
940	mAbs 610 nm
950	mAbs 660 nm

Select the desired wavelength from the method list or by entering the corresponding method number directly.

900 mAbs 430 nm
-2600 mAbs - + 2600 mAbs
prepare Zero
press ZERO

The display shows e.g.:

Always carry out zeroing using a filled (e.g. deionised water) vial.

Zero accepted
prepare Test
press TEST

The display shows:

Carry out measurement of the sample.

500 mAbs

The display shows e.g.:

TIP: To ensure complete reaction times the user countdown may be helpful (chapter 2.2.3, page 210).

2.4 Photometer settings: Table of Mode-Functions

MODE-Function	No.	Description	Page
Calibration	40	Special method calibration	232
Clear calibration	46	Deleting user calibration	240
Clock	12	Setting date and time	219
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Langelier	70	Calculation of Langelier saturation Index (Water Balance)	252
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Method list	60	User method list, adaption	242
M list all on	61	User method list, switching on all methods	243
M list all off	62	User method list, switching off all methods	243
Print	20	Printing all stored results	222
Print, code no.	22	Print only results of a selected code no. range	224
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System info	91	Information about the instrument e.g. current software version	254
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User concentration	64	Entering the data necessary to run a user concentration method	244
User polynoms	65	Entering the data necessary to run a user polynomial	246
User methods clear	66	Delete all data of a user polynomial or of a concentration method	249
User methods print	67	Print out all data stored with mode 64 (concentration) or mode 65 (polynomial)	250
User methods init	69	Initialise the user method system (polynomial and concentration)	251

The selected settings are kept by the photometer even when switched off. To change photometer settings a new setting is required.

2.4.1 blank because of technical requirements

2.4.2 Instrument basic settings 1

Selecting a language

Press [MODE] [1] [0] keys.

Confirm with [↵] key.

```
<Language>
        Deutsch
   >>   English
        Francais
```

The display shows:

Press arrow key [▼] or [▲] to select the required language from the displayed list.

Confirm with [↵] key.

Key beep

Press [MODE] [1] [1] keys.

Confirm with [↵] key.

<Key Beep> **ON: 1 OFF: 0**

The display shows:

- Press [0] key to switch the key beep off.

- Press [1] key to switch the key beep on.

Confirm with [↵] key.

Note:
In the case of methods with reaction periods, an acoustic signal still sounds during the last 10 seconds of the countdown even if the key beep is switched off.

Setting date and time

Press [MODE] [1] [2] keys.

Confirm with [↵] key.

<clock>	
yy-mm-dd	**hh:mm**
_ _-_ _-_ _	_ _:_ _

The display shows:

The entry comprises two digits each.

yy-mm-dd	**hh:mm**
09-05-14	_ _:_ _

Enter year, month and day,
e.g.: 14. May 2009 = [0][9][0][5][1][4]

yy-mm-dd	**hh:mm**
09-05-14	**15:07**

Enter hours and minutes
e.g.: 3.07 p.m. = [1][5][0][7]

Confirm with [↵] key.

Note:
While confirming date and time with [↵] key the seconds are adjusted to zero automatically.

Countdown (Ensuring reaction periods)

Some methods require a reaction period. This reaction period is incorporated in the method as standard with the countdown function.
It is possible to switch the countdown off for all methods:

Press [MODE] [1] [3] keys.

Confirm with [↵] key.

<Countdown>
ON: 1 OFF: 0

The display shows:

- Press [0] key to switch the countdown off.

- Press [1] key to switch the countdown on.

Confirm with [↵] key.

Notes:

1. It is possible to interrupt the working countdown by pressing the [↵] key (application e.g. serial analysis).
   The "user countdown" is also available if the countdown is switched off.
2. If the countdown function is switched off, the operator is responsible for ensuring the necessary reaction period.
   Non-compliance with reaction periods leads to incorrect test results.

Signal beep

Performing a zero or a measurement takes 8 seconds. The photometer indicates the end of zeroing or measuring by a short beep.

Press [MODE] [1] [4] keys.

Confirm with [↵] key.

<Signal Beep>
ON: 1 OFF: 0

The display shows:

- Press [0] key to switch the signal beep off.

- Press [1] key to switch the signal beep on.

Confirm with [↵] key.

Note:
In the case of methods with reaction periods, an acoustic signal still sounds during the last 10 seconds of the countdown even if the key beep is switched off.

2.4.3 Printing of stored results

Printing all results

Press [MODE] [2] [0] keys.

Confirm with [↵] key.

<Print>
Print all Data
Start: ↵
cancel: ESC

The display shows:

Press [↵] key for printing out all stored test results.

Test No.:

The display shows e.g.:

After printing the photometer goes back to <Mode-Menu> automatically.

Note:
It is possible to cancel the entry by [ESC].
All stored data are printed out.

Printing results of a selected time period

Press [MODE] [2] [1] keys.

Confirm with [↲] key.

<Print>
sorted: date
from yy-mm-dd
__-__-__

The display shows:

Enter year, month and day for the first day of the required period, e.g.: 14 May 2009 = [0][9][0][5][1][4]

Confirm with [↲] key.

to yy-mm-dd
__-__-__

The display shows:

Enter year, month and day for the last day of the required period, e.g.: 19 May 2009 = [0][9][0][5][1][9]

Confirm with [↲] key.

from 2009-05-14
to 2009-05-19
Start: ↲
cancel: ESC

The display shows:

Press [↲] key and all stored results in the selected date range are printed.

After printing the photometer goes back to mode menu automatically.

Note:
It is possible to cancel the entry by [ESC].
If you want to print only results of one day enter the same date twice to determine the period.

Printing results of a selected Code No. range

Press [MODE] [2] [2] keys.

Confirm with [↲] key.

<Print>
sorted: Code-No.
from _ _ _ _ _ _

The display shows:

Enter numeric code number (up to 6 places) for the first required Code No., e.g.: [1].

Confirm with [↲] key.

to _ _ _ _ _ _

The display shows:

Enter numeric code number (up to 6 places) for the last required Code No., e.g.: [1] [0].

Confirm with [↲] key.

from 000001
to 000010
Start: ↲
cancel: ESC

The display shows:

Press [↲] key and all stored results in the selected code number range are printed.

