

Lovibond® Water Testing

Tintometer® Group



Manual of Methods

MD 100 • MD 110 • MD 200

Iron

(EN) Manual of Methods

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(ES) Manual de Métodos

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(IT) Manuale dei Metodi

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(NL) Handboek Methoden

Zijde 122

(DE) Methodenhandbuch

Seite 22

(FR) Méthodes Manuel

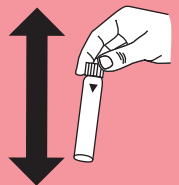
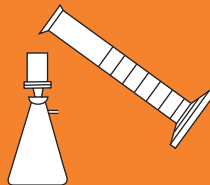
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(PT) Métodos Manual

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(ZH) 方法手册

Page 142



KS4.3 T / 20


Method name

Method number

Bar code for the detection of the methods

$K_{S4.3 T}$
20

0.1 - 4 mmol/l $K_{S4.3}$
S:4.3

Acid / Indicator

Measuring range

Chemical Method

Display in the MD 100 / MD 110 / MD 200

Instrument specific information

The test can be performed on the following devices. In addition, the required cuvette and the absorption range of the photometer are indicated.

Instrument Type	Cuvette	λ	Measuring Range
MD 200, MD 600, MD 610, MD 640, MultiDirect, PM 620, PM 630	ø 24 mm	610 nm	0.1 - 4 mmol/l $K_{S4.3}$
SpectroDirect, XD 7000, XD 7500	ø 24 mm	615 nm	0.1 - 4 mmol/l $K_{S4.3}$

Material

Required material (partly optional):

Reagents	Packaging Unit	Part Number
Alka-M-Photometer	Tablet / 100	513210BT
Alka-M-Photometer	Tablet / 250	513211BT

Application List

- Waste Water Treatment
- Drinking Water Treatment
- Raw Water Treatment

Notes

1. The terms Alkalinity-m, m-Value, total alkalinity and Acid demand to $K_{S4.3}$ are identical.
2. For accurate results, exactly 10 ml of water sample must be used for the test.

Language codes ISO 639-1

Revision status

EN Handbook of Methods 01/20

Performing test procedure

Implementation of the provision Acid capacity $K_{S_{4.3}}$ with Tablet

Select the method on the device

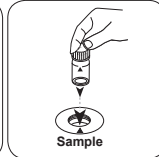
For this method, no ZERO measurements are to be carried out with the following devices: XD 7000, XD 7500



Fill 24 mm vial with **10 ml sample**.



Close vial(s).



Place **sample vial** in the sample chamber. • Pay attention to the positioning.

• • •



Dissolve tablet(s) by inverting.

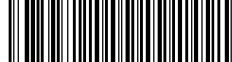


Place **sample vial** in the sample chamber. • Pay attention to the positioning.



Press the **TEST** (XD: **START**) button.

The result in Acid Capacity $K_{S_{4.3}}$ appears on the display.



Iron T

M220

0.02 - 1 mg/L Fe

FE

Ferrozine / Thioglycolate

EN

Material

Required material (partly optional):

Reagents	Packaging Unit	Part Number
Iron II LR (Fe ²⁺)	Tablet / 100	515420BT
Iron II LR (Fe ²⁺)	Tablet / 250	515421BT
Iron LR (Fe ²⁺ und Fe ³⁺)	Tablet / 100	515370BT
Iron LR (Fe ²⁺ und Fe ³⁺)	Tablet / 250	515371BT

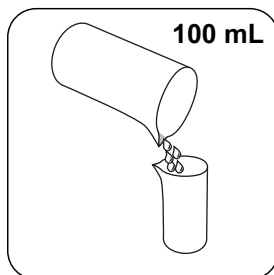
Preparation

1. Water that has been treated with organic compounds such as corrosion inhibitors, must be oxidised where necessary to break down the iron complex. 1 ml of concentrated Sulphuric acid ($\geq 95\%$) and 1 ml concentrated Nitric acid ($\geq 65\%$) is therefore added to 100 ml water sample and boiled down to approximately half the volume. After cooling down, the digestion procedure is continued.

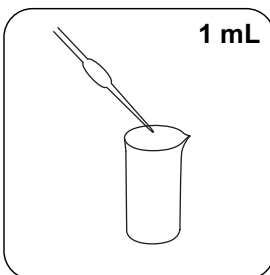
Notes

1. This method is for the determination of total dissolved Fe²⁺ and Fe³⁺.
2. For the determination of Fe²⁺, the IRON (II) LR Tablet, instead of the IRON LR Tablet is used.

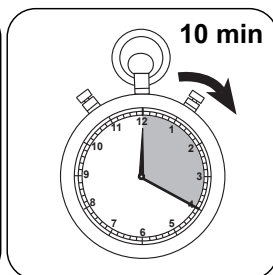
Digestion



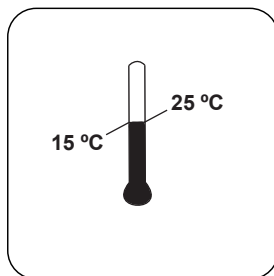
Fill a suitable sample vessel with **100 mL sample**.



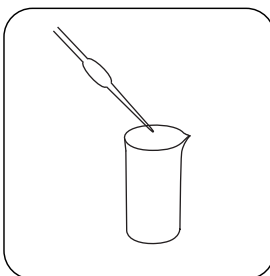
Add **1 mL concentrated sulfuric acid ($\geq 95\%$)**.



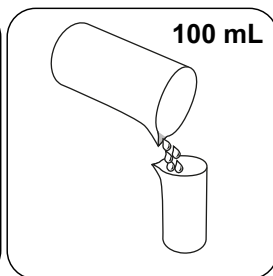
The sample is to be **heated for 10 minutes**, or for as long as it takes for everything to be completely dissolved.



Allow the sample to cool to room temperature.



Adjust **pH-value** of the sample with **ammonia solution (10-25 %)** to 3-5.



Fill the sample with **deionised water to 100 mL**.

This sample is used for the analysis of total solved and dissolved Iron.

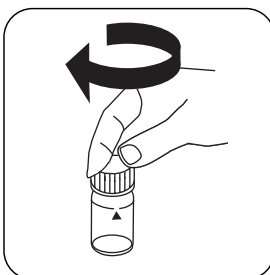
Determination of Iron (II,III), dissolved with Tablet

Select the method on the device.

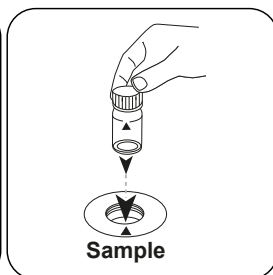
For testing of **dissolved and undissolved Iron**, carry out the described **digestion**.



Fill 24 mm vial with **10 mL sample**.



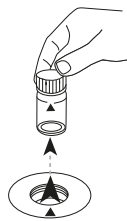
Close vial(s).



Place **sample vial** in the sample chamber. Pay attention to the positioning.



Zero

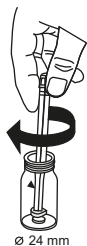


Press the **ZERO** button.

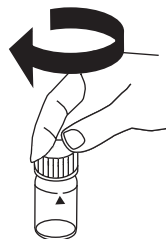
Remove the vial from the sample chamber.



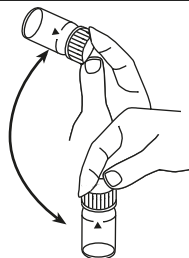
Add **IRON LR tablet**.



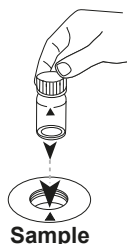
Crush tablet(s) by rotating slightly.



Close vial(s).



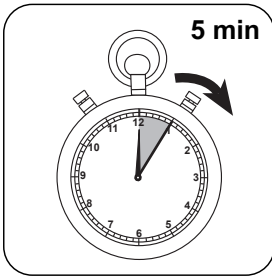
Dissolve tablet(s) by inverting.



Place **sample vial** in the sample chamber. Pay attention to the positioning.

Test

Press the **TEST** (XD: **START**) button.

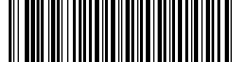


Wait for **5 minute(s)**
reaction time.

Once the reaction period is finished, the measurement takes place automatically.

The result in mg/L Iron appears on the display.

EN



Chemical Method

Ferrozine / Thioglycolate

Appendix

EN

Interferences

Removeable Interferences

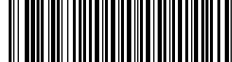
1. The presence of copper increases the test result by 10 %. At a concentration of 10 mg/L copper in the sample, the measurement result is increased by 1 mg/L iron. The interference can be eliminated by the addition of thiourea

Method Validation

Limit of Detection	0.01 mg/L
Limit of Quantification	0.016 mg/L
End of Measuring Range	1 mg/L
Sensitivity	0.92 mg/L / Abs
Confidence Intervall	0.013 mg/L
Standard Deviation	0.005 mg/L
Variation Coefficient	1.23 %

Bibliography

Photometrische Analyse, Lange/ Vjedelek, Verlag Chemie 1980, p. 102



Iron PP

M222

0.02 - 3 mg/L Fe^g)

FE1

1,10-Phenanthroline

EN

Material

Required material (partly optional):

Reagents	Packaging Unit	Part Number
VARIO Ferro F10	Powder / 100 pc.	530560
VARIO Ferro F10	Powder / 1000 pc.	530563

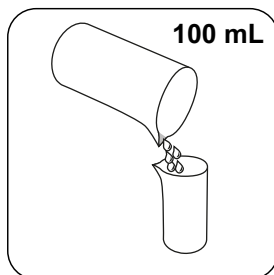
Preparation

1. Iron oxide requires mild, strong or Digesdahl digestion before the analysis (digestion process with acid).
2. Very strong alkaline or acidic water samples should be adjusted to between pH 3 and pH 5 before the analysis.
3. Water samples containing visible rust should be allowed to react for at least five minutes.
4. Water that has been treated with organic compounds such as corrosion inhibitors, must be oxidised where necessary to break down the iron complex. 1 ml of concentrated Sulphuric acid ($\geq 95\%$) and 1 ml concentrated Nitric acid ($\geq 65\%$) is therefore added to 100 ml water sample and boiled down to approximately half the volume. After cooling down, the digestion procedure is continued.

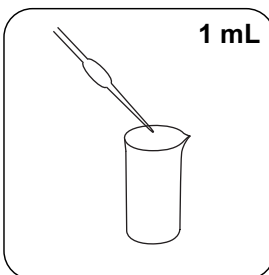
Notes

1. This method is for the determination of all forms of dissolved iron and most forms of undissolved iron.
2. Accuracy is not affected by undissolved powder.

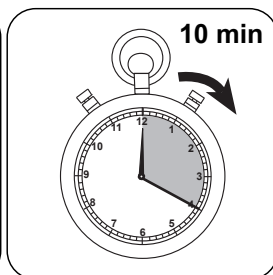
Digestion



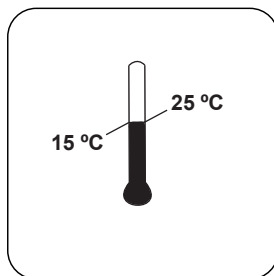
Fill a suitable sample vessel with **100 mL sample**.



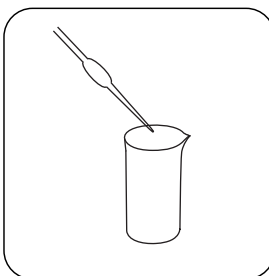
Add **1 mL concentrated sulfuric acid ($\geq 95\%$)**.



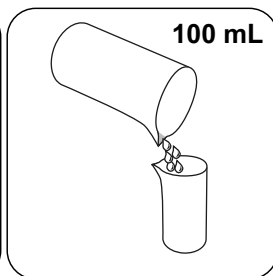
The sample is to be **heated for 10 minutes**, or for as long as it takes for everything to be completely dissolved.



Allow the sample to cool to room temperature.



Adjust **pH-value** of the sample with **ammonia solution (10-25 %)** to 3-5.



Fill the sample with **deionised water to 100 mL**.

This sample is used for the analysis of total solved and dissolved Iron.

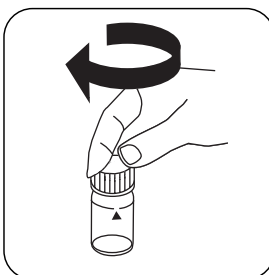
Determination of Iron (II,III), dissolved with Vario Powder Packs

Select the method on the device.

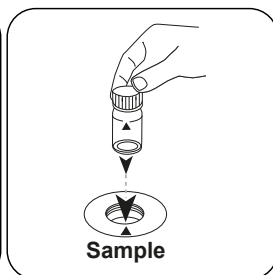
For testing of **Iron with tablet**, carry out the described **digestion**.



Fill 24 mm vial with **10 mL sample**.



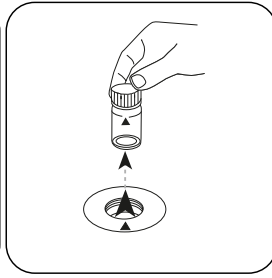
Close vial(s).



Place **sample vial** in the sample chamber. Pay attention to the positioning.

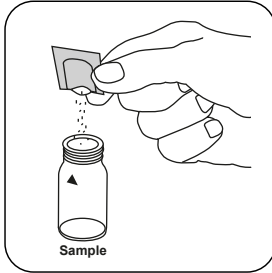


Zero

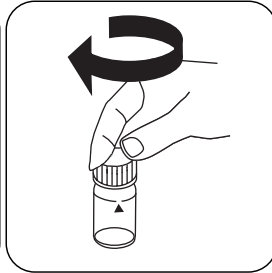


Press the **ZERO** button.

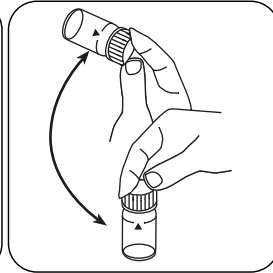
Remove the vial from the sample chamber.



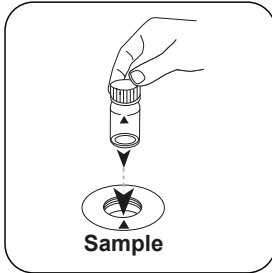
Add **Vario FERRO F10 powder pack**.



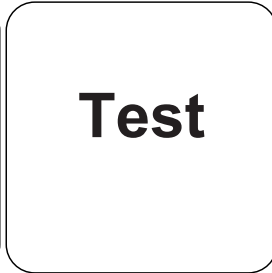
Close vial(s).



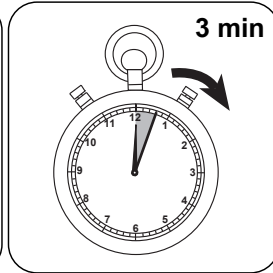
Invert several times to mix the contents.



Place **sample vial** in the sample chamber. Pay attention to the positioning.



Press the **TEST (XD: START)** button.



Wait for **3 minute(s) reaction time**.

Once the reaction period is finished, the measurement takes place automatically.
The result in mg/L Iron appears on the display.



Chemical Method

1,10-Phenanthroline

Appendix

Interferences

Persistent Interferences

1. Iridium interferes with the test.

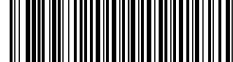
According to

DIN 38406-E1

Standard Method 3500-Fe-1997

US EPA 40 CFR 136

⁹⁾ Reagent recovers most insoluble iron oxides without digestion



Iron (TPTZ) PP

M223

0.02 - 1.8 mg/L Fe

FE2

TPTZ

EN

Material

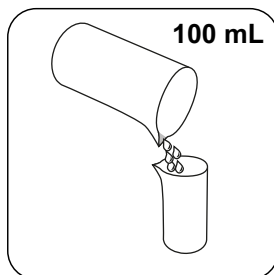
Required material (partly optional):

Reagents	Packaging Unit	Part Number
VARIO Iron TPTZ F10	Powder / 100 pc.	530550

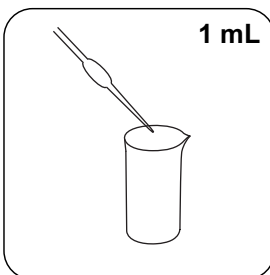
Preparation

1. Digestion is required for the determination of total Iron. The TPTZ reagent recovers most iron oxides without digestion.
2. All glassware must first be rinsed with diluted 1:1 Hydrochloric acid solution before the analysis and then rinsed with deionised water to remove iron deposits that can cause slightly high results.
3. Strong alkaline or acidic water samples should be adjusted between pH 3 and pH 8 before the analysis (use 0.5 mol/l Sulphuric acid or 1 mol/l Sodium hydroxide).
4. Water that has been treated with organic compounds such as corrosion inhibitors, must be oxidised where necessary to break down the iron complex. 1 ml of concentrated Sulphuric acid ($\geq 95\%$) and 1 ml concentrated Nitric acid ($\geq 65\%$) is therefore added to to 100 ml water sample and boiled down to approximately half the volume. After cooling down, the digestion procedure is continued.

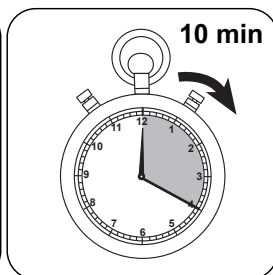
Digestion



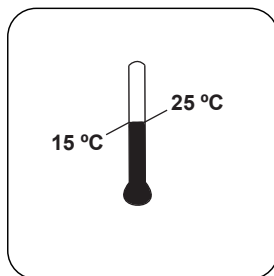
Fill a suitable sample vessel with **100 mL sample**.



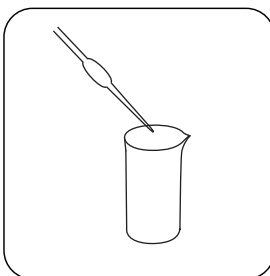
Add **1 mL concentrated sulfuric acid ($\geq 95\%$)**.



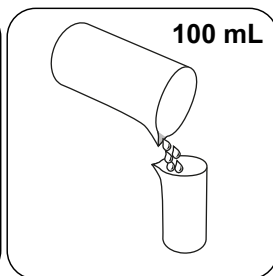
The sample is to be **heated for 10 minutes**, or for as long as it takes for everything to be completely dissolved.



Allow the sample to cool to room temperature.



Adjust **pH-value** of the sample with **ammonia solution (10-25 %)** to 3-5.



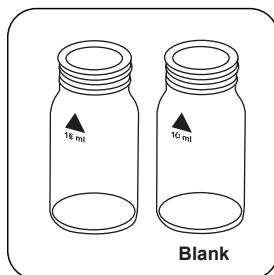
Fill the sample with **deionised water to 100 mL**.

This sample is used for the analysis of total solved and dissolved Iron.

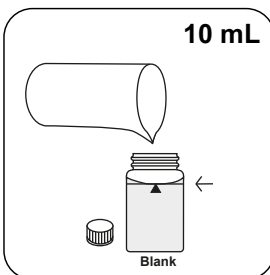
Determination of Iron, total with Vario Powder Pack

Select the method on the device.

For testing of **total Iron**, carry out the described **digestion**.



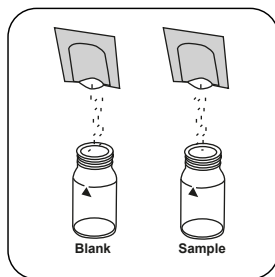
Prepare two clean 24 mm vials. Mark one as a blank.



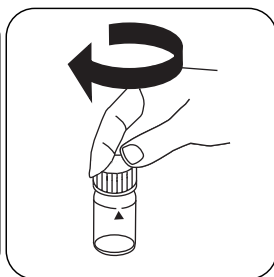
Put **10 mL deionised water** in the blank.



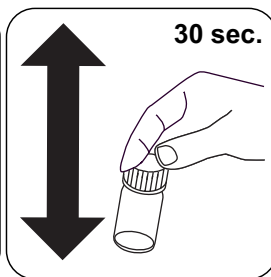
Put **10 mL sample** in the sample vial.



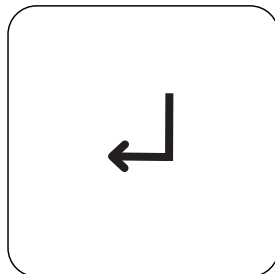
Add a **Vario IRON TPTZ F10 powder pack** in each vial.



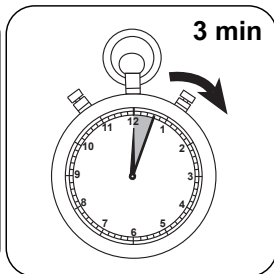
Close vial(s).



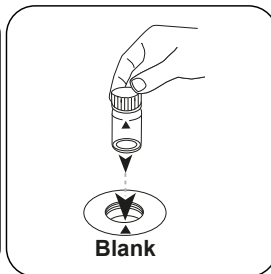
Mix the contents by shaking. (30 sec.).



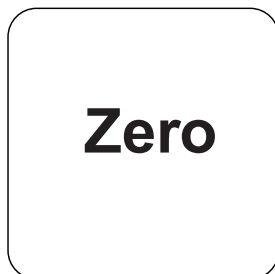
Press the **ENTER** button.



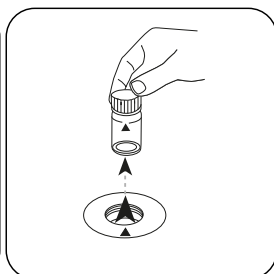
Wait for **3 minute(s) reaction time**.



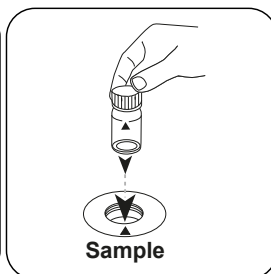
Place **blank** in the sample chamber. Pay attention to the positioning.



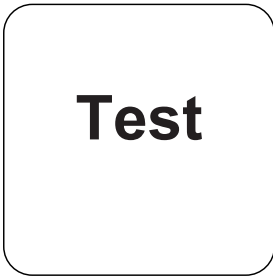
Press the **ZERO** button.



Remove the vial from the sample chamber.



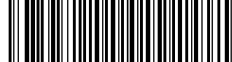
Place **sample vial** in the sample chamber. Pay attention to the positioning.



Press the **TEST** (XD:
START) button.

The result in mg/L Iron appears on the display.

EN



Chemical Method

TPTZ

Appendix

EN

Interferences


Persistent Interferences

When interferences occur, colour development is inhibited or a precipitate is formed. The values refer to a standard with an iron concentration of 0.5 mg/L.

Interference	from / [mg/L]
Cd	4
Cr ³⁺	0.25
Cr ⁶⁺	1.2
Co	0.05
Cu	0.6
CN ⁻	2.8
Mn	50
Hg	0.4
Mo	4
Ni	1
NO ₂ ⁻	0.8

Bibliography

G. Frederic Smith Chemical Co., The Iron Reagents, 3rd ed. (1980)

KS4.3 T / 20


Methoden Name

Methodennummer

Barcode zur Methodenerkennung

Messbereich

20

S:4.3

Chemische Methode

$K_{S_{4.3} T}$
0,1 - 4 mmol/l $K_{S_{4.3}}$
Säure / Indikator

Instrumentenspezifische Informationen

Der Test kann auf den folgenden Geräten durchgeführt werden. Zusätzlich sind die benötigte Küvette und der Absorptionsbereich der Photometer angegeben.

Geräte	Küvette	λ	Messbereich
MD 200, MD 600, MD 610, MD 640, MultiDirect, PM 620, PM 630	ø 24 mm	610 nm	0,1 - 4 mmol/l $K_{S_{4.3}}$
SpectroDirect, XD 7000, XD 7500	ø 24 mm	615 nm	0,1 - 4 mmol/l $K_{S_{4.3}}$

Material

Benötigtes Material (zum Teil optional):

Reagenzien	Form/Menge	Bestell-Nr.
Alka-M-Photometer	Tablette / 100	513210BT
Alka-M-Photometer	Tablette / 250	513211BT

Anwendungsbereich

- Abwasserbehandlung
- Trinkwasseraufbereitung
- Rohwasserbehandlung

Anmerkungen

1. Die Begriffe Alkalität-m, m-Wert, Gesamtalkalität und Säurekapazität $K_{S_{4.3}}$ sind identisch.
2. Die exakte Einhaltung des Probevolumens von 10 ml ist für die Genauigkeit des Analyseergebnisses entscheidend.

Sprachkürzel nach ISO 639-1

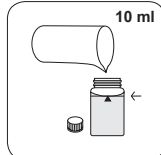
Revisionsstand

DE Methodenhandbuch 01/20

Durchführung der
Messung**Durchführung der Bestimmung Säurekapazität $K_{s4,3}$ mit Tablette**

Die Methode im Gerät auswählen.

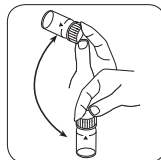
Für diese Methode muss bei folgenden Geräten keine ZERO-Messung durchgeführt werden: XD 7000, XD 7500

24-mm-Küvette mit **10 ml Probe** füllen.

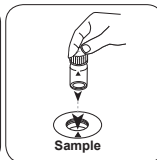
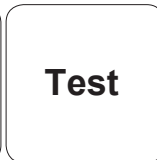
Küvette(n) verschließen.

Die **Probeküvette** in den Messschacht stellen. Positionierung beachten.

• • •



Tablette(n) durch Umschwenken lösen.

Die **Probeküvette** in den Messschacht stellen. Positionierung beachten.Taste **TEST** (XD: **START**) drücken.In der Anzeige erscheint das Ergebnis als Säurekapazität $K_{s4,3}$.



Eisen T

M220

0,02 - 1 mg/L Fe

FE

Ferrozine / Thioglycolat

Material

DE

Benötigtes Material (zum Teil optional):

Reagenzien	Form/Menge	Bestell-Nr.
Iron II LR (Fe ²⁺)	Tablette / 100	515420BT
Iron II LR (Fe ²⁺)	Tablette / 250	515421BT
Iron LR (Fe ²⁺ und Fe ³⁺)	Tablette / 100	515370BT
Iron LR (Fe ²⁺ und Fe ³⁺)	Tablette / 250	515371BT

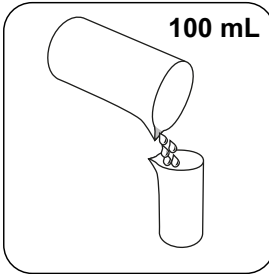
Vorbereitung

1. Wässer, die mit organischen Verbindungen als Korrosionsschutz usw. behandelt worden sind, müssen ggf. oxidiert werden, um die Eisenkomplexe zu zerstören. Dazu wird eine 100 ml Probe mit 1 ml konzentrierter Schwefelsäure ($\geq 95\%$) und 1 ml konzentrierter Salpetersäure ($\geq 65\%$) versetzt und auf die Hälfte eingedampft. Nach dem Abkühlen wird der Aufschluss durchgeführt.

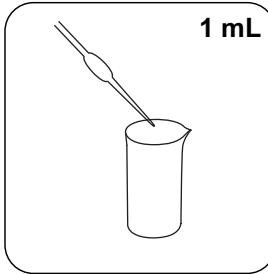
Anmerkungen

1. Bei dieser Methode erfolgt die Bestimmung von gesamt gelösten Fe²⁺ und Fe³⁺.
2. Zur Bestimmung von Fe²⁺ wird die IRON (II) LR Tablette, anstelle der IRON LR Tablette verwendet.

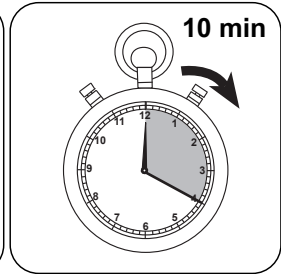
Aufschluss



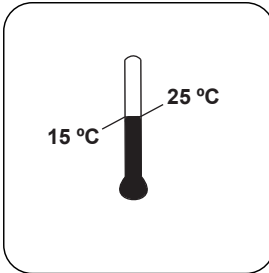
Ein geeignetes
Probengefäß mit **100 mL**
Probe füllen.



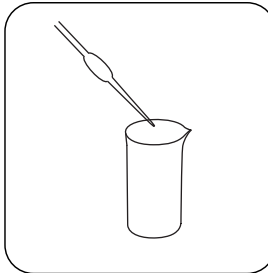
1 mL konzentrierte
Schwefelsäure (≥ 95 %)
zugeben.



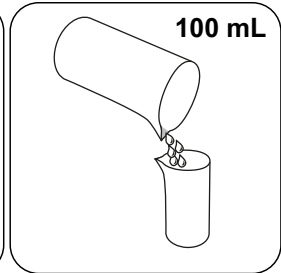
Die Probe für **10 Minuten**
erhitzen, oder so lange, bis
sich alles vollständig gelöst
hat.



Die Probe auf
Raumtemperatur
abkühlen lassen.



Den **pH-Wert** der Probe
mit **Ammoniaklösung**
(10-25 %) auf 3-5
einstellen.



Die Probe mit **VE-Wasser**
auf **100 mL** auffüllen.

Diese Probe für die Analyse von gesamten gelösten und ungelösten Eisen verwenden.
Durchführung der Bestimmung Eisen(II,III), gelöst mit Tablette

Die Methode im Gerät auswählen.

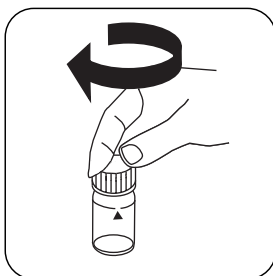
Für die Bestimmung von **gelöstem und ungelöstem Eisen** den beschriebenen
Aufschluss durchführen.



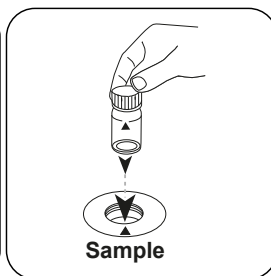
DE



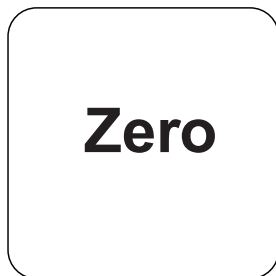
24-mm-Küvette mit **10 mL**
Probe füllen.



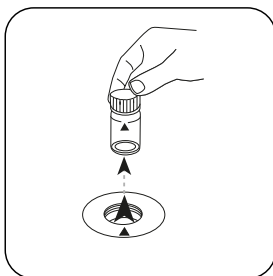
Küvette(n) verschließen.



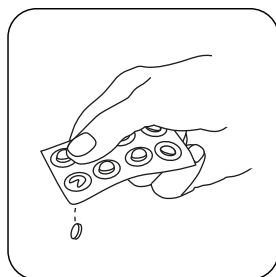
Die **Probeküvette** in
den Messschacht stellen.
Positionierung beachten.



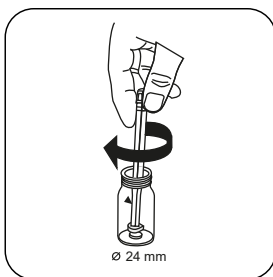
Taste **ZERO** drücken.



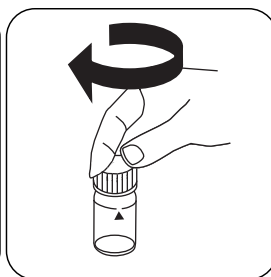
Küvette aus dem
Messschacht nehmen.



Eine **IRON LR** Tablette
zugeben.



Tablette(n) unter leichter
Drehung zerdrücken.



Küvette(n) verschließen.



Tablette(n) durch Umschwenken lösen.

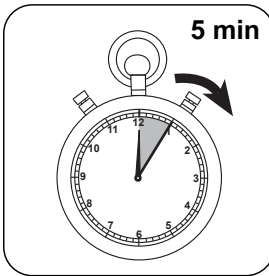


Die **Probenküvette** in den Messschacht stellen. Positionierung beachten.



Taste **TEST (XD: START)** drücken.

DE



5 Minute(n) Reaktionszeit abwarten.

Nach Ablauf der Reaktionszeit erfolgt automatisch die Messung.
In der Anzeige erscheint das Ergebnis in mg/L Eisen.



Chemische Methode

Ferrozine / Thioglycolat

Appendix

DE

Störungen

Ausschließbare Störungen

1. Die Anwesenheit von Kupfer erhöht das Messergebnis um 10 %. Bei einer Konzentration von 10 mg/L Kupfer in der Probe wird das Messergebnis um 1 mg/L Eisen erhöht.
Die Störung kann durch die Zugabe von Thioharnstoff beseitigt werden

Methodenvalidierung

Nachweisgrenze	0.01 mg/L
Bestimmungsgrenze	0.016 mg/L
Messbereichsende	1 mg/L
Empfindlichkeit	0.92 mg/L / Abs
Vertrauensbereich	0.013 mg/L
Verfahrensstandardabweichung	0.005 mg/L
Verfahrensvariationskoeffizient	1.23 %

Literaturverweise

Photometrische Analyse, Lange/ Vjedelek, Verlag Chemie 1980, S. 102



Eisen PP

M222

0,02 - 3 mg/L Fe⁹⁾

FE1

1,10-Phenanthroline

Material

DE

Benötigtes Material (zum Teil optional):

Reagenzien	Form/Menge	Bestell-Nr.
VARIO Ferro F10	Pulver / 100 St.	530560
VARIO Ferro F10	Pulver / 1000 St.	530563

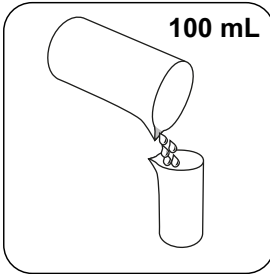
Vorbereitung

1. Eisenoxid erfordert vor der Analyse einen schwachen, starken oder Digesdahl Aufschluss (saurer Aufschlussverfahren).
2. Sehr stark alkalische oder saure Wässer sollten vor der Analyse auf einen pH-Wert zwischen 3 und 5 eingestellt werden.
3. Bei Proben die sichtbaren Rost enthalten sollte eine Reaktionszeit von mindestens 5 Minuten eingehalten werden.
4. Wässer, die mit organischen Verbindungen als Korrosionsschutz usw. behandelt worden sind, müssen ggf. oxidiert werden, um die Eisenkomplexe zu zerstören. Dazu wird eine 100 ml Probe mit 1 ml konzentrierter Schwefelsäure ($\geq 95\%$) und 1 ml konzentrierter Salpetersäure ($\geq 65\%$) versetzt und auf die Hälfte eingedampft. Nach dem Abkühlen wird der Aufschluss durchgeführt.

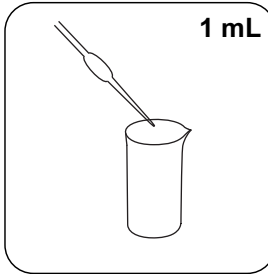
Anmerkungen

1. Bei dieser Methode erfolgt die Bestimmung von allen Formen gelösten Eisens und den meisten Formen von ungelöstem Eisen.
2. Die Genauigkeit wird durch ungelöstes Pulver nicht vermindert.

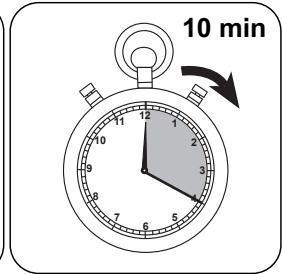
Aufschluss



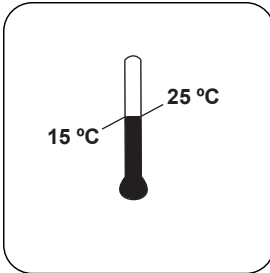
Ein geeignetes
Probengefäß mit **100 mL**
Probe füllen.



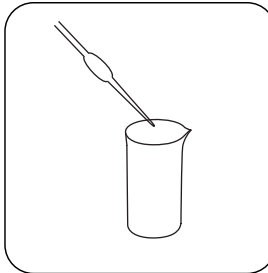
1 mL konzentrierte
Schwefelsäure (≥ 95 %)
zugeben.



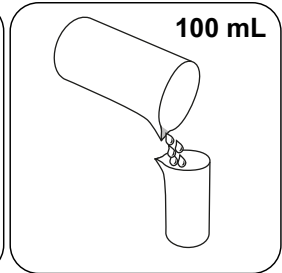
Die Probe für **10 Minuten**
erhitzen, oder so lange, bis
sich alles vollständig gelöst
hat.



Die Probe auf
Raumtemperatur
abkühlen lassen.



Den **pH-Wert** der Probe
mit **Ammoniaklösung**
(10-25 %) auf 3-5
einstellen.



Die Probe mit **VE-Wasser**
auf **100 mL** auffüllen.

Diese Probe für die Analyse von gesamten gelösten und ungelösten Eisen verwenden.

Durchführung der Bestimmung Eisen(II,III), gelöst mit Vario Pulverpäckchen

Die Methode im Gerät auswählen.

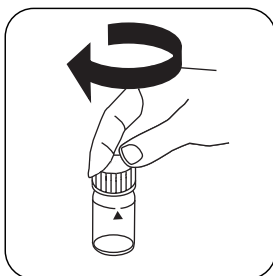
Für die Bestimmung von **Eisen mit Tablette** den beschriebenen **Aufschluss**
durchführen.



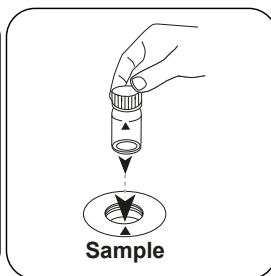
DE



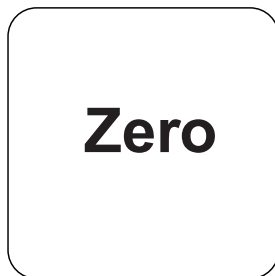
24-mm-Küvette mit **10 mL**
Probe füllen.



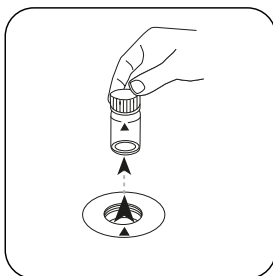
Küvette(n) verschließen.



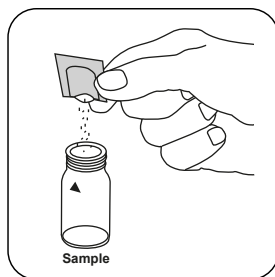
Die **Probeküvette** in
den Messsacht stellen.
Positionierung beachten.



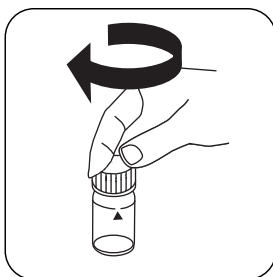
Taste **ZERO** drücken.



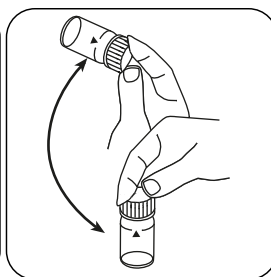
Küvette aus dem
Messsacht nehmen.



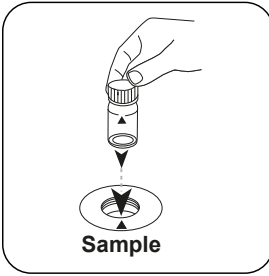
Ein **Vario FERRO**
F10 Pulverpäckchen
zugeben.



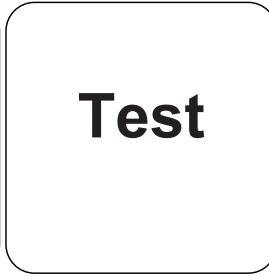
Küvette(n) verschließen.



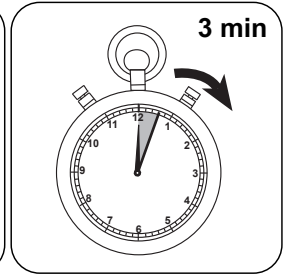
Inhalt durch Umschwenken
mischen.



Die **Probenküvette** in den Messschacht stellen. Positionierung beachten.



Taste **TEST** (XD: **START**) drücken.



3 Minute(n) Reaktionszeit abwarten.

Nach Ablauf der Reaktionszeit erfolgt automatisch die Messung.

In der Anzeige erscheint das Ergebnis in mg/L Eisen.



Chemische Methode

1,10-Phenanthroline

Appendix

DE

Störungen

Permanente Störungen

1. Iridium stört die Bestimmung.

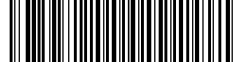
Gemäß

DIN 38406-E1

Standard Method 3500-Fe-1997

US EPA 40 CFR 136

⁹⁾ Reagenz erfasst die meisten Eisenoxide



Eisen (TPTZ) PP

M223

0,02 - 1,8 mg/L Fe

FE2

TPTZ

Material

DE

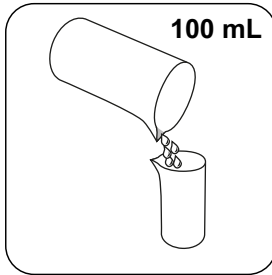
Benötigtes Material (zum Teil optional):

Reagenzien	Form/Menge	Bestell-Nr.
VARIO Iron TPTZ F10	Pulver / 100 St.	530550

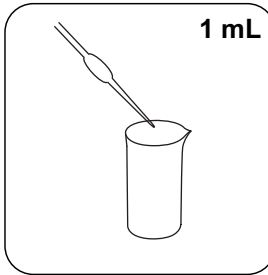
Vorbereitung

1. Für die Bestimmung von Gesamteisen ist ein Aufschluss erforderlich. Das TPTZ Reagenz erfasst die meisten Eisenoxide ohne Aufschluss.
2. Sämtliche Laborgläser vor der Analyse mit verdünnter Salzsäurelösung (1:1) und anschließend mit VE-Wasser spülen, um Eisenablagerungen zu beseitigen, die zu geringfügig höheren Ergebnissen führen können.
3. Stark alkalische oder saure Wässer sollten vor der Analyse in einen pH-Bereich zwischen 3 und 8 gebracht werden (mit 0,5 mol/l Schwefelsäure bzw. 1 mol/l Natronlauge).
4. Wässer, die mit organischen Verbindungen als Korrosionsschutz usw. behandelt worden sind, müssen ggf. oxidiert werden, um die Eisenkomplexe zu zerstören. Dazu wird eine 100 ml Probe mit 1 ml konzentrierter Schwefelsäure ($\geq 95\%$) und 1 ml konzentrierter Salpetersäure ($\geq 65\%$) versetzt und auf die Hälfte eingedampft. Nach dem Abkühlen wird der Aufschluss durchgeführt.

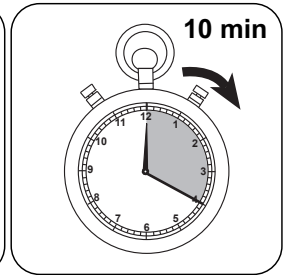
Aufschluss



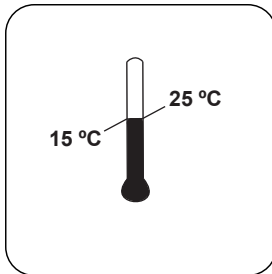
Ein geeignetes
Probengefäß mit **100 mL**
Probe füllen.



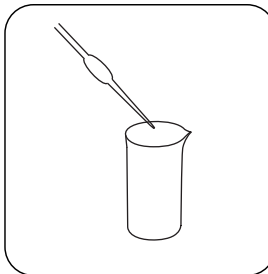
1 mL konzentrierte
Schwefelsäure (≥ 95 %)
zugeben.