After printing the photometer goes back to mode menu automatically.

Note:
It is possible to cancel the entry by [ESC].
If you want to print only results of one code number enter the same code number twice.
If you want to print all results without code no. (code no. is 0) enter Zero [0] twice.

Printing results of one selected method

Press [MODE] [2] [3] keys.

Confirm with [↵] key.

```
<Print>
>>20 Acid demand
   30 Alkalinity-tot
   40 Aluminium T
```

The display shows:

Select the required method from the displayed list or enter the method number directly.

Confirm with [↵] key.

In case of differentiated methods select the required kind of determination and confirm with [↵] key.

```
<Print>
method
30 Alkalinity-tot
Start:  ↵
cancel: ESC
```

The display shows:

Press [↵] key and all stored results of the selected method are printed.

After printing the photometer goes back to mode menu automatically.

Printing Parameter

Press [MODE] [2] [9] keys.

Confirm with [↲] key.

```
<printing parameter>
1: Flow control
2: Baud rate

cancel:              ESC
```

The display shows:

Press [1] key to select "Flow control".

```
<Flow Control>
is: Hardware
select:  [▲] [▼]
save:                ↲
cancel:              ESC
```

The display shows:

Press arrow key [▼] or [▲] to select the required Protocol.
(Xon/Xoff, Hardware, no control)

Confirm with [↲] key.

Finish with [ESC] key.
Flow Control will be set to the selection displayed at "is".

Press [2] key to select "Baud rate".

```
<Baud rate>
is: 19200
select:  [▲] [▼]
save:                ↲
cancel:              ESC
```

The display shows:

Press arrow key [▼] or [▲] to select the required baud rate. (600, 1200, 2400, 4800, 9600, 14400, 19200)

Confirm with [↲] key.

End with [ESC] key.

Back to Mode Menu with [ESC] key.

Back to method selection with [ESC] key.

Note:
Select "Hardware" as Protocol and "19200" as baud rate if you use the printer **DP 1012**.
Select "Hardware" as Protocol and "9600" as baud rate if you use the printer **DPN 2335**.
For setting of the printer see chapter 2.5.1 Connection to a printer.

2.4.4 Recall / delete stored results

Recall all stored results

Press [MODE] [3] [0] keys.

Confirm with **[↲]** key.

<Storage>
display all data
Start: ↲ cancel: ESC
print: F3
print all: F2

The display shows:

The stored data sets are displayed in chronological order, starting with the latest stored test result. Press **[↲]** key and all stored results are displayed.

- Press [F3] key to print the displayed result.
- Press [F2] key to print all results.
- End with [ESC].
- Press arrow key [▼] to display the following test result.
- Press arrow key [▲] to display the previous test result.

no data

If there are no test results in memory the display shows:

Recall results of a selected time period

Press [MODE] [3] [1] keys.

Confirm with [↲] key.

<Storage>
sorted: date
from yy-mm-dd
__-__-__

The display shows:

Enter year, month and day for the first day of the required period, e.g.: 14 May 2009 = [0][9][0][5][1][4]

Confirm with [↲] key.

to yy-mm-dd
__-__-__

The display shows:

Enter year, month and day for the last day of the required period, e.g.: 19 May 2009 = [0][9][0][5][1][9]

Confirm with [↲] key.

from 2009-05-14
to 2009-05-19
Start: ↲ cancel: ESC
print: F3
print all: F2

The display shows:

- Press [↲] key and all stored results in the selected date range are displayed.
- Press [F3] key to print the displayed result.
- Press [F2] key to print all selected results.
- End with [ESC].

Note:
It is possible to cancel the entry by [ESC].
If you want to recall only results of one day enter the same date twice to determine the time period.

Recall results of a selected Code No. range

Press [MODE] [3] [2] keys.

Confirm with [↵] key.

<Storage>
sorted: Code No.
from _ _ _ _ _ _

The display shows:

Enter numeric code number (up to 6 places) for the first required Code No., e.g.: [1].

Confirm with [↵] key.

to _ _ _ _ _ _

The display shows:

Enter numeric code number (up to 6 places) for the last required Code No., e.g.: [1] [0].

Confirm with [↵] key.

from 000001
to 000010
Start: ↵ cancel: ESC
print: F3
print all: F2

The display shows:

- Press [↵] key and all stored results in the selected Code No. range are displayed.
- Press [F3] key to print the displayed result.
- Press [F2] key to print all selected results.
- End with [ESC].

Note:
It is possible to cancel the entry by [ESC].
If you want to recall only results of one code number enter the same code number twice.
If you want to recall all results without code no. (code no. is 0) enter Zero [0] twice.

Recall results of one selected method

Press [MODE] [3] [3] keys.

Confirm with [↲] key.

<Storage>
>>20 Acid demand
30 Alkalinity-tot
40 Aluminium T

The display shows:

Select the required method from the displayed list or enter the method number directly.

Confirm with [↲] key.

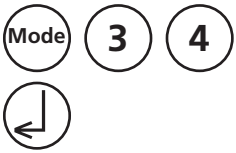
In case of differentiated methods select the required kind of determination and confirm with [↲] key.

<Storage>
method
30 Alkalinity-tot
Start: ↲ cancel: ESC
print: F3
print all: F2

The display shows:

- Press [↲] key and all stored results of the selected method are displayed.
- Press [F3] key to print the displayed result.
- Press [F2] key to print all selected results.
- End with [ESC].

Delete stored results



Press [MODE] [3] [4] keys.

Confirm with [↲] key.

```
<Delete data>
Delete all data?
 YES : 1   NO : 0
```

The display shows:

- Press [0] key to retain the data sets in memory.

- After pressing key [1] the following acknowledgment is displayed:

```
<Delete data>
Delete data ↲
Do not delete: ESC
```

Press [↲] key to delete.

ATTENTION:
All stored test results are deleted

or cancel without deleting data by pressing [ESC] key.

Note:
All stored test results are deleted.

2.4.5 Calibration

Calcium Hardness Method 191 – Calibration of a method blank

Mode 4 0

Press [MODE] [4] [0] keys.

Confirm with [↲] key.

<Calibration>
1: M 191 Ca-Hardness 2
2: M 191 reset 0 cali.
3: M 170 Fluoride L

The display shows:

1

Press [1] key.