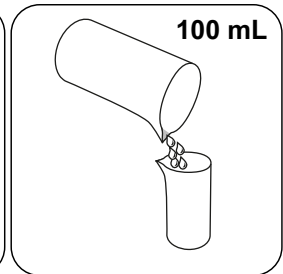
Die Probe für **10 Minuten**
erhitzen, oder so lange, bis
sich alles vollständig gelöst
hat.



Die Probe auf
Raumtemperatur
abkühlen lassen.



Den **pH-Wert** der Probe
mit **Ammoniaklösung**
(10-25 %) auf 3-5
einstellen.



Die Probe mit **VE-Wasser**
auf **100 mL** auffüllen.

Diese Probe für die Analyse von gesamten gelösten und ungelösten Eisen verwenden.

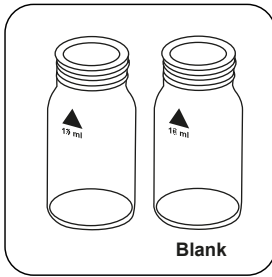
Durchführung der Bestimmung Eisen, gesamt mit Vario Pulverpäckchen

Die Methode im Gerät auswählen.

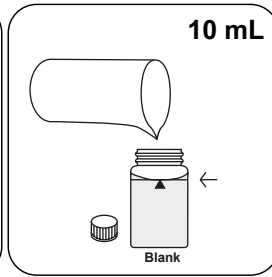
Für die Bestimmung von **Gesamteisen** den beschriebenen **Aufschluss** durchführen.



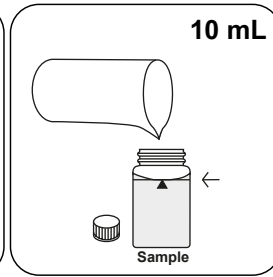
DE



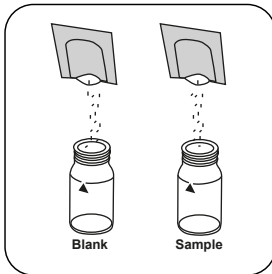
Zwei saubere 24-mL-Küvetten bereitstellen. Eine als Nullküvette kennzeichnen.



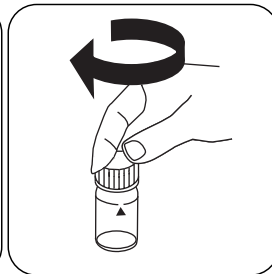
10 mL VE-Wasser in die Nullküvette geben.



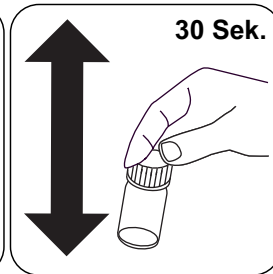
10 mL Probe in die Probenküvette geben.



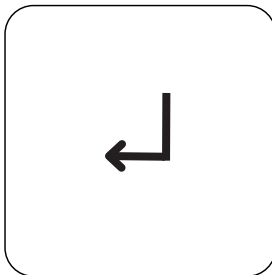
In jede Küvette ein **Vario IRON TPTZ F10 Pulverpäckchen** geben.



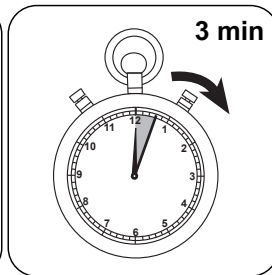
Küvette(n) verschließen.



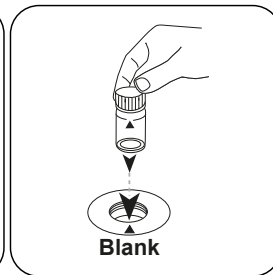
Inhalt durch Schütteln mischen (30 Sek.).



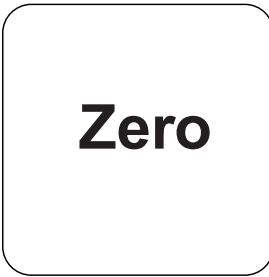
Taste **ENTER** drücken.



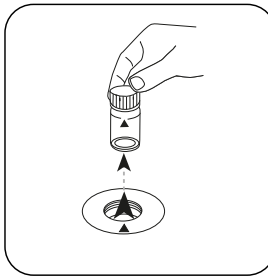
3 Minute(n) Reaktionszeit abwarten.



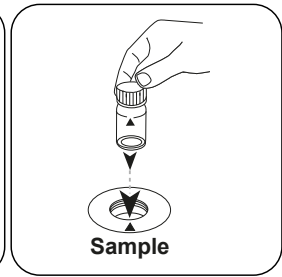
Die **Nullküvette** in den Messschacht stellen. Positionierung beachten.



Taste **ZERO** drücken.

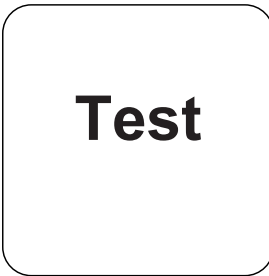


Küvette aus dem
Messschacht nehmen.



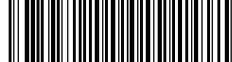
Die **Probenküvette** in
den Messschacht stellen.
Positionierung beachten.

DE



Taste **TEST** (XD: **START**)
drücken.

In der Anzeige erscheint das Ergebnis in mg/L Eisen.



Chemische Methode

TPTZ

Appendix

DE

Störungen


Permanente Störungen

Bei Auftretenden Störungen ist die Farbbildung gehemmt oder es bildet sich ein Niederschlag. Die Angaben beziehen sich auf einen Standard mit einer Eisenkonzentration von 0,5 mg/L.

Störung	Stört ab / [mg/L]
Cd	4
Cr ³⁺	0.25
Cr ⁶⁺	1.2
Co	0.05
Cu	0.6
CN ⁻	2.8
Mn	50
Hg	0.4
Mo	4
Ni	1
NO ₂ ⁻	0.8

Literaturverweise

G. Frederic Smith Chemical Co., The Iron Reagents, 3rd ed. (1980)

KS4.3 T / 20


Nombre del método

Número de método

Código de barras para reconocer el método

Rango de medición

20

S:4.3

Indicación en la pantalla de MD 100 / MD 110 / MD 200

Método químico

Información específica del instrumento

La prueba puede realizarse en los siguientes dispositivos. Además, se muestran la cubeta requerida y el rango de absorción del fotómetro.

Dispositivos	Cubeta	λ	Rango de medición
MD 200, MD 600, MD 610, MD 640, MultiDirect, PM 620, PM 630	\varnothing 24 mm	610 nm	0.1 - 4 mmol/l $K_{S4.3}$
SpectroDirect, XD 7000, XD 7500	\varnothing 24 mm	615 nm	0.1 - 4 mmol/l $K_{S4.3}$

Material

Material requerido (parcialmente opcional):

Título	Unidad de embalaje	Referencia No
Fotómetro alca-M	Tabletas / 100	513210BT
Fotómetro alca-M	Tabletas / 250	513211BT

Lista de aplicaciones

- Tratamiento de aguas residuales
- Tratamiento de aguas potables
- Tratamiento de aguas de aporte

Notas

1. Las definiciones de alcalinidad-m, valor-m y capacidad ácida $K_{S4.3}$ son idénticas.
2. Añadir un volumen de muestra de exactamente 10 ml, ya que este volumen influye de forma decisiva en la exactitud del resultado.

Códigos de idioma ISO 639-1

Estado de revisión

ES Manual de Métodos 01/20

Realización de la determinación

Ejecución de la determinación Capacidad ácida $K_{24.3}$ con tableta

Seleccionar el método en el aparato.

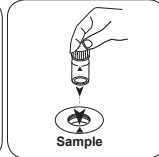
Para este método no es necesario realizar medición CERO en los aparatos siguientes: XD 7000, XD 7500



Llenar la cubeta de 24 mm con 10 ml de muestra .

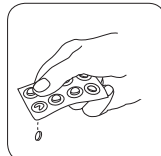


Cerrar la(s) cubeta(s).

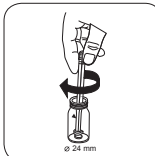


Poner la **cubeta de muestra** en el compartimiento de medición. ¡Debe tenerse en cuenta el posicionamiento!

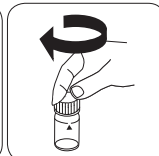
• • •



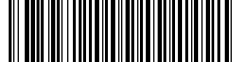
Añadir **tableta ALKA-M-PHOTOMETER**.



Triturar la(s) tableta(s) girando ligeramente.



Cerrar la(s) cubeta(s).



Hierro T

M220

0.02 - 1 mg/L Fe

FE

Ferrocina / Tioglicolato

ES

Material

Material requerido (parcialmente opcional):

Reactivos	Unidad de embalaje	No. de referencia
Hierro II LR (Fe ²⁺)	Tabletas / 100	515420BT
Hierro II LR (Fe ²⁺)	Tabletas / 250	515421BT
Hierro LR (Fe ²⁺ und Fe ³⁺)	Tabletas / 100	515370BT
Hierro LR (Fe ²⁺ und Fe ³⁺)	Tabletas / 250	515371BT

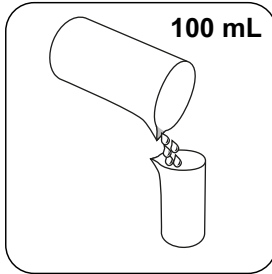
Preparación

1. Las aguas que han sido tratadas con compuestos orgánicos como protección contra la corrosión, etc., pueden oxidarse para destruir los complejos de hierro. Para ello se disuelve una muestra de 100 ml con 1 ml de ácido sulfúrico concentrado y 1 ml de ácido nítrico concentrado y se evapora a la mitad. Después de enfriarse se realiza la disgregación.

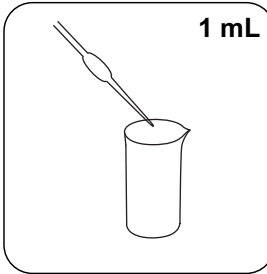
Notas

1. Con este método se realiza la determinación del Fe²⁺ y Fe³⁺ total disuelto.
2. Para determinar Fe²⁺ se utiliza la tableta IRON (II) LR, en lugar de la tableta IRON LR.

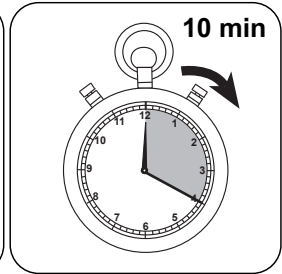
Disgregación



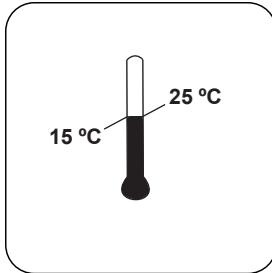
Llenar un recipiente de muestra apropiado con **100 mL de muestra**.



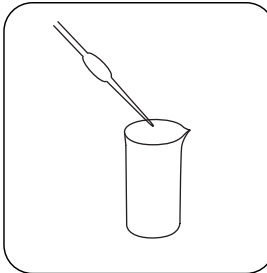
Añadir **1 mL de ácido sulfúrico concentrado**.



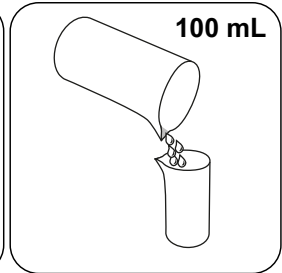
Calentar la muestra durante 10 minutos, o hasta que se haya disuelto totalmente.



Dejar enfriar la muestra a **temperatura ambiente**.



Ajustar el **valor de pH** de la muestra con **solución amoniacal a 3-5**.



Rellenar la muestra con **agua desionizada hasta 100 mL**.

Utilizar esta muestra para el análisis de total de hierro disuelto y no disuelto.

Ejecución de la determinación Hierro (II,III), disuelto con tableta

Seleccionar el método en el aparato.

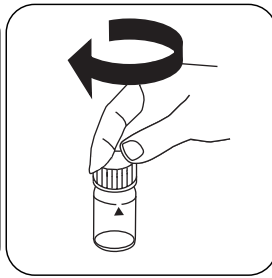
Para la determinación de **Hierro disuelto y sin disolver** realizar la disgregación descrita.



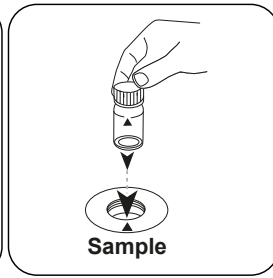
ES



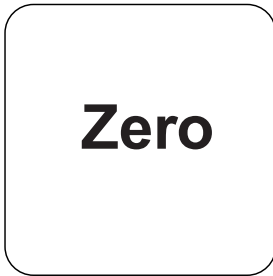
Llenar la cubeta de 24 mm con **10 mL de muestra** .



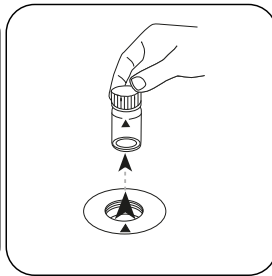
Cerrar la(s) cubeta(s).



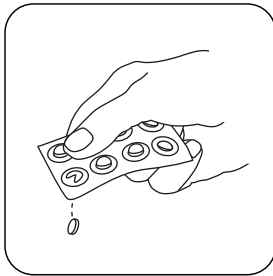
Poner la **cubeta de muestra** en el compartimiento de medición. ¡Debe tenerse en cuenta el posicionamiento!



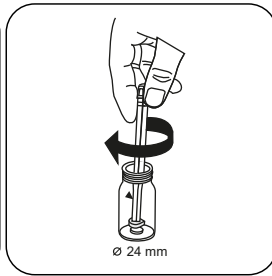
Pulsar la tecla **ZERO**.



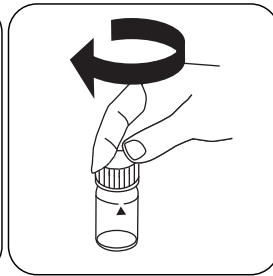
Extraer la cubeta del compartimiento de medición.



Añadir **tableta IRON LR**.



Triturar la(s) tableta(s) girando ligeramente.



Cerrar la(s) cubeta(s).



Disolver la(s) tableta(s) girando.

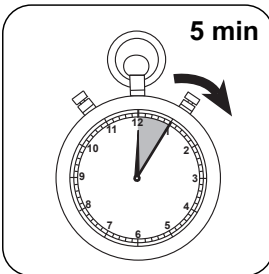


Poner la **cubeta de muestra** en el compartimiento de medición. ¡Debe tenerse en cuenta el posicionamiento!



Pulsar la tecla **TEST** (XD: **START**).

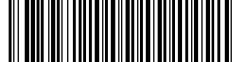
ES



Esperar **5 minutos como período de reacción**.

Finalizado el periodo de reacción se realizará la determinación automáticamente.

A continuación se visualizará el resultado en mg/L Hierro.



Método químico

Ferrocina / Tioglicolato

Apéndice

ES

Interferencia

Interferencias extraíbles

1. La presencia de cobre aumenta el resultado de medición un 10 %. Con una concentración de 10 mg/L de cobre en la muestra, el resultado de la medición se aumenta en 1 mg/L de hierro.
La perturbación puede eliminarse añadiendo tiourea.

Validación del método

Límite de detección	0.01 mg/L
Límite de determinación	0.016 mg/L
Límite del rango de medición	1 mg/L
Sensibilidad	0.92 mg/L / Abs
Intervalo de confianza	0.013 mg/L
Desviación estándar	0.005 mg/L
Coficiente de variación	1.23 %

Bibliografía

Photometrische Analyse, Lange/ Vjedelek, Verlag Chemie 1980, S. 102



Hierro PP

M222

0.02 - 3 mg/L Fe^g)

FE1

1,10-Fenantrolina

ES

Material

Material requerido (parcialmente opcional):

Reactivos	Unidad de embalaje	No. de referencia
Hierro F10 VARIO	Polvos / 100 Cantidad	530560
Hierro F10 VARIO	Polvos / 1000 Cantidad	530563

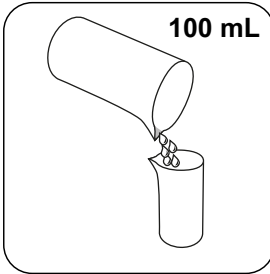
Preparación

1. El óxido de hierro necesita antes de la determinación una disgregación leve, fuerte o según Digesdahl (véase disgregación ácida).
2. Las muestras acuosas muy ácidas o muy básicas se han de diluir a un pH entre 3 y 5.
3. Las muestras que contengan óxidos visibles deberán mantener un periodo de reacción mínimo de 5 minutos.
4. Las aguas que han sido tratadas con compuestos orgánicos como protección contra la corrosión, etc., pueden oxidarse para destruir los complejos de hierro. Para ello se disuelve una muestra de 100 ml con 1 ml de ácido sulfúrico concentrado y 1 ml de ácido nítrico concentrado y se evapora a la mitad. Después de enfriarse se realiza la disgregación.

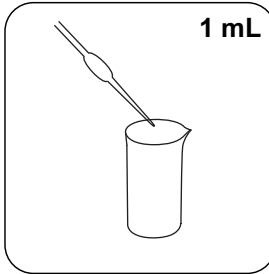
Notas

1. Con este método se realiza la determinación de todas las formas de hierro disuelto y la mayoría de hierro no disuelto.
2. Los polvos no disueltos no influyen en la exactitud del método.

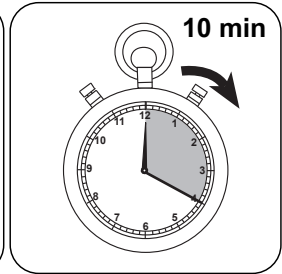
Disgregación



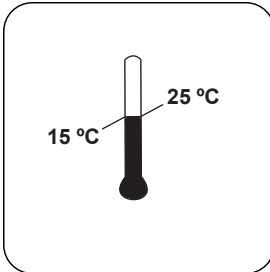
Llenar un recipiente de muestra apropiado con **100 mL de muestra**.



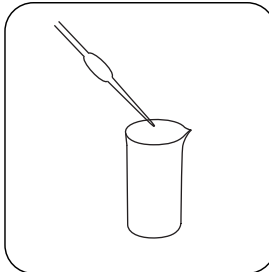
Añadir **1 mL de ácido sulfúrico concentrado**.



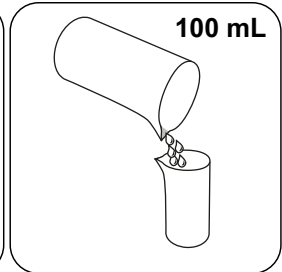
Calentar la muestra durante 10 minutos, o hasta que se haya disuelto totalmente.



Dejar enfriar la muestra a **temperatura ambiente**.



Ajustar el **valor de pH** de la muestra con **solución amoniacal a 3-5**.



Rellenar la muestra con **agua desionizada hasta 100 mL**.

Utilizar esta muestra para el análisis de total de hierro disuelto y no disuelto.

Ejecución de la determinación Hierro (II,III), disuelto con sobres de polvos Vario

Seleccionar el método en el aparato.

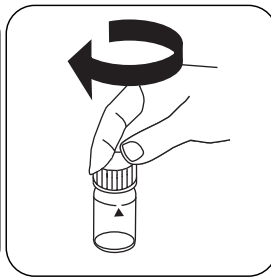
Para la determinación de **Hierro con tableta realizar la disgregación** descrita.



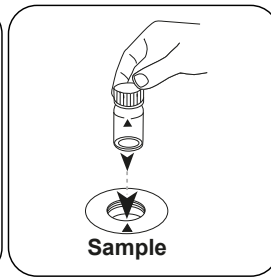
ES



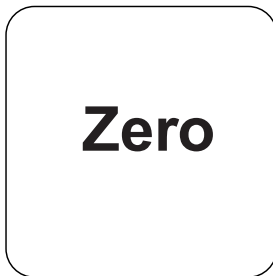
Llenar la cubeta de 24 mm con **10 mL de muestra** .



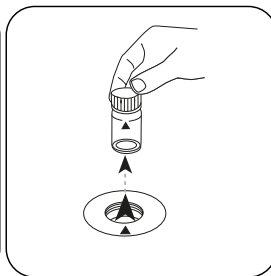
Cerrar la(s) cubeta(s).



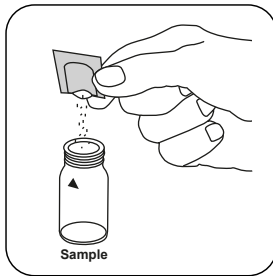
Poner la **cubeta de muestra** en el compartimiento de medición. ¡Debe tenerse en cuenta el posicionamiento!



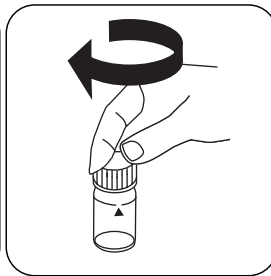
Pulsar la tecla **ZERO**.



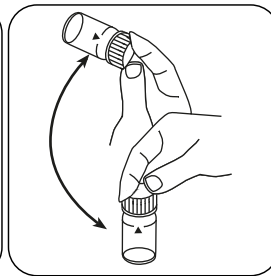
Extraer la cubeta del compartimiento de medición.



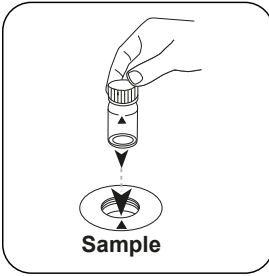
Añadir un **sobre de polvos Vario FERRO F10** .



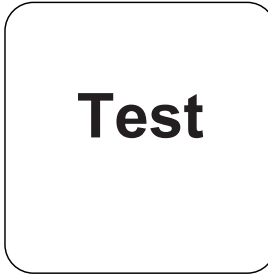
Cerrar la(s) cubeta(s).