<Calibration>
M191 Ca-Hardness 2T
prepare ZERO
press ZERO

The display shows:

1. Fill a clean vial (24 mm Ø) with exactly **10 ml of deionised water,** close tightly with the cap.
2. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero

3. Press **ZERO** key.
4. Remove the vial from the sample chamber.
5. Pipette 100 ml of water free of calcium to an appropriate beaker (note 2, 3).
6. Add **10 CALCIO H No. 1 tablets** straight from the foil to the 100 ml of water, crush the tablets using a clean stirring rod and dissolve the tablets completely.
7. Add **10 CALCIO H No. 2 tablets** straight from the foil to the same water, crush the tablets using a clean stirring rod and dissolve the tablets completely.

8. Press [↲] key.

Zero accepted
Countdown
2:00
start: ↲

Wait for a **reaction period of 2 minutes**.

After the reaction period is finished proceed as follows:

9. Rinse the vial (24 mm Ø) with the coloured sample from the beaker and fill with 10 ml of the sample.

prepare TEST **press TEST**	10. Press **TEST** key.
stored	The batch related method blank is saved.
	Press [↵] key, to go back to mode menu.

Notes:

1. If a new batch of CALCIO tablets is used a calibration of the method blank has to be performed to optimise the results.
2. Deionised or tap water
3. If no water free of Calcium is available these ions can be masked by using EDTA. Preparation: Add 50 mg (a spatula-tipful) EDTA to 100 ml water and dissolve.
4. To achieve the most accurate method blank it is important to adhere exactly to the sample volume of 100 ml.

Calcium Hardness Method 191 – Reset method blank to factory calibration

	Press [MODE] [4] [0] keys.
	Confirm with [↵] key.
<Calibration> **1: M 191 Ca-Hardness 2** **2: M 191 reset 0 cali.** **3: M 170 Fluoride L**	The display shows:
	Press [2] key.
<Calibration> **M191 Ca-Hardness 2T** **Reset ?** **YES: 1 , NO: 0**	The display shows:

0

Press [0] key to keep the method blank.

1

Press [1] key to erase the method blank and set the value back to factory calibration.

The instrument goes back to mode menu automatically.

Fluoride Method 170

Mode 4 0

Press [MODE] [4] [0] keys.

Confirm with [↵] key.

<Calibration>
1: M191 Ca-Hardness 2
2: M191 reset 0 cali.
3: M170 Fluoride L

The display shows:

3

Press [3] key.

<Calibration>
M170 Fluoride L
ZERO: deionised water
press ZERO

The display shows:

1. Fill a clean vial (24 mm Ø) with exactly **10 ml of deionised water**, close tightly with the cap.
2. Place the vial in the sample chamber making sure that the marks ⧖ are aligned.
3. Press **ZERO** key.
4. Remove the vial from the sample chamber.
5. Add **exactly 2 ml SPADNS reagent solution** to the water sample. **Caution: Vial is filled up to the top!**
6. Close the vial tightly with the cap and swirl gently several times to mix the contents.

7. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

Zero accepted
T1: 0 mg/l F
press TEST

8. Press **TEST** key.

9. Remove the vial from the sample chamber, empty the vial, rinse vial and cap several times and then fill the vial with exactly **10 ml Fluoride standard** (Concentration 1 mg/l F).

10. Add **exactly 2 ml SPADNS reagent solution** to the Fluoride standard.
Caution: Vial is filled up to the top!

11. Place the vial in the sample chamber making sure that the ⧖ marks are aligned.

T1 accepted
T2: 1 mg/l F
press TEST

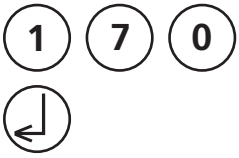
12. Press **TEST** key.

Calibration
accepted
↵

The display shows:

Confirm with [↵] key.

Back to method selection with **ESC** key.



Select Fluoride method with keys [1][7][0] and [↵].

Error, absorbance
T2 > T1
↵

If an error message appears please repeat adjustment.

Notes:

1. The same batch of SPADNS reagent solution must be used for adjustment and test. The adjustment process needs to be performed for each new batch of SPADNS reagent solution (see Standard methods 20th, 1998, APHA, AWWA, WEF 4500 F D., S. 4-82).
2. As the test result is highly dependent on exact sample and reagent volumes, the sample and reagent volumes should always be metered by using a 10 ml resp. 2 ml volumetric pipette (class A).

User Calibration

If a test method is user calibrated the method name is displayed inverse.

Procedure:

- Prepare a standard of known concentration and use this standard instead of the sample according to the test procedure.
- It is recommend to use well known standards which are formulated according to DIN EN, ASTM or other international norms or to use certified standards which are commercially available.
- After measuring this standard solution it is possible to change the displayed results to the required value.
- If a method uses a mathematic equation for the calculation of the result, it is only possible to calibrate the basic tests since all the other tests use the same polynomial.
- The same applies for some test procedures which use a polynomial from another test procedure.

Return to factory calibration:

If the user calibration is deleted the factory calibration is automatically activated.

Remarks:

The method "Fluoride" cannot be calibrated with mode 45 since the test requires a calibration related to the batch of the liquid reagent (SPADNS) (mode 40, chapter "Fluoride Method 170").

Table

No.	Method	Recommended range for user calibration
20	Acid demand	1–3 mmol/l
35	Alkalinity-p	100–300 mg/l $CaCO_3$
30	Alkalinity-total	50–150 mg/l $CaCO_3$
40	Aluminium T	0.1–0.2 mg/l Al
50	Aluminium PP	0.1–0.2 mg/l Al
60	Ammonia T	0.3–0.5 mg/l N
62	Ammonia PP	0.3–0.5 mg/l N
65	Ammonia LR TT	1 mg/l N
66	Ammonia HR TT	20 mg/l N
85	Boron	1 mg/l B
80	Bromine	Calibration with basic test 100 Chlorine free
63	Chloramine, mono	3–4 mg/l Cl_2
90	Chloride	10–20 mg/l Cl
100	Chlorine T	0.5–1.5 mg/l Cl
101	Chlorine L	Calibration with basic test 100 Chlorine free
110	Chlorine PP	0.5–1 mg/l Cl_2
105	Chlorine (KI) HR	70–150 mg/l Cl
120	Chlorine dioxide	Calibration with basic test 100 Chlorine free