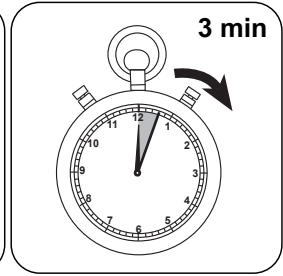
Mezclar el contenido girando.



Poner la **cupeta de muestra** en el compartimiento de medición. ¡Debe tenerse en cuenta el posicionamiento!



Pulsar la tecla **TEST** (XD: **START**).



Esperar **3 minutos como periodo de reacción**.

Finalizado el periodo de reacción se realizará la determinación automáticamente.
A continuación se visualizará el resultado en mg/L Hierro.



Método químico

1,10-Fenantrolina

Apéndice

ES

Interferencia

Interferencias persistentes

1. El iridio perturba la determinación.

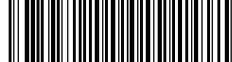
De acuerdo a

DIN 38406-E1

Método estándar 3500-Fe-1997

US EPA 40 CFR 136

⁹⁾ Reagente prende la mayor parte de los óxidos de hierro



Hierro (TPTZ) PP

M223

0.02 - 1.8 mg/L Fe

FE2

TPTZ

ES

Material

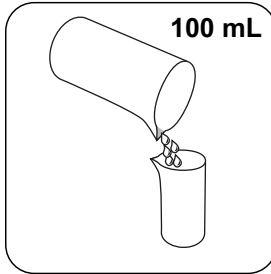
Material requerido (parcialmente opcional):

Reactivos	Unidad de embalaje	No. de referencia
Hierro TPTZ F10 VARIO	Polvos / 100 Cantidad	530550

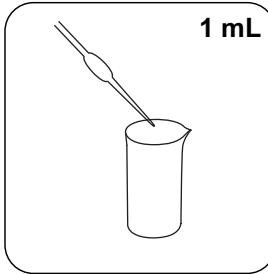
Preparación

1. Para la determinación del hierro total es necesaria una disgregación. El reactivo TPTZ registra la mayoría de los óxidos de hierro sin disgregación.
2. Para eliminar residuos férricos que pueden producir resultados más elevados, lavar todos los aparatos antes de su uso con una solución de ácido clorhídrico (1:1) enjuagándolos a continuación con agua desionizada.
3. Las muestras acuosas muy ácidas o muy básicas se deberán neutralizar a un valor de pH entre 3 y 8 antes de realizar el análisis (con 0,5 mol/l de ácido sulfúrico o 1 mol/l de hidróxido sódico).
4. Las aguas que han sido tratadas con compuestos orgánicos como protección contra la corrosión, etc., pueden oxidarse para destruir los complejos de hierro. Para ello se disuelve una muestra de 100 ml con 1 ml de ácido sulfúrico concentrado y 1 ml de ácido nítrico concentrado y se evapora a la mitad. Después de enfriarse se realiza la disgregación.

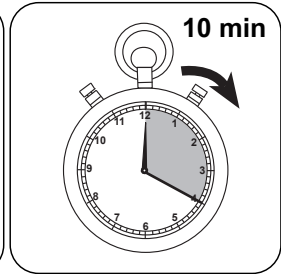
Disgregación



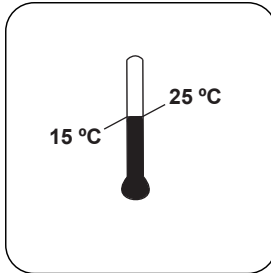
Llenar un recipiente de muestra apropiado con **100 mL de muestra**.



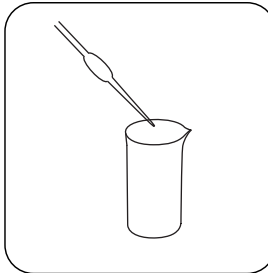
Añadir **1 mL de ácido sulfúrico concentrado**.



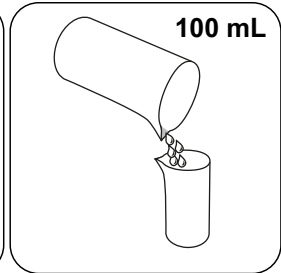
Calentar la muestra durante 10 minutos, o hasta que se haya disuelto totalmente.



Dejar enfriar la muestra a **temperatura ambiente**.



Ajustar el **valor de pH** de la muestra con **solución amoniacal a 3-5**.



Rellenar la muestra con **agua desionizada hasta 100 mL**.

Utilizar esta muestra para el análisis de total de hierro disuelto y no disuelto.

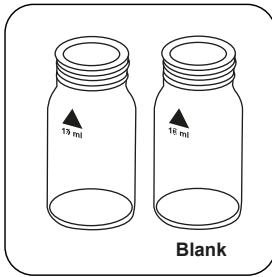
Ejecución de la determinación Hierro, total con sobres de polvos Vario

Seleccionar el método en el aparato.

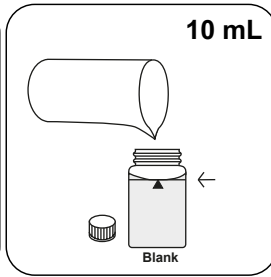
Para la determinación de **Hierro total realizar la disgregación** descrita.



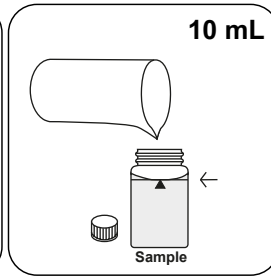
ES



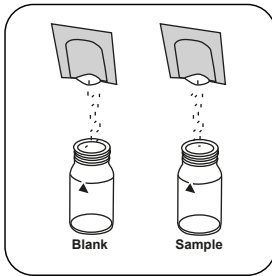
Preparar dos cubetas limpias de 24 mm. Identificar una como cubeta en blanco.



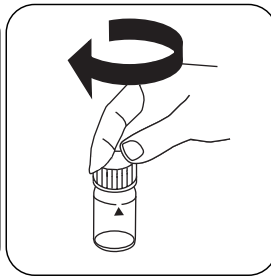
Añadir **10 mL de agua desionizada** en la cubeta en blanco.



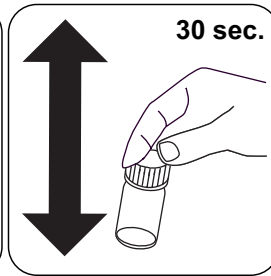
Añadir **10 mL de muestra** en la cubeta con la muestra.



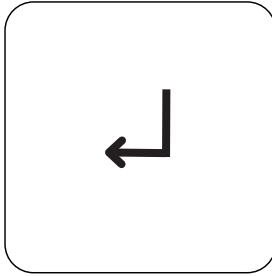
Añadir un sobre de polvos de Vario IRON TPTZ F10 en cada cubeta.



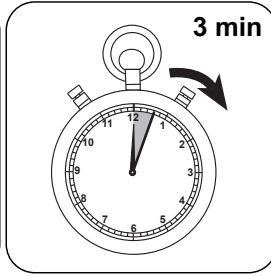
Cerrar la(s) cubeta(s).



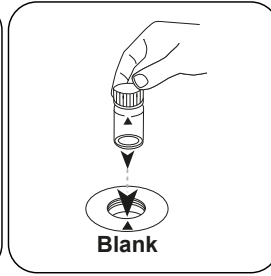
Mezclar el contenido agitando (30 sec.).



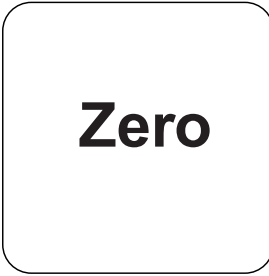
Pulsar la tecla **ENTER**.



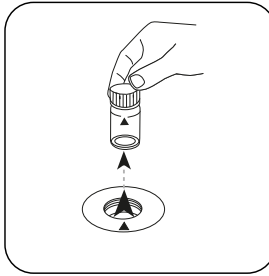
Esperar **3 minutos como periodo de reacción**.



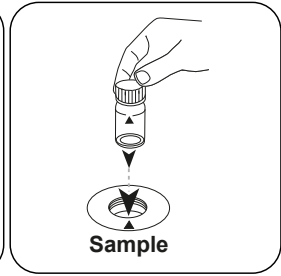
Poner la **cubeta en blanco** en el compartimiento de medición. ¡Debe tenerse en cuenta el posicionamiento!



Pulsar la tecla **ZERO**.

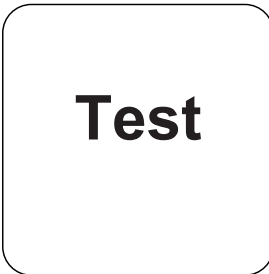


Extraer la cubeta del compartimiento de medición.



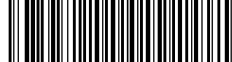
Poner la **cubeta de muestra** en el compartimiento de medición. ¡Debe tenerse en cuenta el posicionamiento!

ES



Pulsar la tecla **TEST** (XD: **START**).

A continuación se visualizará el resultado en mg/L Hierro.



Método químico

TPTZ

Apéndice

ES

Interferencia

Interferencias persistentes


Si se producen perturbaciones se inhibe la coloración o se forma precipitación. Los datos se refieren a un estándar con una concentración de hierro de 0,5 mg/L.

Interferencia	de / [mg/L]
Cd	4
Cr ³⁺	0.25
Cr ⁶⁺	1.2
Co	0.05
Cu	0.6
CN ⁻	2.8
Mn	50
Hg	0.4
Mo	4
Ni	1
NO ₂ ⁻	0.8

Bibliografía

G. Frederic Smith Chemical Co., The Iron Reagents, 3rd ed. (1980)

KS4.3 T / 20



Nom de la méthode → KS4.3 T

Numéro de méthode → 20

Code à barres pour reconnaître la méthode → [Barcode]

Plage de mesure → 0.1 - 4 mmol/l $K_{S4.3}$

Méthode chimique → Acide / Indicateur

Affichage dans le MD 100 / MD 110 / MD 200 → S:4.3

Informations spécifiques à l'instrument

Le test peut être effectué sur les appareils suivants. De plus, la cuvette requise et la plage d'absorption du photomètre sont indiquées.

Appareils	Cuvette	λ	Gamme de mesure
MD 200, MD 600, MD 610, MD 640, MultiDirect, PM 620, PM 630	ø 24 mm	610 nm	0.1 - 4 mmol/l $K_{S4.3}$
SpectroDirect, XD 7000, XD 7500	ø 24 mm	615 nm	0.1 - 4 mmol/l $K_{S4.3}$

Matériel

Matériel requis (partiellement optionnel):

Titre	Pack contenant	Code
Alka-M-Photometer	Pastilles / 100	513210BT
Alka-M-Photometer	Pastilles / 250	513211BT

Liste d'applications

- Traitement des eaux usées
- Traitement de l'eau potable
- Traitement de l'eau brute

Indication

1. Les termes Alcalinité-m, Valeur m, Alcalinité totale et Capacité acide $K_{S4.3}$ sont identiques.
2. L'observation exacte du volume d'échantillon de 10 ml est décisive pour l'exactitude du résultat de l'analyse.

Codes de langue ISO 639-1 → FR

État de révision → 01/20

FR Méthodes Manuel 01/20

Procédure du test

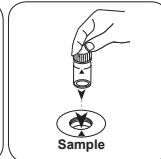
Réalisation de la quantification Capacité acide $K_{s4.3}$ avec pastille

Sélectionnez la méthode sur l'appareil.

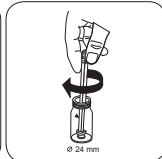
Cette méthode ne nécessite aucune mesure du zéro sur les appareils suivants : XD 7000, XD 7500

Remplissez une cuvette de 24 mm de **10 ml d'échantillon**.

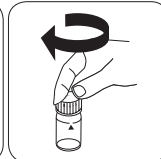
Fermez la(les) cuvette(s).

Placez la **cuvette réservée à l'échantillon** dans la chambre de mesure. Attention à la positionner correctement.

• • •

Ajoutez une **pastille de ALKA-M-PHOTOMETER**.

Écrasez la(les) pastille(s) en la(les) tournant un peu.



Fermez la(les) cuvette(s).



Fer T

M220

0.02 - 1 mg/L Fe

FE

Ferrozine / Thioglycolate

FR

Matériel

Matériel requis (partiellement optionnel):

Réactifs	Pack contenant	Code
Fer II LR (Fe ²⁺)	Pastilles / 100	515420BT
Fer II LR (Fe ²⁺)	Pastilles / 250	515421BT
Fer LR (Fe ²⁺ und Fe ³⁺)	Pastilles / 100	515370BT
Fer LR (Fe ²⁺ und Fe ³⁺)	Pastilles / 250	515371BT

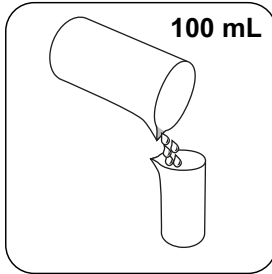
Préparation

1. Il faudra éventuellement oxyder les eaux auparavant traitées avec des composés organiques pour les protéger de la corrosion etc. afin de détruire les complexes du fer. À un échantillon de 100 ml, on ajoutera 1 ml d'acide sulfurique concentré et 1 ml d'acide nitrique concentré pour réduire le tout de moitié par évaporation. Le fractionnement est effectué après le refroidissement.

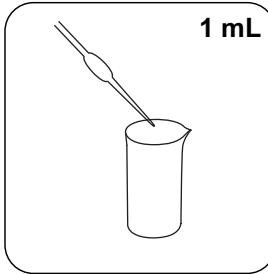
Indication

1. Cette méthode permet de quantifier le Fe²⁺ total dissous et Fe³⁺.
2. Pour la quantification du Fe²⁺, la pastille IRON LR est remplacée par la pastille IRON (II) LR.

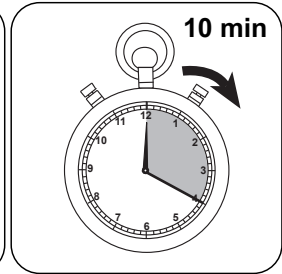
Fractionnement



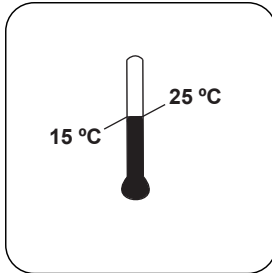
Versez **100 mL d'échantillon** dans un tube pour échantillon adéquat.



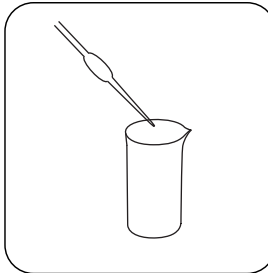
Ajoutez **1 mL de d'acide sulfurique concentré**.



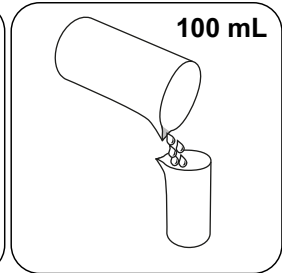
Réchauffez l'échantillon pendant **10 minutes**, ou jusqu'à ce que tout soit entièrement dissous.



Laissez refroidir l'échantillon à **température ambiante**.



Réglez le **pH** de l'échantillon avec **solution ammoniacquée** pour obtenir 3-5.



Complétez l'échantillon en ajoutant **d'eau déminéralisée** pour obtenir **100 mL**.

Utilisez cet échantillon pour analyser Fer total résolu et dissous.

Réalisation de la quantification Fer (II,III), dissous avec pastille

Sélectionnez la méthode sur l'appareil.

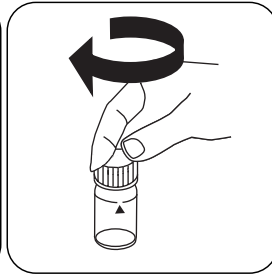
Pour la quantification de **Fer dissous et non dissous**, procédez au fractionnement décrit.



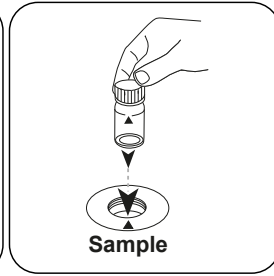
FR



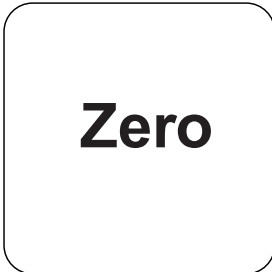
Remplissez une cuvette de 24 mm de **10 mL d'échantillon**.



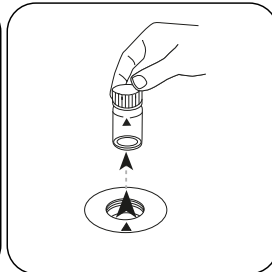
Fermez la(les) cuvette(s).



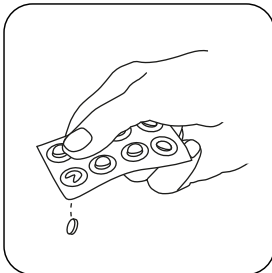
Placez la **cuvette réservée à l'échantillon** dans la chambre de mesure. Attention à la positionner correctement.



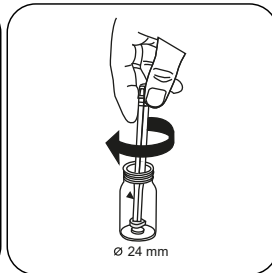
Appuyez sur la touche **ZERO**.



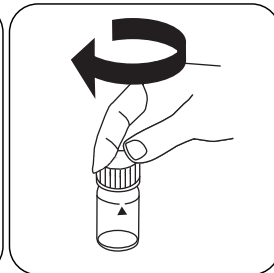
Retirez la cuvette de la chambre de mesure.



Ajoutez une **pastille de IRON LR**.



Écrasez la(les) pastille(s) en la(les) tournant un peu.



Fermez la(les) cuvette(s).



Dissolvez la(les) pastille(s) en mettant le tube plusieurs fois à l'envers.

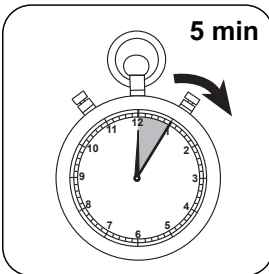


Placez la **cuvette réservée à l'échantillon** dans la chambre de mesure. Attention à la positionner correctement.



Appuyez sur la touche **TEST** (XD: **START**).

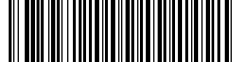
FR



Attendez la fin du **temps de réaction de 5 minute(s)**.

À l'issue du temps de réaction, la mesure est effectuée automatiquement.

Le résultat s'affiche à l'écran en mg/L fer.



Méthode chimique

Ferrozine / Thioglycolate

Appendice

FR

Interférences

Interférences exclues

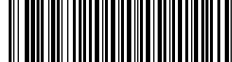
1. La présence de cuivre augmente le résultat de 10 %. À une concentration de cuivre égale à 10 mg/L dans l'échantillon, le résultat augmente de 1 mg/L de fer. La perturbation peut être éliminée par un apport de thiocarbamide.

Méthode Validation

Limite de détection	0.01 mg/L
Limite de détermination	0.016 mg/L
Fin de la gamme de mesure	1 mg/L
Sensibilité	0.92 mg/L / Abs
Intervalle de confiance	0.013 mg/L
Déviatoin standard	0.005 mg/L
Coefficient de variation	1.23 %

Bibliographie

Photométrische Analyse, Lange/ Vjedelek, Verlag Chemie 1980, S. 102



Fer PP

M222

0.02 - 3 mg/L Fe⁹⁾

FE1

1,10-Phénanthroline

FR

Matériel

Matériel requis (partiellement optionnel):

Réactifs	Pack contenant	Code
VARIO Ferro F10	Poudre / 100 Pièces	530560
VARIO Ferro F10	Poudre / 1000 Pièces	530563

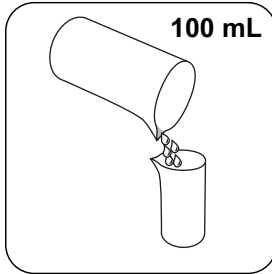
Préparation

1. Avant de pouvoir être analysé, l'oxyde de fer nécessite un fractionnement faible, fort ou Digesdahl (procédure de fractionnement acide).
2. Avant l'analyse, les eaux fortement alcalines ou acides devraient être ajustées sur un pH compris entre 3 et 5.
3. Pour les échantillons ayant des traces visibles de rouille, il est recommandé de respecter un temps de réaction d'au moins 5 minutes.
4. Il faudra éventuellement oxyder les eaux auparavant traitées avec des composés organiques pour les protéger de la corrosion etc. afin de détruire les complexes du fer. À un échantillon de 100 ml, on ajoutera 1 ml d'acide sulfurique concentré et 1 ml d'acide nitrique concentré pour réduire le tout de moitié par évaporation. Le fractionnement est effectué après le refroidissement.

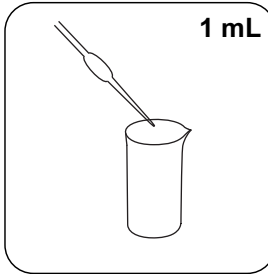
Indication

1. Cette méthode permet de quantifier toutes les formes de fer dissous et la plupart des formes de fer non dissous.
2. L'exactitude n'est pas diminuée par de la poudre non dissoute.

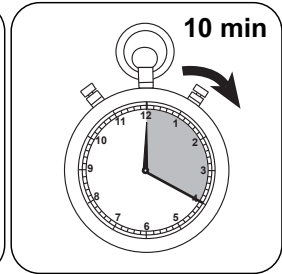
Fractionnement



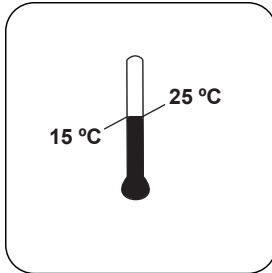
Versez **100 mL d'échantillon** dans un tube pour échantillon adéquat.



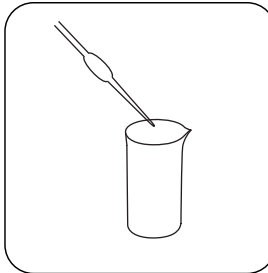
Ajoutez **1 mL de d'acide sulfurique concentré**.



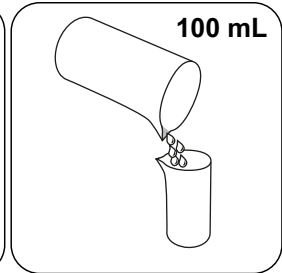
Réchauffez l'échantillon pendant **10 minutes**, ou jusqu'à ce que tout soit entièrement dissous.



Laissez refroidir l'échantillon à **température ambiante**.



Régalez le **pH** de l'échantillon avec **solution ammoniacuée** pour obtenir 3-5.



Complétez l'échantillon en ajoutant **d'eau déminéralisée** pour obtenir **100 mL**.

Utilisez cet échantillon pour analyser Fer total résolu et dissous.

Réalisation de la quantification Fer (II,III), dissous avec sachet de poudre Vario

Sélectionnez la méthode sur l'appareil.

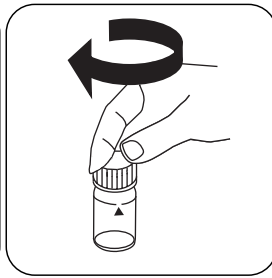
Pour la quantification de **Fer avec pastille**, procédez au fractionnement décrit.



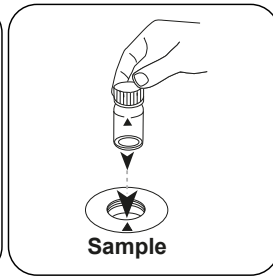
FR



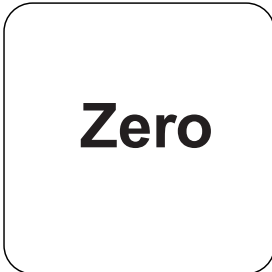
Remplissez une cuvette de 24 mm de **10 mL d'échantillon**.



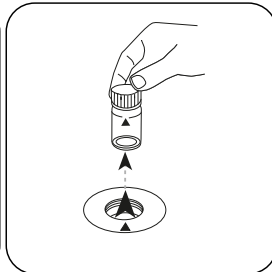
Fermez la(les) cuvette(s).



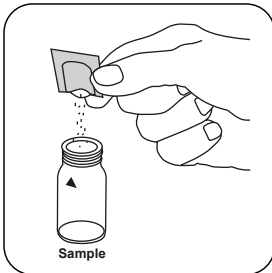
Placez la **cuvette réservée à l'échantillon** dans la chambre de mesure. Attention à la positionner correctement.



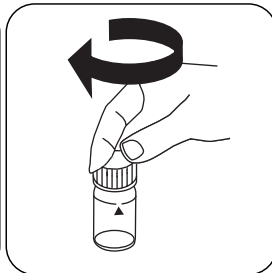
Appuyez sur la touche **ZERO**.



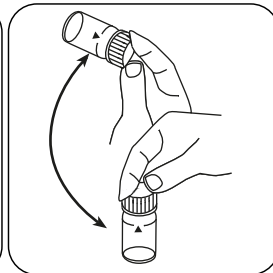
Retirez la cuvette de la chambre de mesure.



Ajoutez un **sachet de poudre Vario FERRO F10**.



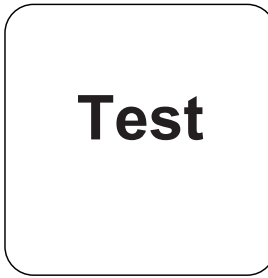
Fermez la(les) cuvette(s).



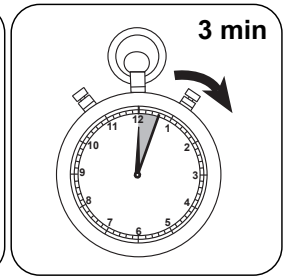
Mélangez le contenu en mettant le tube plusieurs fois à l'envers puis à l'endroit.



Placez la **cuvette réservée à l'échantillon** dans la chambre de mesure. Attention à la positionner correctement.

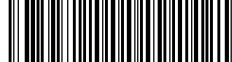


Appuyez sur la touche **TEST** (XD: **START**).



Attendez la fin du **temps de réaction de 3 minute(s)**.

À l'issue du temps de réaction, la mesure est effectuée automatiquement. Le résultat s'affiche à l'écran en mg/L fer.



Méthode chimique

1,10-Phénanthroline

Appendice

FR

Interférences

Interférences persistantes

1. L'iridium perturbe la quantification.

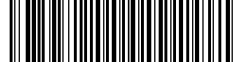
Selon

DIN 38406-E1

Standard Method 3500-Fe-1997

US EPA 40 CFR 136

⁹⁾Ce réactif réagit à la plupart des oxydes de fer



Fer (TPTZ) PP

M223

0.02 - 1.8 mg/L Fe

FE2

TPTZ

FR

Matériel

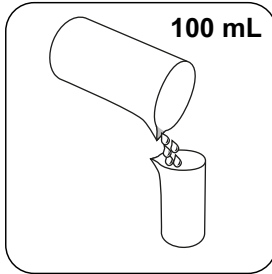
Matériel requis (partiellement optionnel):

Réactifs	Pack contenant	Code
VARIO Fer TPTZ F10	Poudre / 100 Pièces	530550

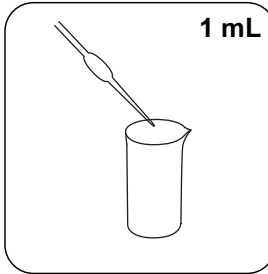
Préparation

1. La quantification du fer total nécessite un fractionnement. Le réactif TPTZ détecte la plupart des oxydes de fer sans fractionnement.
2. Avant l'analyse, lavez tous les instruments en verre en utilisant une solution d'acide chlorhydrique diluée (1:1) puis rincez-les à l'eau déminéralisée afin d'éliminer les dépôts de fer susceptibles d'augmenter légèrement les résultats.
3. Avant l'analyse, les eaux fortement alcalines ou acides devraient être ajustées sur un pH compris entre 3 et 8 (avec 0,5 mol/l d'acide sulfurique ou 1 mol/l de soude caustique).
4. Il faudra éventuellement oxyder les eaux auparavant traitées avec des composés organiques pour les protéger de la corrosion etc. afin de détruire les complexes du fer. À un échantillon de 100 ml, on ajoutera 1 ml d'acide sulfurique concentré et 1 ml d'acide nitrique concentré pour réduire le tout de moitié par évaporation. Le fractionnement est effectué après le refroidissement.

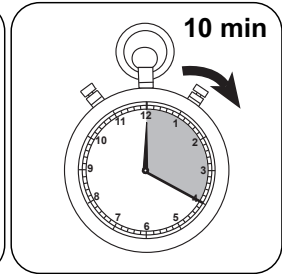
Fractionnement



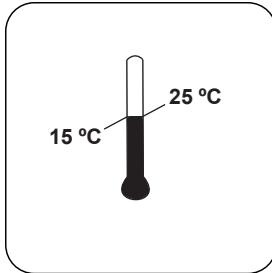
Versez **100 mL** d'échantillon dans un tube pour échantillon adéquat.



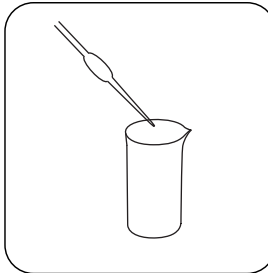
Ajoutez **1 mL de d'acide sulfurique concentré**.



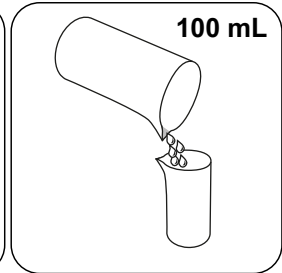
Réchauffez l'échantillon pendant **10 minutes**, ou jusqu'à ce que tout soit entièrement dissous.



Laissez refroidir l'échantillon à **température ambiante**.



Régalez le **pH** de l'échantillon avec **solution ammoniacquée** pour obtenir 3-5.



Complétez l'échantillon en ajoutant **d'eau déminéralisée** pour obtenir **100 mL**.

Utilisez cet échantillon pour analyser Fer total résolu et dissous.

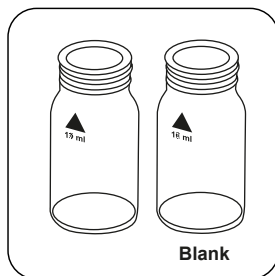
Réalisation de la quantification Fer, total avec sachet de poudre Vario

Sélectionnez la méthode sur l'appareil.

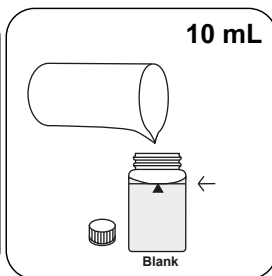
Pour la quantification de **Fer total**, procédez au fractionnement décrit.



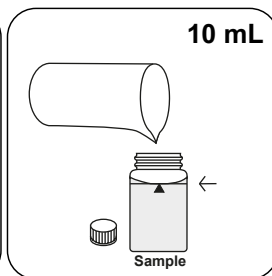
FR



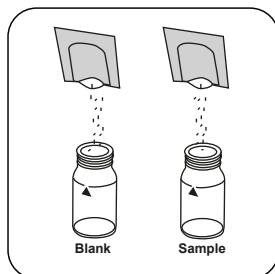
Préparez deux cuvettes propres de 24 mm. L'une des deux cuvettes sera la cuvette du blanc. Étiquetez-la.



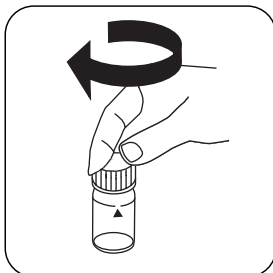
Versez **10 mL d'eau déminéralisée** dans la cuvette du blanc.



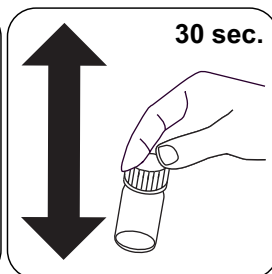
Versez **10 mL d'échantillon** dans la cuvette réservée à l'échantillon.



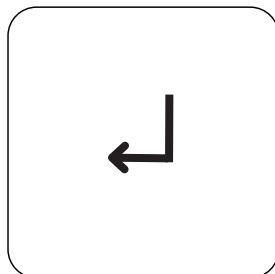
Dans chaque cuvette, versez **un sachet de poudre Vario IRON TPTZ F10**.



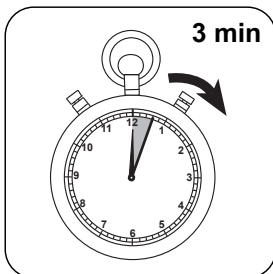
Fermez la(les) cuvette(s).



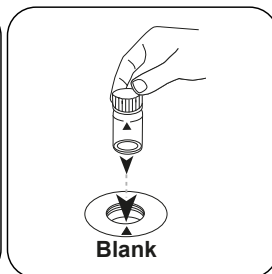
Mélangez le contenu en agitant (30 sec.).



Appuyez sur la touche **ENTER**.



Attendez la fin du **temps de réaction de 3 minute(s)**.

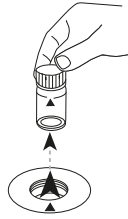


Placez la **cuvette du blanc** dans la chambre de mesure. Attention à la positionner correctement.

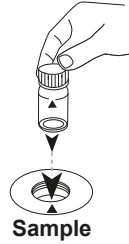


Zero

Appuyez sur la touche **ZERO**.



Retirez la cuvette de la chambre de mesure.



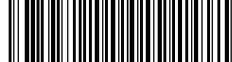
Placez la **cuvette réservée à l'échantillon** dans la chambre de mesure. Attention à la positionner correctement.

FR

Test

Appuyez sur la touche **TEST** (XD: **START**).

Le résultat s'affiche à l'écran en mg/L fer.



Méthode chimique

TPTZ

Appendice

FR

Interférences

Interférences persistantes

En cas de perturbation, la coloration est bloquée ou il se forme un précipité. Les indications s'appuient sur un étalon caractérisé par une concentration de fer égale à 0,5 mg/L.

Interférences	de / [mg/L]
Cd	4
Cr ³⁺	0.25
Cr ⁶⁺	1.2
Co	0.05
Cu	0.6
CN ⁻	2.8
Mn	50
Hg	0.4
Mo	4
Ni	1
NO ₂ ⁻	0.8

Bibliographie

G. Frederic Smith Chemical Co., The Iron Reagents, 3rd ed. (1980)

KS4.3 T / 20

Denominazione metodo

Numero metodo

Codice a barre per riconoscere il metodo

Range di misura

Indicazione sul display del MD 100 / MD 110 / MD 200

Metodo chimico

Informazioni specifiche dello strumento

Il test può essere eseguito sui seguenti dispositivi. Inoltre, sono indicate la cuvetta richiesta e il range di assorbimento del fotometro.

Dispositivi	Cuvetta	λ	Campo di misura
MD 200, MD 600, MD 610, MD 640, MultiDirect, PM 620, PM 630	ø 24 mm	610 nm	0.1 - 4 mmol/l $K_{S_{4.3}}$
SpectroDirect, XD 7000, XD 7500	ø 24 mm	615 nm	0.1 - 4 mmol/l $K_{S_{4.3}}$

Materiale

Materiale richiesto (in parte facoltativo):

Titolo	Unità di imballaggio	N. ordine
Alka-M-Photometer	Pastiglia / 100	513210BT
Alka-M-Photometer	Pastiglia / 250	513211BT

Campo di applicazione

- Trattamento acqua di scarico
- Trattamento acqua potabile
- Trattamento acqua non depurata

Note

1. I termini alcalinità M, valore M, alcalinità totale e capacità acida $K_{S_{4.3}}$ sono equivalenti.
2. Per l'accuratezza del risultato dell'analisi è fondamentale che il volume del campione misuri esattamente 10 ml.

ISO 639-1 codici linguistici

Stato di revisione

IT Manuale dei Metodi 01/20

**Svolgimento della
misurazione**

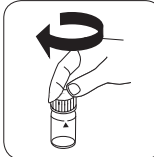
Esecuzione della rilevazione Capacità acida $K_{s4,3}$ con pastiglia

Selezionare il metodo nel dispositivo.

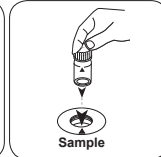
Con i seguenti dispositivi, per questo metodo non è necessario eseguire una misurazione ZERO: XD 7000, XD 7500



Riempire una cuvetta da 24 mm con **10 ml di campione**.

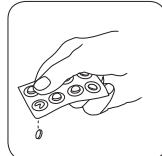


Chiudere la/e cuvetta/e.

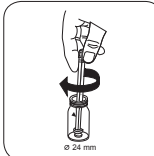


Posizionare la **cuvetta del campione** nel vano di misurazione. Fare attenzione al posizionamento.

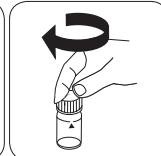
• • •



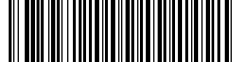
Aggiungere una **pastiglia ALKA-M-PHOTOMETER**.



Frantumare la/e pastiglia/e con una leggera rotazione.



Chiudere la/e cuvetta/e.



Ferro T

M220

0.02 - 1 mg/L Fe

FE

Ferrozine / acido tioglicolico

IT

Materiale

Materiale richiesto (in parte facoltativo):

Reagenti	Unità di imballaggio	N. ordine
Ferro II LR (Fe ²⁺)	Pastiglia / 100	515420BT
Ferro II LR (Fe ²⁺)	Pastiglia / 250	515421BT
Ferro LR (Fe ²⁺ und Fe ³⁺)	Pastiglia / 100	515370BT
Ferro LR (Fe ²⁺ und Fe ³⁺)	Pastiglia / 250	515371BT

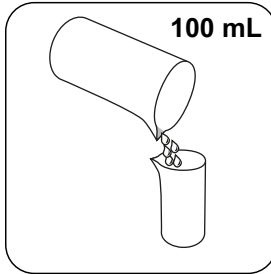
Preparazione

1. Le acque che sono state trattate con composti organici che proteggono dalla corrosione devono essere eventualmente ossidate per disgregare i complessi di ferro. A tale scopo si aggiunge un campione da 100 ml con 1 ml di acido solforico concentrato e 1 ml di acido nitrico concentrato e lo si fa evaporare fino alla metà. Dopo il raffreddamento viene eseguita la digestione.

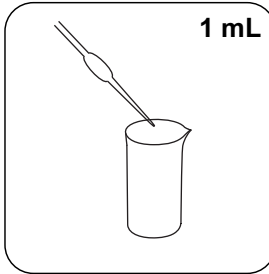
Note

1. Con questo metodo viene rilevato il totale del Fe²⁺ e del Fe³⁺ disciolti.
2. Per rilevare il Fe²⁺ si utilizza, invece della pastiglia IRON LR, la pastiglia IRON (II) LR.

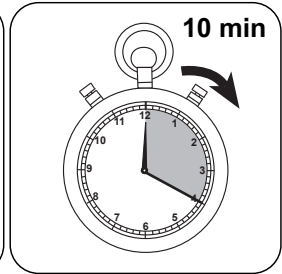
Digestione



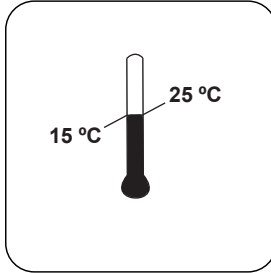
Riempire un recipiente per campioni adeguato con **100 mL di campione**.



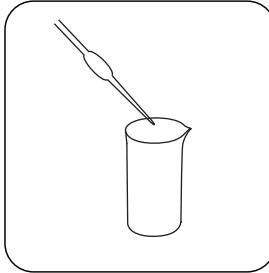
Aggiungere **1 mL di acido solforico concentrato**.



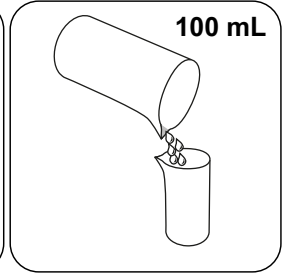
Riscaldare il campione per 10 minuti o finché non si sarà sciolto completamente.



Lasciar raffreddare il campione a **temperatura ambiente**.



Regolare il **valore di pH** del campione con **soluzione di ammoniaca su 3-5**.



Aggiungere al campione **acqua demineralizzata fino a raggiungere i 100 mL**.

Utilizzare questo campione per l'analisi di Ferro soluto e disciolto totale.

Esecuzione della rilevazione Ferro(II,III), disciolto con pastiglia

Selezionare il metodo nel dispositivo.

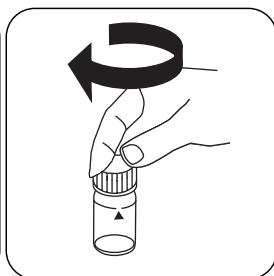
Per la determinazione di **Ferro disciolto e non disciolto** eseguire la **digestione** descritta.



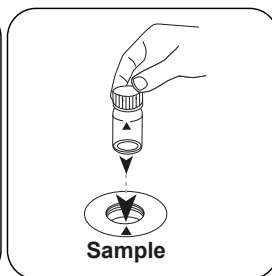
IT



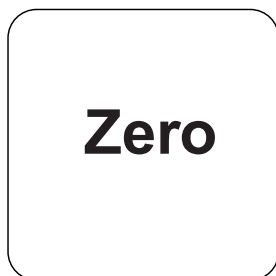
Riempire una cuvetta da 24 mm con **10 mL di campione**.



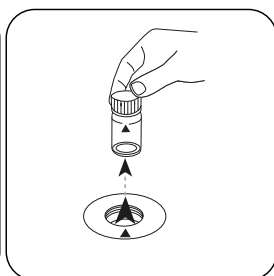
Chiudere la/e cuvetta/e.



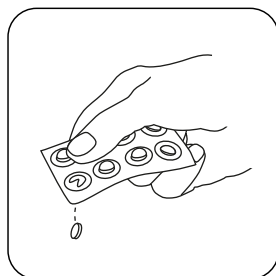
Posizionare la **cuvetta del campione** nel vano di misurazione. Fare attenzione al posizionamento.



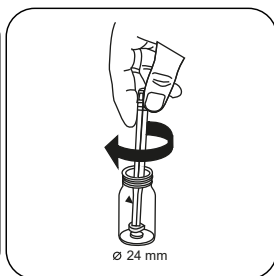
Premere il tasto **ZERO**.



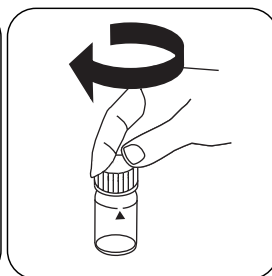
Prelevare la cuvetta dal vano di misurazione.



Aggiungere **una pastiglia IRON LR**.



Frantumare la/e pastiglia/e con una leggera rotazione.



Chiudere la/e cuvetta/e.



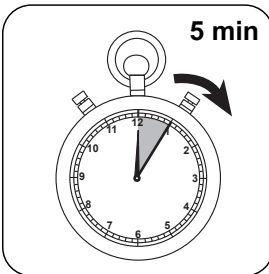
Far sciogliere la/e pastiglia/e agitando.



Posizionare la **cuvetta del campione** nel vano di misurazione. Fare attenzione al posizionamento.