No.	Method	Recommended range for user calibration
130	COD LR	100 mg/l O_2
131	COD MR	500 mg/l O_2
132	COD HR	5 g/l O_2 = 5000 mg/lO_2
204	Colour	operating range
150	Copper T	0.5–1.5 mg/l Cu
153	Copper PP	0.5–1.5 mg/l Cu
157	Cyanide	0.1–0.3 mg/l CN
160	Cyanuric acid	30–60 mg/l Cys
165	DEHA T	200–400 µg/l DEHA
167	DEHA PP	200 µg/l DEHA
170	Fluoride	Calibration with 0 and 1 mg/l F through Mode 40
190	Hardness, Calcium	100–200 mg/l $CaCO_3$
191	Hardness, Calcium	100–200 mg/l $CaCO_3$
200	Hardness, total T	15–25 mg/l $CaCO_3$
201	Hardness, total HR T	Calibration with basic test 200 Hardness, total
205	Hydrazine P	0.2–0.4 mg/l N_2H_4
206	Hydrazine L	0.2–0.4 mg/l N_2H_4
207	Hydrazine C	0.2–0.4 mg/l N_2H_4
210	Hydrogen peroxide	Calibration with basic test100 Chlorine free
215	Iodine	Calibration with basic test 100 Chlorine free
220	Iron T	0.3–0.7 mg/l Fe
222	Iron PP	0.1–2 mg/l Fe
223	Iron (TPTZ) PP	0.3–0.7 mg/l Fe
240	Manganese T	1–2 mg/l Mn
242	Manganese PP	0.1–0.4 mg/l Mn
243	Manganese HR PP	4–6 mg/l Mn
250	Molybdate T	5–15 mg/l Mo
252	Molybdate HR PP	10–30 mg/l Mo
265	Nitrate TT	10 mg/l N
270	Nitrite T	0.2–0.3 mg/l N
272	Nitrite LR PP	0.1–0.2 mg/l N
280	Nitrogen, total LR	10 mg/l N
281	Nitrogen, total HR	50–100 mg/l N
300	Ozone (DPD)	Calibration with basic test 100 Chlorine free
290	Oxygen, active	Calibration with basic test 100 Chlorine free
292	Oxygen, dissolved	possible against meter for dissolved oxygen
329	pH-Value LR	6.0–6.6
330	pH-Value T	7.6–8.0
331	pH-Value L	7.6–8.0
332	pH-Value HR	8.6–9.0
70	PHMB	15–30 mg/l
320	Phosphate LR T	1–3 mg/l PO_4
321	Phosphate HR T	30–50 mg/l PO_4

No.	Method	Recommended range for user calibration
323	Phosphate, ortho PP	0.1–2 mg/l PO_4
324	Phosphate, ortho TT	3 mg/l PO_4
327	Phosphate 1, ortho C	20–30 mg/l PO_4
328	Phosphate 2, ortho C	1–3 mg/l PO_4
325	Phosphate, total TT	0.3–6 mg/l P
326	Phosphate, hydr. TT	0.3–0.6 mg/L P
316	Phosphonate	1–2 mg/l PO_4
340	Potassium	3 mg/l K
350	Silica	0.5–1.5 mg/l SiO_2
351	Silica LR PP	1 mg/l SiO_2
352	Silica HR PP	50 mg/l SiO_2
212	Sodium hypochlorite	8 %
360	Sulfate PP	50 mg/l SO_4
355	Sulfate T	50 mg/l SO_4
365	Sulfide	0.2–0.4 mg/l S
370	Sulfite	3–4 mg/l SO_3
384	Suspended Solids	operating range
386	Turbidity	operating range
390	Urea	1–2 mg/l CH_4N_2O
400	Zinc	0.2–0.4 mg/L Zn

Store user calibration

100 Chlorine T **0.02-6 mg/l Cl2** **0.90 mg/l free Cl2**

Perform the required method as described in the manual using a standard of known concentration instead of the water sample.

If the test result is displayed press [MODE] [4] [5] keys and confirm with [↲] key.

The display shows:

<user calibration> **100 Chlorine T** **0.02-6 mg/l Cl2** **0.90 mg/l free Cl2** **up: ↑, down: ↓** **save: ↲**

Pressing the arrow key [▲] once increases the displayed result.

Pressing the arrow key [▼] once decreases the displayed result.

Press keys till the displayed result corresponds to the value of the standard.

Confirm with [↲] key to store the new calibration factor.

Cancel user calibration by pressing [ESC] key.

Jus Factor **saved**

The display shows:

100 Chlorine T **0.02-6 mg/l Cl2** **1.00 mg/l free Cl2**

Now the method name is displayed inverse and the test result is calculated with the new calibration factor.

Delete user calibration

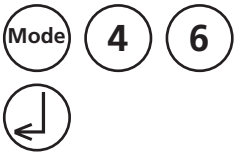
This chapter only applies for methods which can be user calibrated.

100 Chlorine T
0.02-6 mg/l Cl2

Select the required method.

prepare ZERO
press ZERO

Instead of zeroing the instrument press [MODE] [4] [6] keys and confirm with [↵] key.



<user calibration>
100 Chlorine T
0.02-6 mg/l Cl2
clear user
calibration?
YES: 1, NO: 0

The display shows:

Press [1] key to delete user calibration.

Press [0] key to keep the valid user calibration.

The instrument goes back to Zero-query automatically.

2.4.6 Lab function

Reduced operator guidance => “Profi-Mode“

This function may be used for routine analyses with many samples of one method.
The following information is always stored in the methods:

a) Method
b) Range
c) Date and time
d) Differentiation of results
e) Detailed operator instruction
f) Compliance with reaction periods

If the Profi-Mode is active, the photometer provides only a minimum of operator instructions. The criteria specified above in d, e, f are no longer included.

Press [MODE] [5] [0] keys in succession.

Confirm with [↵] key.

<Profi-Mode>
ON : 1 OFF : 0

The display shows:

- Press [0] key to switch the Profi-Mode off.