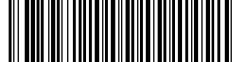
Premere il tasto **TEST** (XD: **START**).



Attendere un **tempo di reazione di 5 minuti/i** .

Allo scadere del tempo di reazione viene effettuata automaticamente la misurazione.

Sul display compare il risultato in mg/L di Ferro.



Metodo chimico

Ferrozine / acido tioglicolico

Appendice

IT

Interferenze

Interferenze escludibili

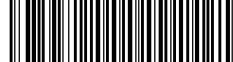
1. La presenza di rame aumenta il risultato della misurazione del 10 %. Con una concentrazione di 10 mg/L di rame nel campione il risultato della misurazione viene aumentato di 1 mg/L di ferro.
L'interferenza può essere eliminata con l'aggiunta di tiourea.

Validazione metodo

Limite di rilevabilità	0.01 mg/L
Limite di quantificazione	0.016 mg/L
Estremità campo di misura	1 mg/L
Sensibilità	0.92 mg/L / Abs
Intervallo di confidenza	0.013 mg/L
Deviazione standard della procedura	0.005 mg/L
Coefficiente di variazione della procedura	1.23 %

Riferimenti bibliografici

Photometrische Analyse, Lange/Vjedelek, Verlag Chemie 1980, pag. 102



Ferro PP

M222

0.02 - 3 mg/L Fe⁹⁾

FE1

1,10-fenantrolina

IT

Materiale

Materiale richiesto (in parte facoltativo):

Reagenti	Unità di imballaggio	N. ordine
VARIO Ferro F10	Polvere / 100 pz.	530560
VARIO Ferro F10	Polvere / 1000 pz.	530563

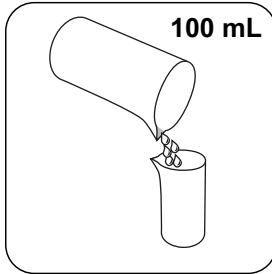
Preparazione

1. Prima dell'analisi, l'ossido di ferro richiede una digestione delicata, vigorosa o Digesdahl (processo di digestione acida).
2. Le acque fortemente alcaline o acide dovrebbero essere regolate prima dell'analisi su un valore di pH compreso tra 3 e 5.
3. Per i campioni con ruggine visibile è necessario osservare un tempo di reazione minimo di 5 minuti.
4. Le acque che sono state trattate con composti organici che proteggono dalla corrosione devono essere eventualmente ossidate per disgregare i complessi di ferro. A tale scopo si aggiunge un campione da 100 ml con 1 ml di acido solforico concentrato e 1 ml di acido nitrico concentrato e lo si fa evaporare fino alla metà. Dopo il raffreddamento viene eseguita la digestione.

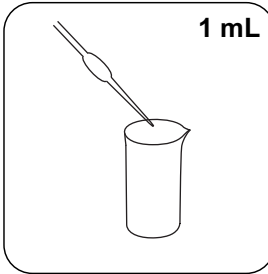
Note

1. Con questo metodo vengono misurate tutte le forme di ferro disciolto e la maggior parte delle forme di ferro non disciolto.
2. L'accuratezza non viene ridotta da eventuale polvere non disciolta.

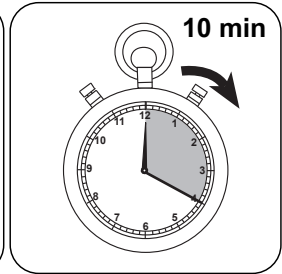
Digestione



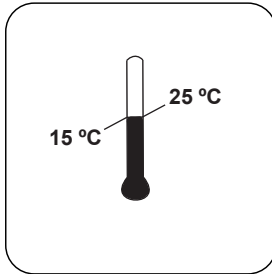
Riempire un recipiente per campioni adeguato con **100 mL di campione**.



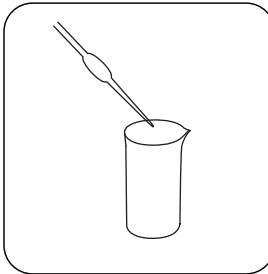
Aggiungere **1 mL di acido solforico concentrato**.



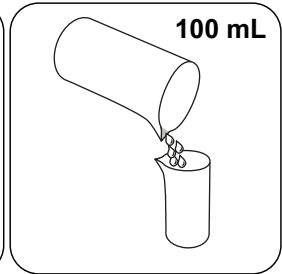
Riscaldare il campione per 10 minuti o finché non si sarà sciolto completamente.



Lasciar raffreddare il campione a **temperatura ambiente**.



Regolare il **valore di pH** del campione con **soluzione di ammoniaca** su 3-5.



Aggiungere al campione **acqua demineralizzata fino a raggiungere i 100 mL**.

Utilizzare questo campione per l'analisi di Ferro soluto e disciolto totale.

Esecuzione della rilevazione Ferro(II,III), disciolto con polvere in bustine Vario

Selezionare il metodo nel dispositivo.

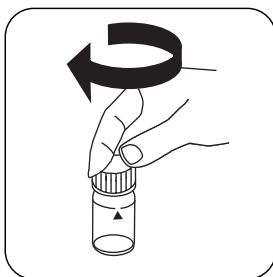
Per la determinazione di **Ferro con pastiglia** eseguire la **digestione** descritta.



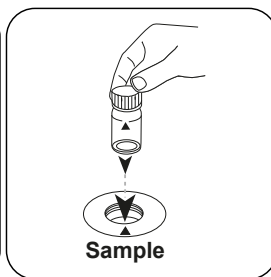
IT



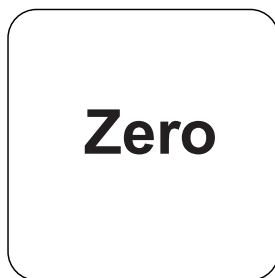
Riempire una cuvetta da 24 mm con **10 mL di campione**.



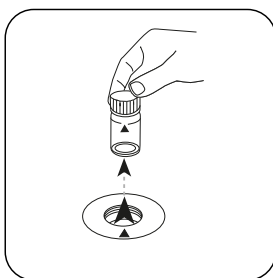
Chiudere la/e cuvetta/e.



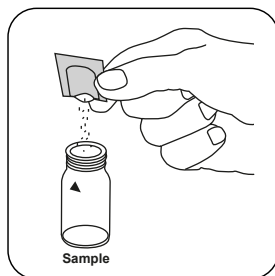
Posizionare la **cuvetta del campione** nel vano di misurazione. Fare attenzione al posizionamento.



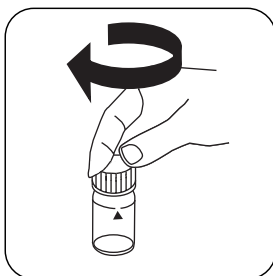
Premere il tasto **ZERO**.



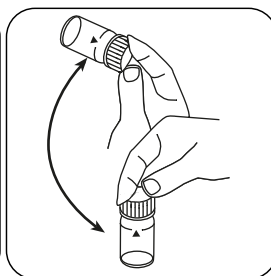
Prelevare la cuvetta dal vano di misurazione.



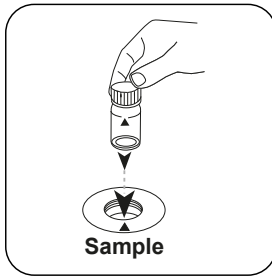
Aggiungere **una bustina di polvere Vario FERRO F10**.



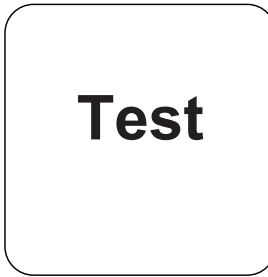
Chiudere la/e cuvetta/e.



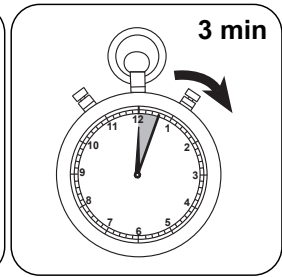
Miscelare il contenuto capovolgendo.



Posizionare la **cuvetta del campione** nel vano di misurazione. Fare attenzione al posizionamento.

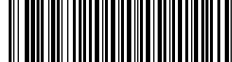


Premere il tasto **TEST** (XD: **START**).



Attendere un **tempo di reazione di 3 minuto/i**.

Allo scadere del tempo di reazione viene effettuata automaticamente la misurazione. Sul display compare il risultato in mg/L di Ferro.



Metodo chimico

1,10-fenantrolina

Appendice

IT

Interferenze

Interferenze permanenti

1. L'iridio interferisce con la rilevazione.

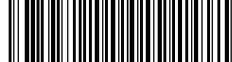
Secondo

DIN 38406-E1

Standard Method 3500-Fe-1997

US EPA 40 CFR 136

⁹⁾ Il reagente cattura la maggior parte degli ossidi di ferro



Ferro (TPTZ) PP

M223

0.02 - 1.8 mg/L Fe

FE2

TPTZ

IT

Materiale

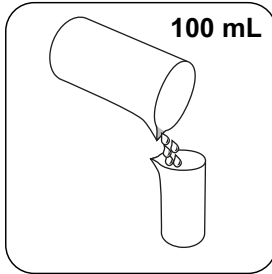
Materiale richiesto (in parte facoltativo):

Reagenti	Unità di imballaggio	N. ordine
VARIO Ferro TPTZ F10	Polvere / 100 pz.	530550

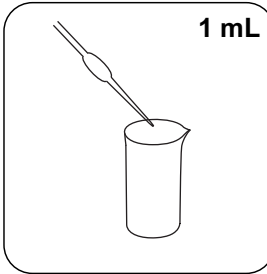
Preparazione

1. Per la rilevazione del ferro totale è necessaria una digestione. Il reagente TPTZ rileva la maggior parte degli ossidi di ferro senza digestione.
2. Prima dell'analisi sciacquare tutti i vetri di laboratorio con una soluzione di acido cloridrico diluita (1:1) e successivamente con acqua demineralizzata per eliminare i depositi di ferro, che potrebbero portare a risultati leggermente maggiorati.
3. Le acque fortemente alcaline o acide dovrebbero essere portate prima dell'analisi entro un range di pH compreso tra 3 e 8 (con 0,5 mol/l di acido solforico o 1 mol/l di liscivia).
4. Le acque che sono state trattate con composti organici che proteggono dalla corrosione devono essere eventualmente ossidate per disgregare i complessi di ferro. A tale scopo si aggiunge un campione da 100 ml con 1 ml di acido solforico concentrato e 1 ml di acido nitrico concentrato e lo si fa evaporare fino alla metà. Dopo il raffreddamento viene eseguita la digestione.

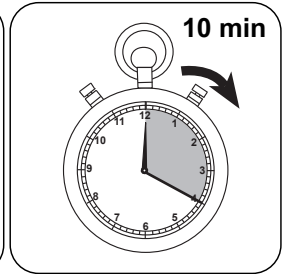
Digestione



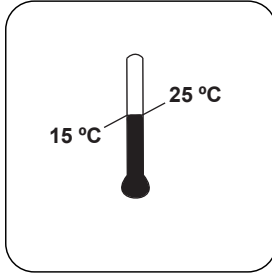
Riempire un recipiente per campioni adeguato con **100 mL di campione**.



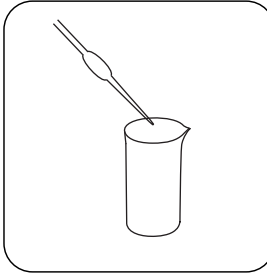
Aggiungere **1 mL di acido solforico concentrato**.



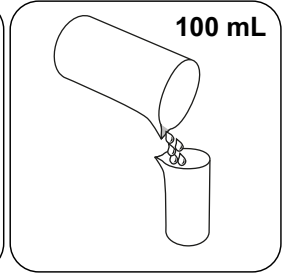
Riscaldare il campione per 10 minuti o finché non si sarà sciolto completamente.



Lasciar raffreddare il campione a **temperatura ambiente**.



Regolare il **valore di pH** del campione con **soluzione di ammoniaca** su 3-5.



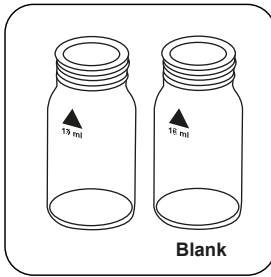
Aggiungere al campione **acqua demineralizzata fino a raggiungere i 100 mL**.

Utilizzare questo campione per l'analisi di Ferro soluto e disciolto totale.

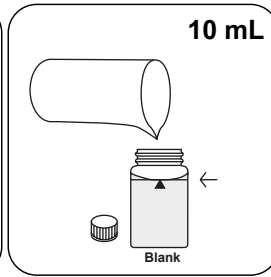
Esecuzione della rilevazione Ferro totale con polvere in bustine Vario

Selezionare il metodo nel dispositivo.

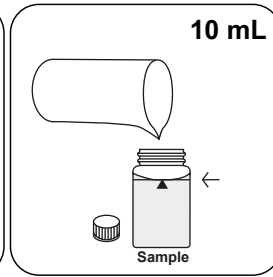
Per la determinazione di **Ferro totale** eseguire la **digestione** descritta.



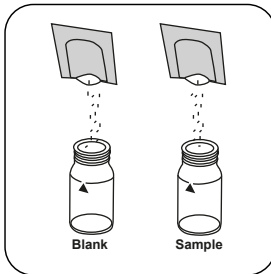
Preparare due cuvette pulite da 24 mm. Contrassegnare una cuvetta come cuvetta zero.



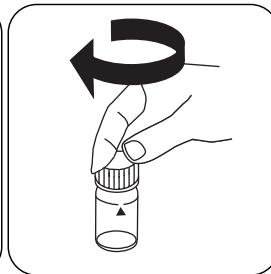
Immettere **10 mL di acqua demineralizzata** nella cuvetta zero.



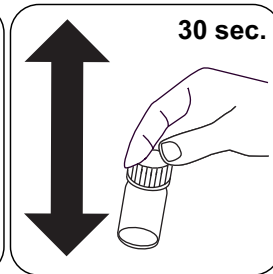
Immettere **10 mL di campione** nella cuvetta del campione.



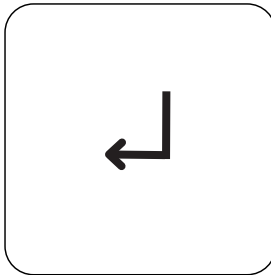
Immettere **una bustina di polvere Vario IRON TPTZ F10** in ogni cuvetta.



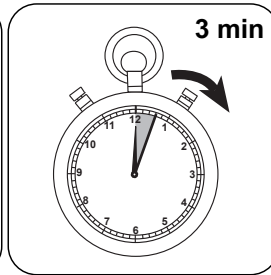
Chiudere la/e cuvetta/e.



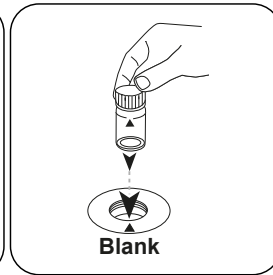
Miscelare il contenuto agitando (30 sec.).



Premere il tasto **ENTER**.



Attendere un **tempo di reazione di 3 minuti/i**.

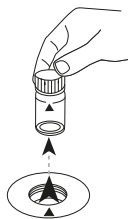


Posizionare la **cuvetta zero** nel vano di misurazione. Fare attenzione al posizionamento.

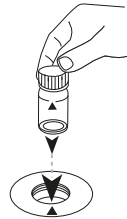


Zero

Premere il tasto **ZERO**.



Prelevare la cuvetta dal vano di misurazione.



Sample

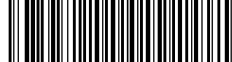
Posizionare la **cuvetta del campione** nel vano di misurazione. Fare attenzione al posizionamento.

IT

Test

Premere il tasto **TEST** (XD: **START**).

Sul display compare il risultato in mg/L di Ferro.



Metodo chimico

TPTZ

Appendice

IT

Interferenze

Interferenze permanenti

In caso di interferenze la formazione del colore viene ostacolata oppure si forma un precipitato. Le indicazioni si riferiscono a una soluzione standard con una concentrazione di ferro di 0,5 mg/L.

Interferenze	da / [mg/L]
Cd	4
Cr ³⁺	0.25
Cr ⁶⁺	1.2
Co	0.05
Cu	0.6
CN ⁻	2.8
Mn	50
Hg	0.4
Mo	4
Ni	1
NO ₂ ⁻	0.8

Riferimenti bibliografici

G. Frederic Smith Chemical Co., The Iron Reagents, 3rd ed. (1980)

Nome do método

Número do método

Código de barras para a deteção dos métodos

Área de medição

Indicado no display: MD 100 / MD 110 / MD 200

KS4.3 T / 20

Nome do método

Número do método

Código de barras para a deteção dos métodos

Área de medição

Método Químico

Indicado no display: MD 100 / MD 110 / MD 200

Informação específica do instrumento

O teste pode ser realizado nos seguintes dispositivos. Além disso, a cubeta necessária e a faixa de absorção do fotómetro são indicadas.

Dispositivos	Cubeta	λ	Faixa de Medição
MD 200, MD 600, MD 610, MD 640, MultiDirect, PM 620, PM 630	ø 24 mm	610 nm	0.1 - 4 mmol/l $K_{S4.3}$
SpectroDirect, XD 7000, XD 7500	ø 24 mm	615 nm	0.1 - 4 mmol/l $K_{S4.3}$

Material

Material necessário (parcialmente opcional):

Título	Unidade de Embalagem	Artigo No
Alka-M-Photometer	Pastilhas / 100	513210BT
Alka-M-Photometer	Pastilhas / 250	513211BT

Lista de Aplicações

- Tratamento de Esgotos
- Tratamento de Água Potável
- Tratamento de Água Bruta

Notas

1. Os termos alcalinidade-m, m-valor, alcalinidade total e capacidade de acidez $K_{S4.3}$ são idênticos.
2. O cumprimento exato do volume da amostra de 10 ml é decisivo para a precisão do resultado de análise.

Códigos de idioma ISO 639-1

Nível de revisão

PT Métodos Manual 01/20

Efetuar a medição

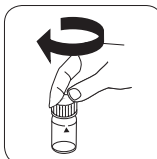
Realização da determinação Capacidade de acidez $K_{s4.3}$ com pastilha

Escolher o método no equipamento.

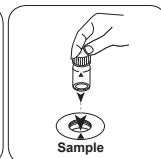
Para este método não tem de ser efetuada uma medição ZERO nos seguintes equipamentos: XD 7000, XD 7500



Encher a célula de 24 mm com 10 ml de amostra .

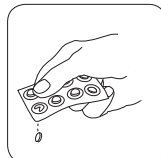


Fechar a(s) célula(s).

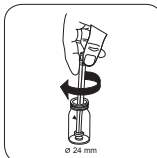


Colocar a **célula de amostra** no compartimento de medição. Observar o posicionamento.

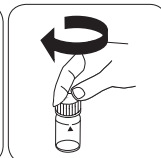
• • •



Pastilha ALKA-M-PHOTO-METER.



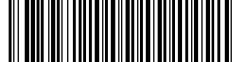
Esmagar a(s) pastilha(s) rodando ligeiramente.



Fechar a(s) célula(s).

PT Métodos Manual 01/20

PT



Ferro T

M220

0.02 - 1 mg/L Fe

FE

Ferrozine / Thioglycolate

PT

Material

Material necessário (parcialmente opcional):

Reagentes	Unidade de Embalagem	Código do Produto
Iron II LR (Fe ²⁺)	Pastilhas / 100	515420BT
Iron II LR (Fe ²⁺)	Pastilhas / 250	515421BT
Iron LR (Fe ²⁺ und Fe ³⁺)	Pastilhas / 100	515370BT
Iron LR (Fe ²⁺ und Fe ³⁺)	Pastilhas / 250	515371BT

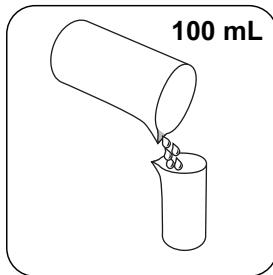
Preparação

1. As águas que foram tratadas com compostos orgânicos como proteção anticorrosiva, etc. têm de ser eventualmente oxidadas para destruir os complexos de ferro. Para isso, transfere-se uma amostra de 100 ml com 1 ml de ácido sulfúrico concentrado e 1 ml de ácido nítrico concentrado e evaporada para metade. Depois de arrefecer, passa-se à digestão.

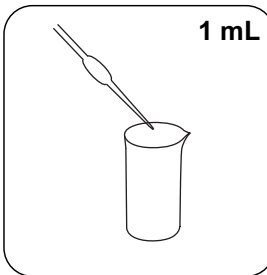
Notas

1. Neste método ocorre a determinação de Fe²⁺ e Fe³⁺ totalmente dissolvido.
2. Para determinar Fe²⁺ usa-se a pastilha IRON (II) LR, em vez da pastilha IRON LR.

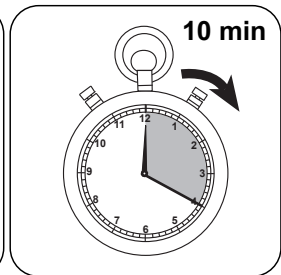
Digestão



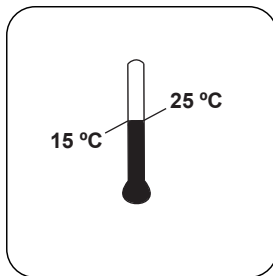
Encher um recipiente de amostra adequado com **100 mL de amostra** .



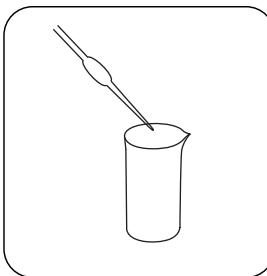
Adicionar **1 mL ácido sulfúrico concentrado** .