- Press [1] key to switch the Profi-Mode on.

switched off

The display shows:

or

switched on

Confirm with [↵] key.

Note:

Storage of test results is possible. When results are stored the display also shows “Profi-Mode”.
The selected settings are kept by the photometer even when it is switched off. To change photometer setting a new setting is required.

2.4.7 User operations

User method list

After switching on the instrument a scroll list of all available methods is automatically shown in the display. To shorten this list according to the requirements of the user it is possible to create a user defined scroll list.

The program structure requires that this list must have at least one active (switched on) method. For this reason it is necessary to activate first all required methods and then to switch off the automatically activated one if this method is not required.

User method list, adaptation

Press [MODE] [6] [0] keys.

Confirm with [↲] key.

```
<Method list>
selected: •
toggle: F2
save: ↲
cancel: ESC
```

The display shows:

Start with [↲] key.

```
<Method list>
>> 30•Alkalinity-tot
   40•Aluminium
   50•Ammonium
....
```

The complete method list is displayed.

Methods with a point [•] behind the method number will be displayed in the method selection list. Methods without a point will not be displayed in the method selection list.

```
>> 30•Alkalinity-tot
```

Press key [▲] or [▼] to select the required method from the displayed list.

Switch with [F2] key between "active" [•] and "inactive" [ ].

```
>> 30 Alkalinity-tot
```

Select next method, activate or inactivate it and continue.

```
>> 30•Alkalinity-tot
```

Confirm with [↲] key.

Cancel without storing by pressing [ESC] key.

Recommendation:

If only a few methods are required it is recommended to perform Mode 62 first, followed by Mode 60.

All user Polynomials (1-25) and Concentrations (1-10) are displayed in the method list, although they are not programmed by the user. Non-programmed user methods can't be activated!

User method list, switch all methods on

This mode function activates all methods. After switching on the instrument a scroll list of all available methods is automatically shown in the display.

Press [MODE] [6] [1] keys.

Confirm with [↵] key.

<Mlist all on>
switch on all methods
YES: 1, NO: 0

The display shows:

- Press [1] key to display all methods in the method selection list.

- Press [0] key to keep the valid method selection list.

The instrument goes back to mode menu automatically.

User method list, switch all methods off

The program structure requires that the method list must have at least one active (switched on) method. For this reason the instrument activates one method automatically.

Press [MODE] [6] [2] keys.

Confirm with [↵] key.

<Mlist all off>
switch off all methods
YES: 1, NO: 0

The display shows:

- Press [1] key to display only one method in the method selection list.

- Press [0] key to keep the valid method selection list.

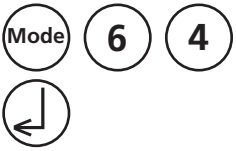
The instrument goes back to mode menu automatically.

User Concentration Methods

It is possible to enter and store up to 10 User Concentration Methods.
Therefor you need 2 to 14 standards of known concentration and one blank (deionised water or reagent blank value). The Standards should be measured with increasing concentrations and from the brightest to the darkest colouration.
The measuring range for „Underrange" and „Overrange" is defined with -2600 mAbs* and +2600 mAbs*. After selection of a method the concentration of the lowest and highest used standard is displayed as measuring range. The operation range should be within this range to achieve best results.

*1000 mAbs = 1 Abs = 1 E (displayed)

Entering a User Concentration:



Press [MODE] [6] [4] keys.

Confirm with [↵] key.

Entry Procedure:

```
< User concentr.>
choose no.: ____
(850-859)
```

The display shows:

Enter a method number in the range from 850 to 859, e.g.: [8] [5] [0]

Confirm with [↵] key.

```
Overwrite conc. meth.?
YES: 1,   NO: 0
```

Note:
if the entered number has already been used to save a concentration the display shows the query:

- Press [0] or [ESC] key to go back to method no. query.
- Press [1] key to start entry mode.

```
wavelength:
1: 530 nm
2: 560 nm
3: 610 nm
```

Enter the required wavelength, e.g.: [2] for 560 nm.


```
choose unit:
>>
        mg/l
        g/l
        mmol/l
        mAbs
        µg/l
        E
        A
        %
```

Press [▲] or [▼] keys to select the required unit.

Confirm with [↵] key.

choose resolution
1: 1
2: 0.1
3: 0.01
4: 0.001

③

Press the appropriate numerical key to select the required resolution, e.g.: for 0.01.

Note:
Please enter the required resolution according to the instrument pre-sets:

range	max. resolutions
0.000 ...9.999	0.001
10.00 ...99.99	0.01
100.0... 999.9	0.1
1000 ...9999	1

Measurement procedure with standards of known concentration:

< User concentr.>
prepare Zero
press ZERO

Zero

The display shows:

Prepare Zero and press [Zero] key.

Note:
Use deionised water or reagent blank value.

< User concentr.>
Zero accepted
S1: +_____
↲ | ESC | F1

The display shows:

Enter the concentration of the first standard;
e.g.: 0.05

- One step back with [ESC].
- Press [F1] key to reset numerical input.

⓪ . ⓪ ⑤

↲

Confirm with [↲] key.

< User concentr.>
S1: 0.05 mg/l
prepare
press TEST

The display shows:

Test

Prepare the first standard and press [Test] key.

S1: 0.05 mg/l
mAbs: 12 ↲

The display shows the input value and the measured absorption value. Confirm with [↲] key.

S1 accepted
S2: +_____
↲ | ESC | F1

Enter the concentration of the second standard;
e.g.: 0.1

- One step back with [ESC].
- Press [F1] key to reset numerical input.

⓪ . ①

↲

Confirm with [↲] key.

S2: 0.10 mg/l **prepare** **press TEST**	Prepare the second standard and press [Test] key.
S2: 0.10 mg/l **mAbs: 150 ↲**	The display shows the input value and the measured absorption value. Confirm with [↲] key.
S2 accepted **S3: +_____** **↲ | ESC | F1 | Store**	**Note:** • Perform as described above to measure further standards. • The minimum of measured standards is 2. • The maximum of measured standards is 14 (S1 to S14).
Store	If all required standards or the maximum value of 14 standards are measured press [Store] key.
stored!	The display shows: The instrument goes back to the mode menu automatically. Now the concentration is stored in the instrument and can be recalled by entering its method number or selecting it from the displayed method list.

TIP:
Save all your concentration data in a written form because in case of power outage (e.g. changing the battery) all concentration data will be lost and must be entered again.
You might want to use Mode 67 to transfer all concentration data to a PC.

User Polynomials

It is possible to enter and store up to 25 User Polynomials.
The program allows the user to apply a Polynomial up to the 5th degree:

$y = A + Bx + Cx^2 + Dx^3 + Ex^4 + Fx^5$

If only a Polynomial of a lower degree is necessary the other coefficients are specified as zero (0), e.g.: for the 2nd degree is D, E, F = 0.
The values of the coefficients A, B, C, D, E, F must be entered in an academic notation with maximal 6 decimal places, e.g.: 121,35673 = 1,213567E+02

Entering a User Polynomial:

Mode 6 5	Press [MODE] [6] [5] keys.
↲	Confirm with [↲] key.
<User polynoms> **choose no.: ____** **(800-824)**	The display shows:
8 0 0	Enter a method number in the range from 800 to 824, e.g.: [8] [0] [0]

Confirm with [↵] key.

Overwrite polynom?
YES: 1, NO: 0

Note:
if the entered number has already been used to save a polynomial the display shows the query:

- Press [0] or [ESC] key to go back to method No. query.
- Press [1] key to start entry mode.

wavelength:
1: 530 nm 4: 430 nm
2: 560 nm 5: 580 nm
3: 610 nm 6: 660 nm

Enter the required wavelength, e.g.: [2] for 560 nm.

< User polynoms>
y = A+Bx+Cx²+Dx³+
Ex⁴+Fx⁵
A: +_____

- Press [▲] or [▼] key to change between plus and minus sign
- Enter data of the coefficient A including decimal point, e.g.: 1.32
- Press [F1] key to reset numerical input.

Confirm with [↵] key.

A: 1.32____ E+____

- Press [▲] or [▼] key to change between plus and minus sign
- Enter the exponent of the coefficient A, e.g.: 3

Confirm with [↵] key.

B: +_____

Successively the instrument queries the data for the other coefficients (B, C, D, E and F).

Note:
If zero [0] is entered for the value of the coefficient, the input of the exponent is omitted automatically.

Confirm every input with [↵] key.

measurement range
Min mAbs: +_____
Max mAbs: +_____

Enter measurement ranges from – 2600 to +2600 mAbs.

- Press [▲] or [▼] key to change between plus and minus sign.
- Enter the values in Absorbance (mAbs) for the upper limit (Max) and the lower limit (Min).

Confirm every input with [↵] key.

choose unit:
>>
mg/l
g/l
mmol/l
mAbs
µg/l
E
A
%

Press [▲] or [▼] keys to select the required unit.

Confirm with [↵] key.

choose resolution
1: 1
2: 0.1
3: 0.01
4: 0.001

③

Press the appropriate numerical key to select the required resolution.

Note:
Please enter the required resolution according to the instrument pre-sets:

range	max. resolutions
0.000 ...9.999	0.001
10.00 ...99.99	0.01
100.0... 999.9	0.1
1000 ...9999	1

stored!

The display shows:

The instrument goes back to the mode menu automatically.

Now the polynomial is stored in the instrument and can be recalled by entering its method number or selecting it from the displayed method list.

TIP:
Save all your polynomial data in a written form because in case of power outage (e.g. changing the battery) all polynomial data will be lost and must be entered again.
You might want to use Mode 67 to transfer all polynomial data to a PC.

Delete User Methods (Polynomial or Concentration)

In principle a valid user method can be overwritten.
An existing user method (Polynomial or Concentration) can be totally deleted as well and is removed out of the method selection list:

Press [MODE] [6] [6] keys.

Confirm with [↵] key.

<User m. clear>
choose no.: _____
(800-824), (850-859)

The display shows:

Enter the number of the User Method you want to delete (in the range from 800 to 824 or 850 to 859),
e.g.: 800

Confirm with [↵] key.

M800
delete?
YES: 1, NO: 0

The query is displayed:

- Press [1] key to delete the selected User Method.

- Press [0] key to keep the valid User Method.

The instrument goes back to mode menu automatically.

Print Data of User Methods (Polynomials & Concentration)

With this Mode function all data (e.g. wavelength, unit ...) of stored user polynomials and concentration methods can be printed out or transferred with HyperTerminal to a PC.

Press [MODE] [6] [7] keys.

Confirm with [↲] key.

<User m. print>
Start: ↲

The display shows:

Press [↲] key to print out the data (e.g. wavelength, unit, ...) of all stored User Methods.

M800
M803
...

The display shows e.g.:

After data transfer the photometer goes back to mode menu automatically.

Initialise User Method System (Polynomials & Concentration)

Power loss will cause incoherent data. The user method system must be initialised with this mode function to set it to a predefined state.

ATTENTION:
All stored user methods (polynomial & concentration) are deleted with initialisation.

Press [MODE] [6] [9] keys.

Confirm with [↲] key.

<User m. init>
Start: ↲

The display shows:

Confirm with [↲] key.

Initialising?
YES: 1, NO: 0

The query is displayed:

- Press [1] key to start initialisation.

- Press [0] key to to cancel without initialisation.

The instrument goes back to mode menu automatically.

2.4.8 Special functions

Langelier Saturation Index (Water Balance)

For calculation the following tests are required:

- pH value
- Temperature
- Calcium hardness
- Total Alkalinity
- TDS (Total Dissolved Solids)

Run each test separately and note the results.
Calculate the Langelier Saturation Index as described:

Calculation of Langelier Saturation Index

With Mode 71 (see below) it is possible to select between degree Celsius or degree Fahrenheit.

Press [MODE] [7] [0] keys.

Confirm with [↵] key.