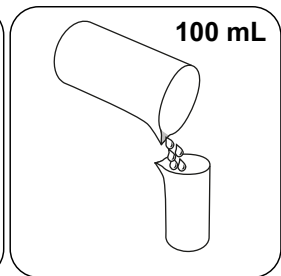
A amostra deve **aquecer durante 10 minutos**, ou até tudo se ter totalmente dissolvido.



Deixar a amostra arrefecer até à **temperatura ambiente** .



Ajustar o **valor pH** da amostra com **solução amoniacal para 3-5**.



Encher a amostra com **água desmineralizada até 100 mL** .

Usar esta amostra para a análise de total de ferro solvido e dissolvido.

Realização da determinação Ferro(II,III), dissolvido com pastilha

Escolher o método no equipamento.

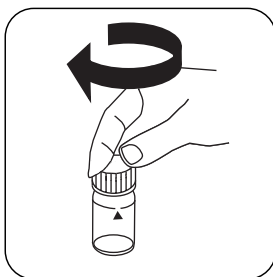
Para a determinação de **Ferro dissolvido e não dissolvido** deve realizar a **digestão** descrita.



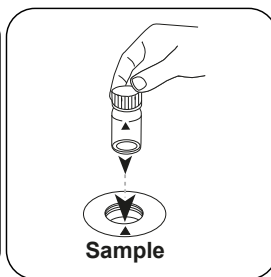
PT



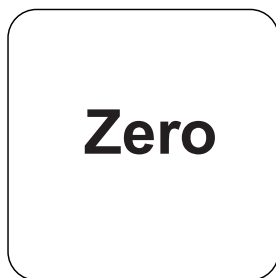
Encher a célula de 24 mm com **10 mL de amostra**.



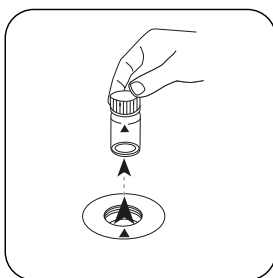
Fechar a(s) célula(s).



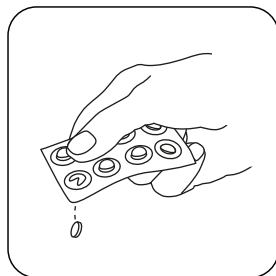
Colocar a **célula de amostra** no compartimento de medição. Observar o posicionamento.



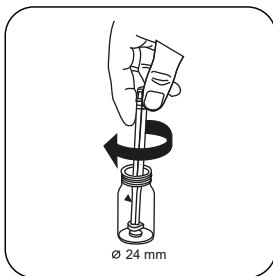
Premir a tecla **ZERO**.



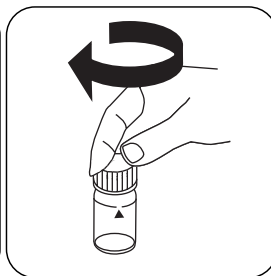
Retirar a célula do compartimento de medição.



Pastilha IRON LR.



Esmagar a(s) pastilha(s) rodando ligeiramente.



Fechar a(s) célula(s).



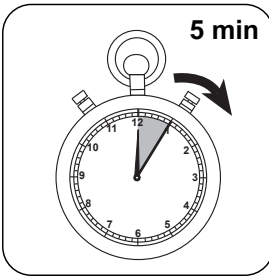
Dissolver a(s) pastilha(s) girando.



Colocar a **célula de amostra** no compartimento de medição. Observar o posicionamento.



Premir a tecla **TEST** (XD: **START**).



Aguardar **5 minuto(s) de tempo de reação**.

Decorrido o tempo de reação, a medição é efetuada automaticamente.

No visor aparece o resultado em mg/L Ferro.



Método Químico

Ferrozine / Thioglycolate

Apêndice

PT

Texto de Interferências

Interferências Removíveis

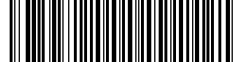
1. A presença de cobre aumenta o resultado de medição em 10 %. Numa concentração de 10 mg/L de cobre na amostra, o resultado de medição aumenta em 1 mg/L de ferro.
A interferência pode ser eliminada com a adição de tiourea

Validação de método

Limite de Detecção	0.01 mg/L
Limite de Determinação	0.016 mg/L
Fim da Faixa de Medição	1 mg/L
Sensibilidade	0.92 mg/L / Abs
Faixa de Confiança	0.013 mg/L
Desvio Padrão	0.005 mg/L
Coefficiente de Variação	1.23 %

Bibliografia

Análise fotométrica, Lange/ Vjedelek, Verlag Chemie 1980, S. 102



Ferro PP

M222

0.02 - 3 mg/L Fe⁹⁾

FE1

1,10-Phenanthroline

PT

Material

Material necessário (parcialmente opcional):

Reagentes	Unidade de Embalagem	Código do Produto
VARIO Iron F10	Pó / 100 pc.	530560
VARIO Iron F10	Pó / 1000 pc.	530563

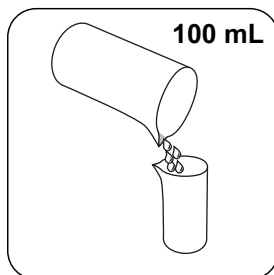
Preparação

1. Óxido de ferro requer, antes da análise, uma digestão fraca, forte ou Digesdahl (processo ácido de digestão).
2. As águas fortemente alcalinas ou ácidas deviam, antes da análise, ser ajustadas para um valor pH entre 3 e 5.
3. No caso de amostras que incluem ferrugem visível, devia manter um tempo de reação mínimo de 5 minutos.
4. As águas que foram tratadas com compostos orgânicos como proteção anticorrosiva, etc. têm de ser eventualmente oxidadas para destruir os complexos de ferro. Para isso, transfere-se uma amostra de 100 ml com 1 ml de ácido sulfúrico concentrado e 1 ml de ácido nítrico concentrado e evaporada para metade. Depois de arrefecer, passa-se à digestão.

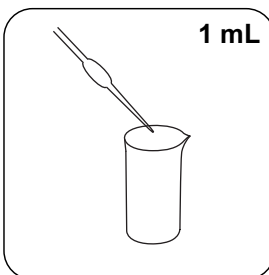
Notas

1. Neste método ocorre a determinação de todas as formas de ferro dissolvido e da maioria das formas de ferro não dissolvido.
2. A precisão não é reduzida pelo pó não dissolvido.

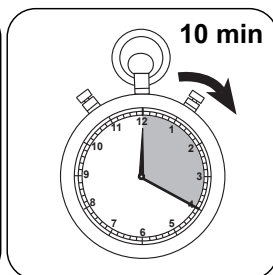
Digestão



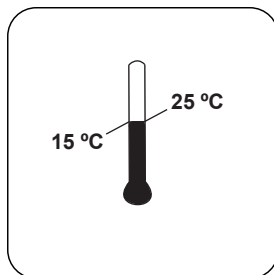
Encher um recipiente de amostra adequado com **100 mL de amostra** .



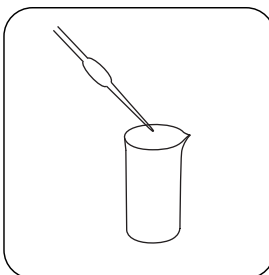
Adicionar **1 mL ácido sulfúrico concentrado** .



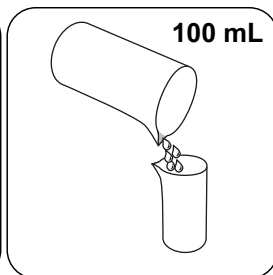
A amostra deve **aquecer durante 10 minutos**, ou até tudo se ter totalmente dissolvido.



Deixar a amostra arrefecer até à **temperatura ambiente** .



Ajustar o **valor pH** da amostra com **solução amoniacal para 3-5**.



Encher a amostra com **água desmineralizada até 100 mL** .

Usar esta amostra para a análise de total de ferro solvido e dissolvido.

Realização da determinação Ferro(II,III), dissolvido com pacote de pó Vario

Escolher o método no equipamento.

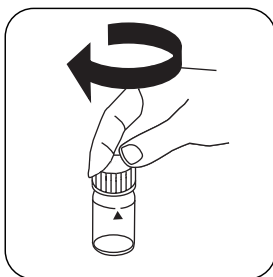
Para a determinação de **Ferro com pastilha** deve realizar a **digestão** descrita.



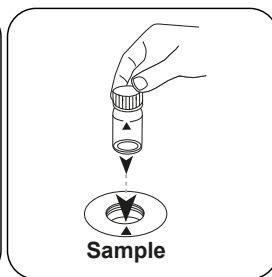
PT



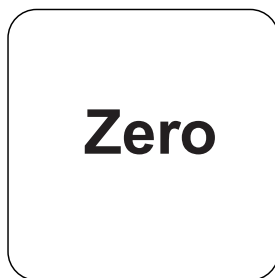
Encher a célula de 24 mm com **10 mL de amostra**.



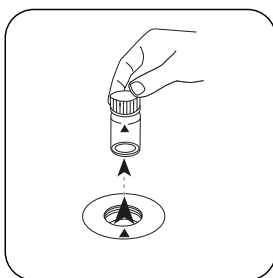
Fechar a(s) célula(s).



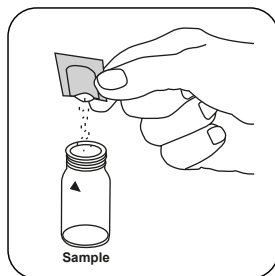
Colocar a **célula de amostra** no compartimento de medição. Observar o posicionamento.



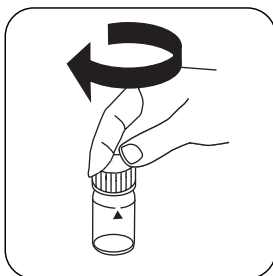
Premir a tecla **ZERO**.



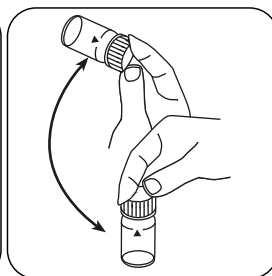
Retirar a célula do compartimento de medição.



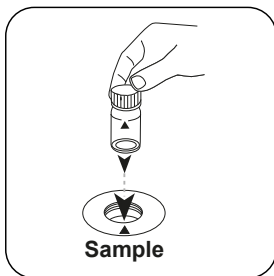
Adicionar um **pacote de pó Vario FERRO F10**.



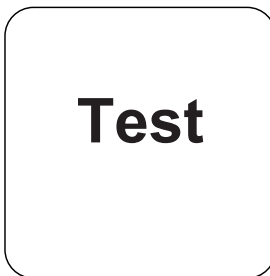
Fechar a(s) célula(s).



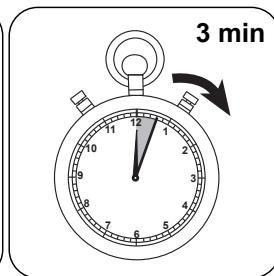
Misturar o conteúdo girando.



Colocar a **célula de amostra** no compartimento de medição. Observar o posicionamento.



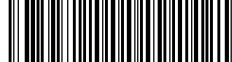
Premir a tecla **TEST** (XD: **START**).



Aguardar **3 minuto(s) de tempo de reação**.

Decorrido o tempo de reação, a medição é efetuada automaticamente.

No visor aparece o resultado em mg/L Ferro.



Método Químico

1,10-Phenanthroline

Apêndice

PT

Texto de Interferências

Interferências Persistentes

1. Iridio interfere na determinação.

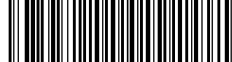
De acordo com

DIN 38406-E1

Standard Method 3500-Fe-1997

US EPA 40 CFR 136

⁹⁾Reagente captura a maioria dos óxidos de ferro



Ferro (TPTZ) PP

M223

0.02 - 1.8 mg/L Fe

FE2

TPTZ

PT

Material

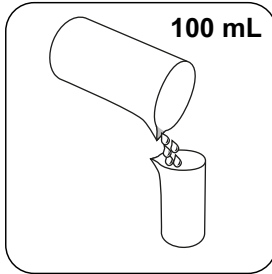
Material necessário (parcialmente opcional):

Reagentes	Unidade de Embalagem	Código do Produto
Iron VARIO TPTZ F10	Pó / 100 pc.	530550

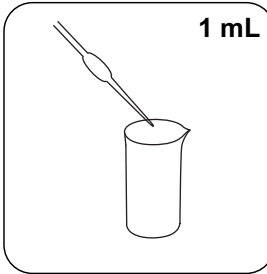
Preparação

1. A determinação de ferro total requer uma digestão. O reagente TPTZ capta a maioria de óxidos de ferro sem digestão.
2. Enxaguar todos os vidros para laboratório, antes da análise, com solução de ácido clorídrico diluído (1:1) e depois com água desmineralizada, para eliminar os depósitos de ferro, que podem causar resultados ligeiramente aumentados.
3. As águas fortemente alcalinas ou ácidas deviam, antes da análise, ser ajustadas para um valor pH entre 3 e 8 (com 0,5 mol/l de ácido sulfúrico ou 1 mol/l soda cáustica).
4. As águas que foram tratadas com compostos orgânicos como proteção anticorrosiva, etc. têm de ser eventualmente oxidadas para destruir os complexos de ferro. Para isso, transfere-se uma amostra de 100 ml com 1 ml de ácido sulfúrico concentrado e 1 ml de ácido nítrico concentrado e evaporada para metade. Depois de arrefecer, passa-se à digestão.

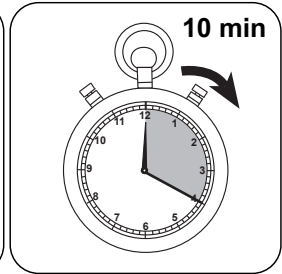
Digestão



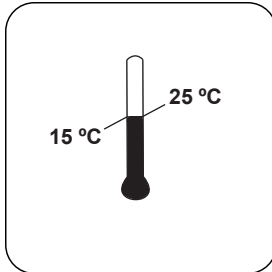
Encher um recipiente de amostra adequado com **100 mL de amostra** .



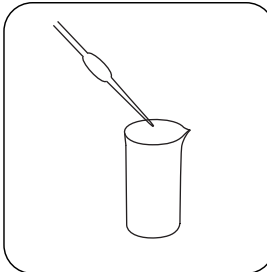
Adicionar **1 mL ácido sulfúrico concentrado** .



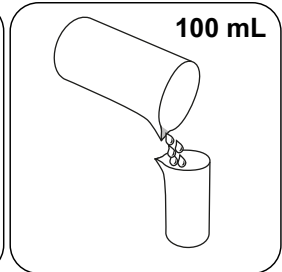
A amostra deve **aquecer durante 10 minutos**, ou até tudo se ter totalmente dissolvido.



Deixar a amostra arrefecer até à **temperatura ambiente** .



Ajustar o **valor pH** da amostra com **solução amoniacal para 3-5**.



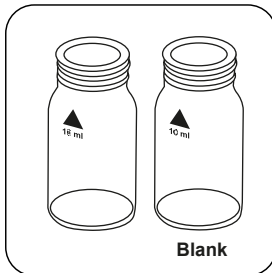
Encher a amostra com **água desmineralizada até 100 mL** .

Usar esta amostra para a análise de total de ferro solvido e dissolvido.

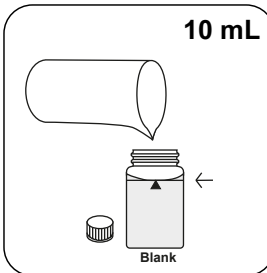
Realização da determinação Ferro, total com pacote de pó Vario

Escolher o método no equipamento.

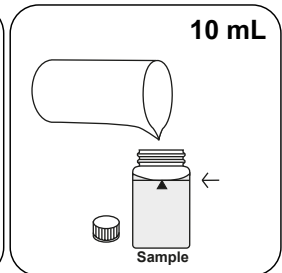
Para a determinação de **Ferro total** deve realizar a **digestão** descrita.



Preparar duas células de 24 mm limpas. Identificar uma célula como célula zero.



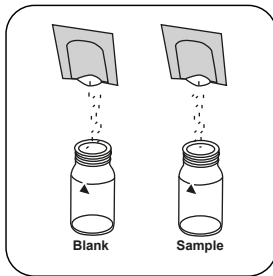
Adicionar **10 mL de água desmineralizada** à célula zero.



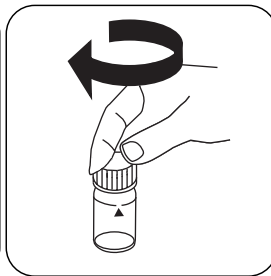
Adicionar **10 mL de amostra** à célula de amostra.



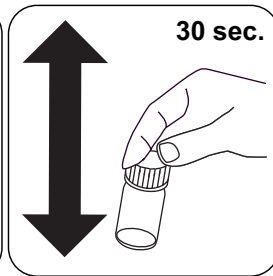
PT



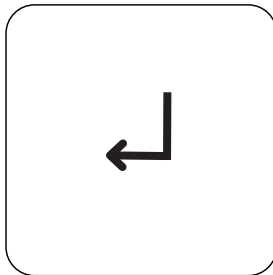
Introduzir em cada célula um pacote de pó Vario IRON TPTZ F10.



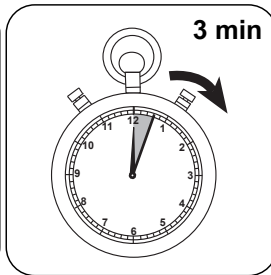
Fechar a(s) célula(s).



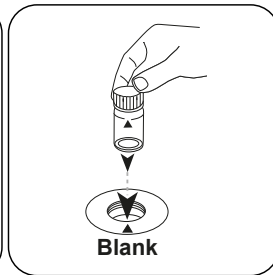
Misturar o conteúdo girando (30 sec.).



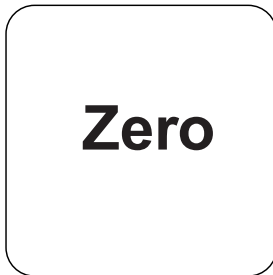
Premir a tecla **ENTER**.



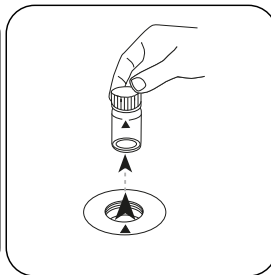
Aguardar **3 minuto(s)** de tempo de reação.



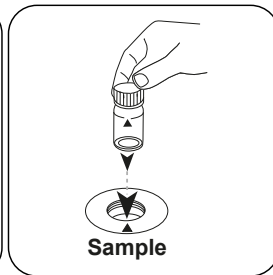
Colocar a **célula zero** no compartimento de medição. Observar o posicionamento.



Premir a tecla **ZERO**.



Retirar a célula do compartimento de medição.



Colocar a **célula de amostra** no compartimento de medição. Observar o posicionamento.

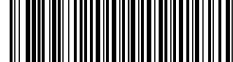


Test

Premir a tecla **TEST** (XD:
START).

No visor aparece o resultado em mg/L Ferro.

PT



Método Químico

TPTZ

Apêndice

PT

Texto de Interferências

Interferências Persistentes


Em caso de interferência, a formação de cor é inibida ou pode formar-se um sedimento. As indicações referem-se a um padrão com uma concentração de ferro de 0,5 mg/L.

Interferências	a partir de / [mg/L]
Cd	4
Cr ³⁺	0.25
Cr ⁶⁺	1.2
Co	0.05
Cu	0.6
CN ⁻	2.8
Mn	50
Hg	0.4
Mo	4
Ni	1
NO ₂ ⁻	0.8

Bibliografia

G. Frederic Smith Chemical Co., The Iron Reagents, 3rd ed. (1980)

KS4.3 T / 20



Naam van de methode

Nummer methode

Streepjescode ter identificatie van de methode

Meetbereik

$K_{S_{4.3}} T$ M20
0.1 - 4 mmol/l $K_{S_{4.3}}$ S:4.3
Zuur / Indicator

Chemische methode

Uitlezing in MD
100 MD 110 / MD 200

Instrument specifieke informatie

De test kan op de volgende apparaten worden uitgevoerd. Bovendien worden de vereiste cuvette en het absorptiebereik van de fotometer aangegeven.

Toestellen	Cuvet	λ	Meetbereik
MD 200, MD 600, MD 610, MD 640, MultiDirect, PM 620, PM 630	\varnothing 24 mm	610 nm	0.1 - 4 mmol/l $K_{S_{4.3}}$
SpectroDirect, XD 7000, XD 7500	\varnothing 24 mm	615 nm	0.1 - 4 mmol/l $K_{S_{4.3}}$

Reagentia

Benodigd materiaal (deels optioneel):

Titel	Verpakkingseenheid	Bestelnr.
Alka-M-Photometer	Tablet / 100	513210BT
Alka-M-Photometer	Tablet / 250	513211BT

Toepassingsbereik

- Afvalwaterzuivering
- Behandeling drinkwater
- Zuivering vervuild water

Aantekeningen

1. De termen alkaliteit-m, m-waarde, totale alkaliteit en zuurcapaciteit_{S_{4.3}} zijn identiek.
2. De exacte naleving van het monstervolume van 10 ml is bepalend voor de nauwkeurigheid van het analysesresultaat.

Beknopte naam conform de norm ISO 639-1

Herziene versie

NL Handboek van Methoden 01/20

Uitvoering van de meting

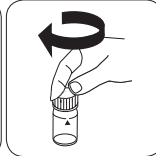
Uitvoering van de bepaling Zuurcapaciteit $K_{s4,3}$ met tablet

De methode in het apparaat selecteren.