<Langelier>
temperature °C:
3°C <=T<=53°C
+_ _ _ _

The display shows:

Enter the temperature value (T) in the range between 3 and 53°C and confirm with [↵] key. If °F was selected, enter the temperature value in the range between 37 and 128°F.

calcium hardness
50<=CH<=1000
+_ _ _ _

The display shows:

Enter the value for Calcium hardness (CH) in the range between 50 and 1000 mg/l $CaCO_3$ and confirm with [↵] key.

tot. alkalinity
5<=TA<=800
+_ _ _ _

The display shows:

Enter the value for Total Alkalinity (TA) in the range between 5 and 800 mg/l $CaCO_3$ and confirm with [↵] key.

total dissol. solids
0<=TDS<=6000
+_ _ _ _

The display shows:

Enter the value for TDS (Total Dissolved Solids) in the range between 0 and 6000 mg/l and confirm with [↵] key.

pH value **0<=pH<=12** **+_ _ _ _**	The display shows:
	Enter the pH value in the range between 0 and 12 and confirm with [↲] key.
<Langelier> **Langelier** **saturation index** **0.00** **Esc ↲**	The display shows the Langelier Saturation Index. Press [↲] key to start new calculation. Return to mode menu by pressing [ESC] key.

Examples:

Operating error:

Values out of defined range:

CH<=1000 mg/l CaCO3!	The entered value is too high.
CH>=50 mg/l CaCO3!	The entered value is too low.
	Confirm display message with [↲] key and enter a value in the defined range.

Selection of temperature unit

Entering the temperature value is possible in degree Celsius or degree Fahrenheit.
Therefore the following preselection is (once) required.

	Press [MODE] [7] [1] keys.
	Confirm with [↲] key.
<temperature> **1: °C 2: °F**	The display shows:
	Press [1] key to select degree Celsius.
	Press [2] key to select degree Fahrenheit.
	The instrument goes back to mode menu automatically.

2.4.9 Instrument basic settings 2

Adjusting display contrast

Press [MODE] [8] [0] keys.

Confirm with [↵] key.

<LCD contrast>
↑ ↓

The display shows:

Press arrow key [▲] to increase contrast of the LCD display.

Press arrow key [▼] to decrease contrast of the LCD display.

Press [Zero] key to increase contrast of the display about ten units.

Press [Test] key to decrease contrast of the display about ten units.

Confirm with [↵] key.

2.4.10 Instrument special functions /service

Photometer-Information

Press [MODE] [9] [1] keys.

Confirm with [↵] key.

<System-Info>
Software:
V201.001.1.001.002
more: ▼, cancel: Esc

This method informs you about the current software version, about the current detected mains power supply, about the number of performed tests and free memory capacity.

Press arrow key [▼] to display the number of performed tests and free memory capacity.

<System-Info>
Number of Tests:
139
free records left
999
cancel: Esc

Finish with [ESC] key.

2.5 Data transfer

Switch the photometer and the personal computer or printer off. Connect the photometer (RS232 interface) and the serial interface of the personal computer or printer using a cable in line with the specified assignment (see technical data). The cable for connection to a personal computer is included in delivery contents.

2.5.1 Connection to a printer

Printer with a serial connection are suitable for connection with the photometer (see chapter 3.4 Technical data interface).

A suitable paper tabel printer is the printer DPN 2335.

Before using the printer **DPN 2335** with the Photometer you should change the following standard adjustments:
(Detailed information of changing the adjustment you will find in the printer manual).

Baud rate:	**9600**
Parity:	**None**
Data bits:	**8**

Note: The printer must be connected and switched on before printing.

Caution: Adjust printing parameter in Mode 29. See chapter 2.4.3 Printing Parameter.

2.5.2 Data transfer to a personal computer

Transferring test results from the photometer to a personal computer requires a transfer program, e.g. HyperTerminal.
Please find detailed information at our homepage in the download-area.

2.5.3 Internet Updates

It is possible to update new software applications and additional languages via the internet.
Please find detailed information at our homepage in the download-area.

Please note:
To prevent loss of stored test results store or print them out before performing an Update.

2.6 blank because of technical requirements

Part 3

Enclosure

3.1 Unpacking

Carefully inspect all items to ensure that every part of the list below is present and no visible damage has occurred during shipment. If there is any damage or something is missing, please contact your local distributor immediately.

3.2 Delivery contents

Standard contents for MultiDirect:

√	
❑	1 Photometer in plastic case
❑	1 Adapter for 16 mm Ø vials
❑	1 Cap for adapter
❑	2 Protective caps for connections
❑	1 Rechargeable battery set (7 Ni-MH-cells; Type AA; 1100 mAh) or 4 batteries (Type AA / LR 6)
❑	1 Lithium battery (CR 2032; 3V)
❑	1 Mains adapter, 100 – 240 V, 50 – 60 Hz (not in basic version)
❑	1 Cable for connection to PC (not in basic version)
❑	3 Round vials with cap, height 48 mm, Ø 24 mm
❑	3 Round vials with cap, height 90 mm, Ø 16 mm
❑	1 Beaker, plastic, 100 ml
❑	1 Cleaning brush
❑	1 Stirring rod, plastic
❑	1 Syringe, plastic, 2 ml
❑	1 Syringe, plastic, 5 ml
❑	1 Syringe, plastic, 10 ml
❑	1 Instruction manual
❑	1 Guarantee declaration

Reagent sets are not part of the standard scope of delivery. Please see the General Catalogue for details of available reagent sets.

3.3 Blank because of technical requirements

3.4 Technical data

Display	Graphic Display (7-line, 21-characters)
Serial Interface	serial RS232 for printer- and PC-connection; 9-pin D-sub-mail connector, data format ASCII, 8-bit Data, no parity, 1 start-bit, 1 stop-bit, baud rate and protocol: adjustable
	Pin assignation: Pin 1 = free Pin 2 = Rx Data Pin 3 = Tx Data Pin 4 = free Pin 5 = GND Pin 6 = free Pin 7 = RTS Pin 8 = CTS Pin 9 = free
Light source	LEDs and photo sensor amplifier in protected cell compartment. Wavelength ranges: λ1 = 530 nm IF Δ λ = 5 nm λ2 = 560 nm IF Δ λ = 5 nm λ3 = 610 nm IF Δ λ = 6 nm λ4 = 430 nm IF Δ λ = 5 nm λ5 = 580 nm IF Δ λ = 5 nm λ6 = 660 nm IF Δ λ = 5 nm IF = Interference filter
Photometric accuracy*	0.100 Abs ± 0.008 Abs 1.000 Abs ± 0.020 Abs
Operation	Acid and solvent resistant touch-sensitive keyboard with integral beeper as acoustic indicator.
Power supply	7 Ni-MH cells (Type AA with 1100 mAh); external main adapter (Input: 100–240 V, 50–60 Hz; Output: 15V=/530 mA) Lithium battery (CR 2032, 3V); for keeping data if there is no power supply from the rechargeable batteries or the main adapter
Auto off	20 minutes after last function, 30 seconds acoustical signal before switch off
Charging time	approx. 10 hours
Dimensions	approx. 265 x 195 x 70 mm (unit) approx. 440 x 370 x 140 mm (case)
Weight (unit) batteries)	approx. 1000 g (with main adapter and rechargeable
Working condition (without condensation)	5 – 40 °C at max. 30–90 % relative humidity
Language options	English, German, French, Spanish, Italian, Portuguese, Polish; further languages via Internet Update
Storage capaity	ca. 1000 data sets

** measured with standard solutions (T = 20°C – 25°C)*

Subject to technical modification!

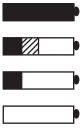
To ensure maximum accuracy of test results, always use the reagent systems supplied by the instrument manufacturer.

3.5 Abbreviations

Abbreviation	Definition
°C	degree Celsius (Centigrade)
°F	degree Fahrenheit °F = (°C x 1.8) + 32
°dH	degree German Hardness
°fH	degree French hardness
°eH	degree English Hardness
°aH	degree American Hardness
Abs	Absorption unit (≙ Extinction E) 1000 mAbs = 1 Abs ≙ 1 A ≙ 1 E
µg/l	(= ppb) Microgram per litre
mg/l	(= ppm) Milligram per litre
g/l	(= ppth) gram per litre
KI	Potassium iodide
$K_{S\,4.3}$	Acid demand to pH 4.3 – this method is similar to Total Alkalinity but converted into the unit "mmol/l", as the German DIN 38409 demand.
TDS	Total Dissolved Solids
LR	Low Range
MR	Medium Range
HR	High Range
C	Reagents from Chemetrics©
L	Liquid reagent
P	Powder (reagent)
PP	Powder Pack
T	Tablet
TT	Tube Test
DEHA	N,N-Diethylhydroxylamine
DPD	Diethyl-p-phenylendiamine
DTNB	Ellmans reagent
PAN	1-(2-Pyridylazo)-2-napthol
PDMAB	Paradimethylaminobenzaldehyde
PPST	3-(2-Pyridyl)-5,6-bis(4-phenylsulfonic acid)1,2,4-triazine
TPTZ	2,4,6-Tri-(2-Pyridyl)-1,3,5-triazine

3.6 Troubleshooting

3.6.1 Operating messages in the display / error display

Display	Possible Causes	Elimination
Overrange	reading is exceeding the range water sample is too cloudy too much light on the photo cell	if possible dilute sample or use other measuring range filtrate water sample seal on the cap? Repeat measurement with seal on the cap of the vial.
Underrange	result is under the detection limit	indicate result with lower x mg/l x = low end of measuring range; if necessary use other analytical method
Storagesystem error use Mode 34	mains power fails or is not connected	insert or change Lithium battery. Delete data with Mode 34
capacity of rechargeable battery 	full capacity warning signal every 3 minutes warning signal every 12 seconds warning signal, the instrument switches itself off	capacity of the rechargeable battery is too low; charge the rechargeable battery; operate instrument with mains adapter
Jus Overrange E4	The user calibration is out of the accepted range	Please check the standard, reaction time and other possible faults. Repeat the user calibration.
Jus Underrange E4		
Overrange E1	The concentration of the standard is too high/too low, so that during user calibration the limit of the range was exceeded	Perform the test with a standard of higher/lower concentration
Underrange E1		
E40 user calibration not possible	If the display shows Overrange/ Underrange for a test result a user calibration is not possible	Perform the test with a standard of higher/lower concentration
Zero not accepted	Light absorption is too great or too low	Refer to chapter 2.3.4 Performing Zero. Clean sample chamber. Repeat zeroing.

Display	Possible Causes	Elimination
???	The calculation of a value (e.g. combined Chlorine) is not possible	Test procedure correct? If not – repeat test
Example 1 0,60 mg/l free Cl ??? comb Cl 0,59 mg/l total Cl		Example 1: The readings for free and total Chlorine are different, but considering the tolerances of each reading they are the same. For this reason the combined Chlorine is most likely zero.
Example 2 Underrange ??? comb Cl 1,59 mg/l total Cl		Example 2: The reading for free Chlorine is under the detection limit. The instrument is not able to calculate the combined Chlorine. In this case the combined Chlorine is most likely the same as the total Chlorine.
Example 3 0,60 mg/l free Cl ??? comb Cl Overrange		Example 3: The reading for total Chlorine is exceeding the range. The instrument is not able to calculate the combined Chlorine. The test should be repeated with a diluted sample.
Error absorbance e.g.: T2>T1	Fluoride calibration was not correct	Repeat calibration
Printer „timeout“	printer switched off; no connection	Connect printer Check connections Switch printer on

3.6.2 General

Finding	Possible Causes	Elimination
Test result deviates from the expected.	Chemical species not as required.	Press arrow keys to select the required chemical species.
No differentiation: e.g. for the Chlorine test there is no selection between differentiated, free or total.	Profi-Mode is switched on.	Switch Profi-Mode off with Mode 50.
The pre-programmed countdown is not displayed.	Countdown is not activated and/or the Profi-Mode is activated.	Switch the countdown on with Mode 13 and/or switch the Profi-Mode off with Mode 50.
It seems that a method is not available.	Method is not activated in the user method list.	Activate the required method in the user method list with Mode 60.
Instrument can be operated with the mains adapter but not with the rechargeable batteries.	Rechargeable batteries are not charged or defect. Fuse (Type A, inert, 20 mm) may be defect.	Charge rechargeable batteries or change them. If the problem still exists change fuse.

3.7 Declaration of CE-Conformity

The manufacturer: **Tintometer GmbH**

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Germany

declares, that this product

Product name: **MultiDirect**

Conforms with EN 61 326 for specific defined electromagnetic environment.
Conforms with EN 61 326 (domestic).

Dortmund, 06. August 2003

Cay-Peter Voss, Managing Director

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