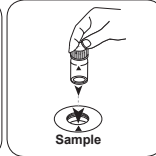
Voor deze methode moet bij de volgende apparaten geen nulmeting worden uitgevoerd:
XD 7000, XD 7500



Spoelbakje van 24 mm met **10 ml** staal vullen.

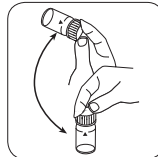


De spoelbakjes afsluiten.

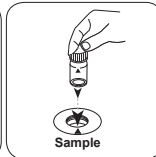


Het **staalspoelbakje** in de meetschacht plaatsen. Op de positionering letten.

• • •



Tabletten oplossen door om te draaien



Het **staalspoelbakje** in de meetschacht plaatsen. Op de positionering letten.



De toets **TEST** (XD: **START**) indrukken.

De display toont het resultaat als Zuurcapaciteit $K_{s4,3}$.



IJzer T

M220

0.02 - 1 mg/L Fe

FE

Ferrozine / Thioglycolaat

NL

Reagentia

Benodigd materiaal (deels optioneel):

Reagentia	Verpakkingseenheid	Bestelnr.
IJzer II LR (Fe ²⁺)	Tablet / 100	515420BT
IJzer II LR (Fe ²⁺)	Tablet / 250	515421BT
IJzer LR (Fe ²⁺ und Fe ³⁺)	Tablet / 100	515370BT
IJzer LR (Fe ²⁺ und Fe ³⁺)	Tablet / 250	515371BT

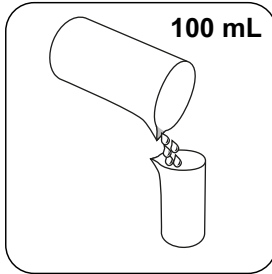
Voorbereiding

1. Water dat is behandeld met organische verbindingen als corrosiebescherming e.d. moet mogelijk worden geoxideerd worden om de ijzercomplexen te vernietigen. Hiertoe wordt een monster van 100 ml gemengd met 1 ml geconcentreerd zwavelzuur ($\geq 95\%$) en 1 ml geconcentreerd salpeterzuur ($\geq 65\%$) en op de helft ingedampd. Na afkoeling wordt de vertering uitgevoerd.

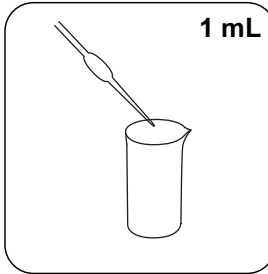
Aantekeningen

1. Deze methode bepaalt het totaal opgeloste Fe²⁺ en Fe³⁺.
2. Voor de bepaling van Fe²⁺ wordt het IRON (II) LR-tablet gebruikt in plaats van het IRON LR-tablet.

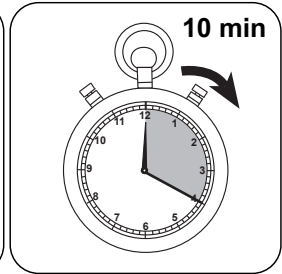
Ontsluiting



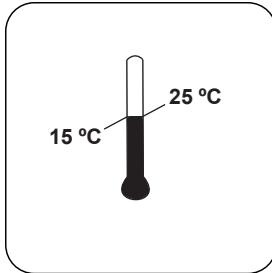
Een geschikte staalbeker met **100 mL staal** vullen.



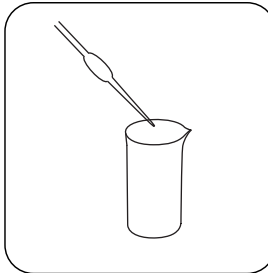
1 mL geconcentreerd zwavelzuur ($\geq 95\%$) toevoegen.



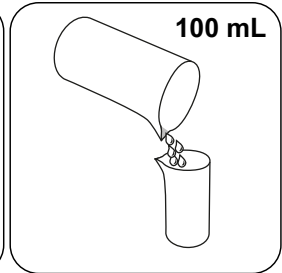
Het staal gedurende **10 minuten verwarmen**, of zolang tot alles volledig is opgelost.



Het staal laten afkoelen tot **kamertemperatuur**.



De **pH-waarde** van het staal met **Ammoniakoplossing (10-25 %)** afstellen op 3-5.



Het staal met **gedeïoniseerd water tot 100 mL** vullen.

Dit staal gebruiken voor de analyse van totaal opgelost en niet-opgelost ijzer.

Uitvoering van de bepaling IJzer (II,III), opgelost met tablet

De methode in het apparaat selecteren.

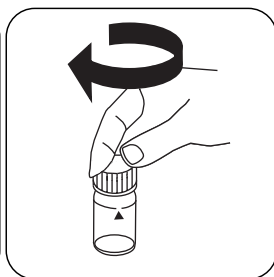
Voor de bepaling van **opgelost en niet-opgelost ijzer** de beschreven **ontsluiting** uitvoeren.



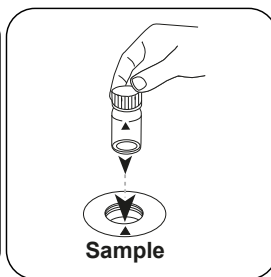
NL



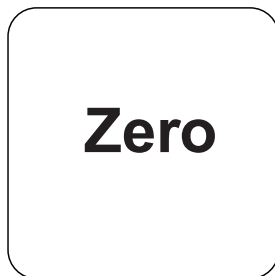
Spoelbakje van 24 mm met **10 mL staal** vullen.



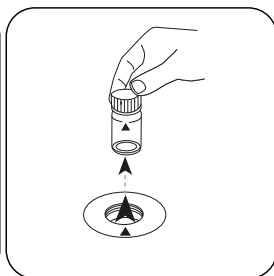
De spoelbakjes afsluiten.



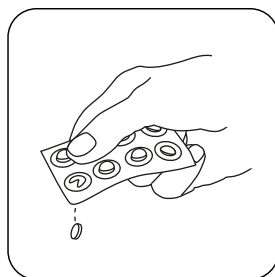
Het **staalspoelbakje** in de meetschacht plaatsen. Op de positionering letten.



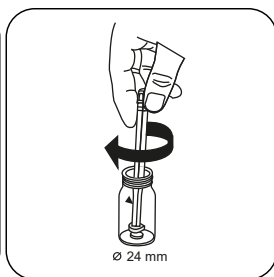
De toets **NUL** indrukken.



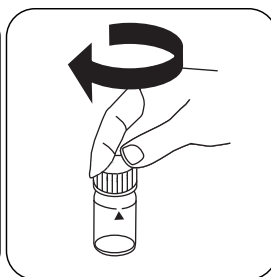
Het spoelbakje uit de meetschacht nemen.



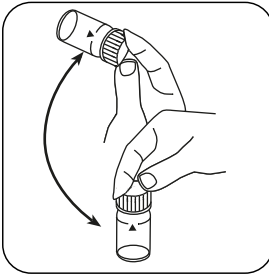
Een **IRON LR tablet** toevoegen.



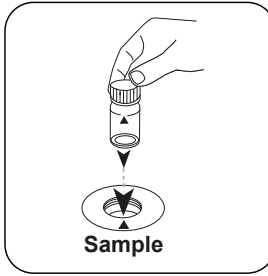
De tabletten onder lichte rotatie verpletteren.



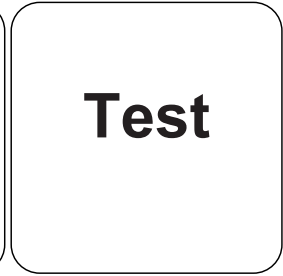
De spoelbakjes afsluiten.



Tabletten oplossen door om te draaien

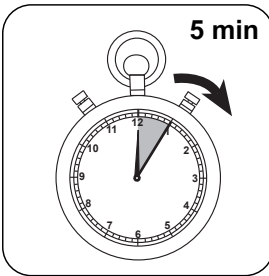


Het **staalspoelbakje** in de meetschacht plaatsen. Op de positionering letten.



De toets **TEST** (XD: **START**) indrukken.

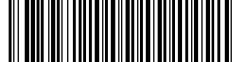
NL



De reactietijd van 5 minuten afwachten.

Na afloop van de reactietijd wordt de meting automatisch uitgevoerd.

De display toont het resultaat in mg/L IJzer.



Chemische methode

Ferrozine / Thioglycolaat

Aanhangsel

NL

Verstoringsen

Uit te sluiten verstoringen

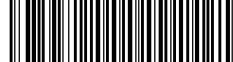
1. De aanwezigheid van koper verhoogt het meetresultaat met 10 %. Bij een concentratie van 10 mg/L koper in het monster wordt het resultaat verhoogd met 1 mg/L ijzer.
De verstoring kan worden geëlimineerd door toevoeging van thiourem

Validatie van de methodes

Aantoonbaarheidsgrens	0.01 mg/L
Bepaalbaarheidsgrens	0.016 mg/L
Einde meetbereik	1 mg/L
Gevoeligheid	0.92 mg/L / Abs
Betrouwbaarheidsgrenzen	0.013 mg/L
Standaardafwijking procedure	0.005 mg/L
Variatiecoëfficiënt procedure	1.23 %

Literatuurverwijzing

Photometrische Analyse, Lange/ Vjedelek, Verlag Chemie 1980, S. 102



IJzer PP

M222

0.02 - 3 mg/L Fe⁹⁾

FE1

1,10-fenantroline

NL

Reagentia

Benodigd materiaal (deels optioneel):

Reagentia	Verpakkingseenheid	Bestelnr.
VARIO Ferro F10	Poeder / 100 St.	530560
VARIO Ferro F10	Poeder / 1000 St.	530563

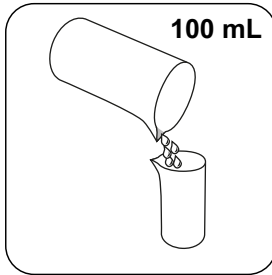
Vorbereiding

1. IJzeroxide vereist een zwakke, sterke of spijsverteringsvertering voor de analyse (zure vertering).
2. Zeer sterk alkalische of zure wateren moeten vóór de analyse op een pH-waarde tussen 3 en 5 worden ingesteld.
3. Voor monsters die zichtbare roest bevatten, moet een reactietijd van ten minste 5 minuten in acht worden genomen.
4. Water dat is behandeld met organische verbindingen als corrosiebescherming e.d. moet mogelijk worden geoxideerd worden om de ijzercomplexen te vernietigen. Hiertoe wordt een monster van 100 ml gemengd met 1 ml geconcentreerd zwavelzuur ($\geq 95\%$) en 1 ml geconcentreerd salpeterzuur ($\geq 65\%$) en op de helft ingedampd. Na afkoeling wordt de vertering uitgevoerd.

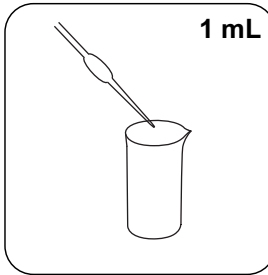
Aantekeningen

1. Deze methode bepaalt alle vormen van opgelost ijzer en de meeste vormen van onopgelost ijzer.
2. De nauwkeurigheid wordt niet verminderd door onopgelost poeder.

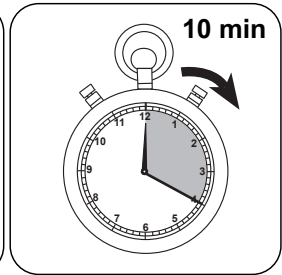
Ontsluiting



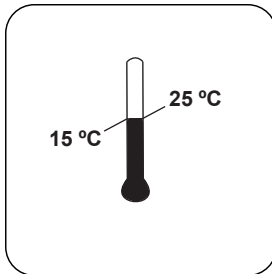
Een geschikte staalbeker met **100 mL** staal vullen.



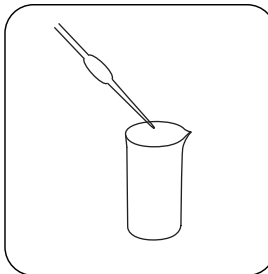
1 mL geconcentreerd zwavelzuur ($\geq 95\%$) toevoegen.



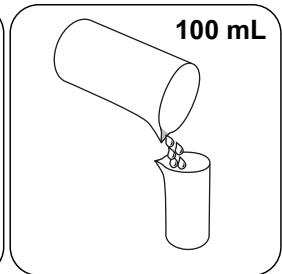
Het staal gedurende **10 minuten verwarmen**, of zolang tot alles volledig is opgelost.



Het staal laten afkoelen tot **kamertemperatuur**.



De **pH-waarde** van het staal met **Ammoniakoplossing (10-25 %)** afstellen op 3-5.



Het staal met **gedeïoniseerd water tot 100 mL** vullen.

Dit staal gebruiken voor de analyse van totaal opgelost en niet-opgelost ijzer.

Uitvoering van de bepaling IJzer(II,III), opgelost met Vario-poederpakje

De methode in het apparaat selecteren.

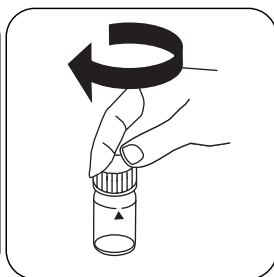
Voor de bepaling van **IJzer met tablet** de beschreven **ontsluiting** uitvoeren.



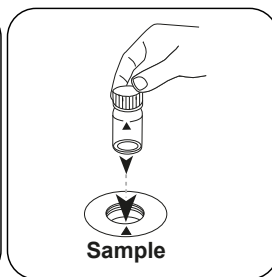
NL



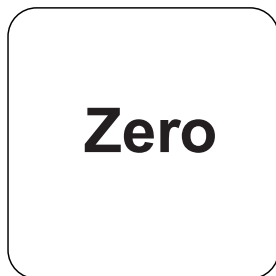
Spoelbakje van 24 mm met
10 mL staal vullen.



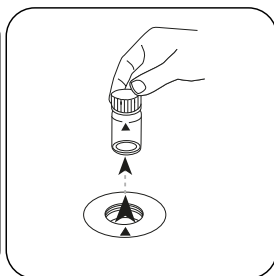
De spoelbakjes afsluiten.



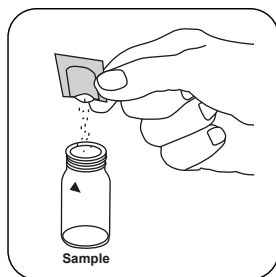
Het **staalspoelbakje** in de
meetschacht plaatsen. Op
de positionering letter.



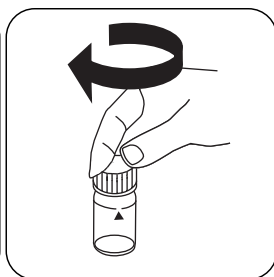
De toets **NUL** indrukken.



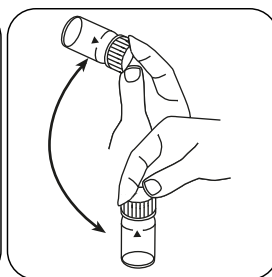
Het spoelbakje uit de
meetschacht nemen.



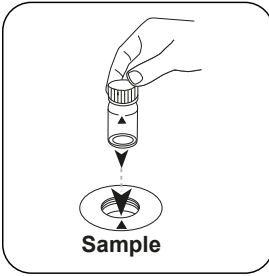
Een **Vario FERRO
F10 poederpakje**
toevoegen.



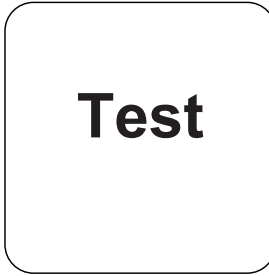
De spoelbakjes afsluiten.



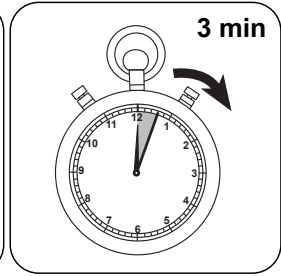
De inhoud mengen door om
te draaien.



Het **staalspoelbakje** in de meetschacht plaatsen. Op de positionering letten.



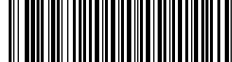
De toets **TEST** (XD: **START**) indrukken.



De reactietijd van **3 minuten** afwachten.

Na afloop van de reactietijd wordt de meting automatisch uitgevoerd.

De display toont het resultaat in mg/L IJzer.



Chemische methode

1,10-fenantroline

Aanhangsel

NL

Verstoringsen

Permanente verstoringen

1. Iridium verstoort de bestemming.

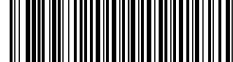
Overeenkomstig

DIN 38406-E1

Standaardmethode 3500-Fe-1997

US EPA 40 CFR 136

⁹⁾ reagens omvat de meeste ijzeroxiden



Ijzer (TPTZ) PP

M223

0.02 - 1.8 mg/L Fe

FE2

TPTZ

NL

Reagentia

Benodigd materiaal (deels optioneel):

Reagentia

VARIO Ijzer TPTZ F10

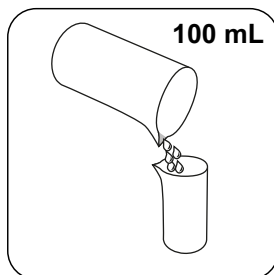
Verpakkingseenheid

Poeder / 100 St. 530550

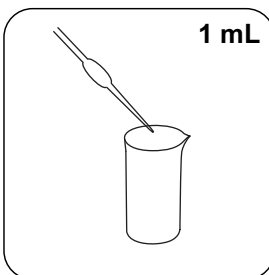
Vorbereiding

1. Voor de bepaling van het totale ijzergehalte is een spijsvertering nodig. Het TPTZ-reagens detecteert de meeste ijzeroxiden zonder spijsvertering.
2. Spoel vóór de analyse al het glaswerk in het laboratorium met verdund zoutzuur (1:1) en vervolgens met gedeïoniseerd water om ijzerdepots, die tot iets hogere resultaten kunnen leiden, te verwijderen.
3. Sterk alkalisch of zuur water moet vóór de analyse in een pH-gebied tussen 3 en 8 (met 0,5 mol/l zwavelzuur of 1 mol/l-natriumhydroxideoplossing) worden geplaatst.
4. Water dat is behandeld met organische verbindingen als corrosiebescherming e.d. moet mogelijk worden geoxideerd worden om de ijzercomplexen te vernietigen. Hiertoe wordt een monster van 100 ml gemengd met 1 ml geconcentreerd zwavelzuur ($\geq 95\%$) en 1 ml geconcentreerd salpeterzuur ($\geq 65\%$) en op de helft ingedampt. Na afkoeling wordt de vertering uitgevoerd.

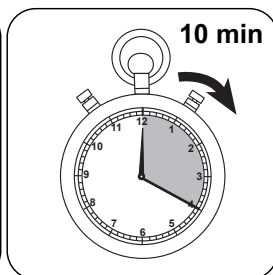
Ontsluiting



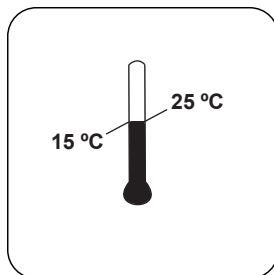
Een geschikte staalbeker met **100 mL staal** vullen.



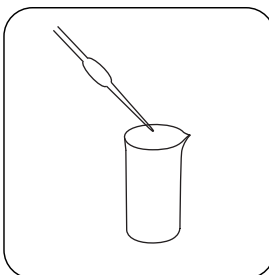
1 mL geconcentreerd zwavelzuur ($\geq 95\%$) toevoegen.



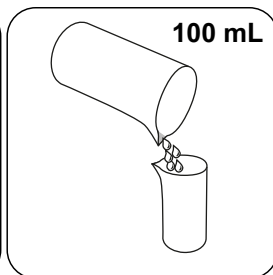
Het staal gedurende **10 minuten verwarmen**, of zolang tot alles volledig is opgelost.



Het staal laten afkoelen tot **kamertemperatuur**.



De **pH-waarde** van het staal met **Ammoniakoplossing (10-25 %)** afstellen op 3-5.



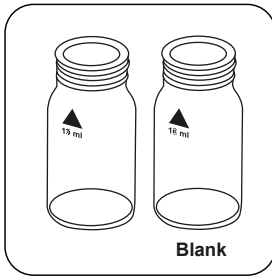
Het staal met **gedeïoniseerd water tot 100 mL** vullen.

Dit staal gebruiken voor de analyse van totaal opgelost en niet-opgelost ijzer.

Uitvoering van de bepaling IJzer, totaal met Vario-poederpakje

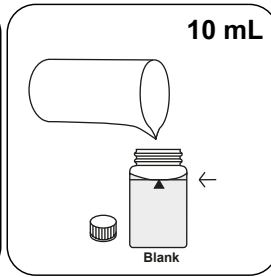
De methode in het apparaat selecteren.

Voor de bepaling van **Totaal ijzer** de beschreven **ontsluiting** uitvoeren.



Blank

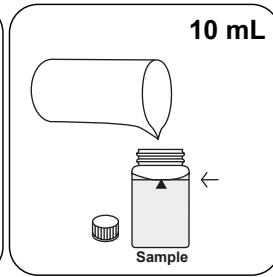
Twee propere spoelbakjes van 24 mm klaarzetten. Een als nulspoelbakje kenmerken.



10 mL

Blank

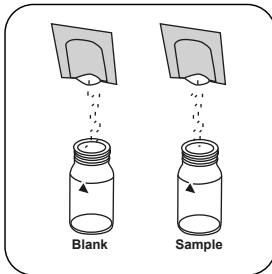
10 mL gedeïoniseerd water in het nulspoelbakje doen.



10 mL

Sample

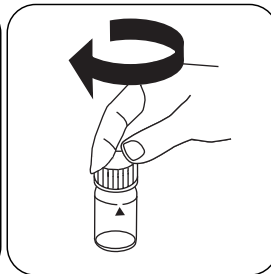
10 mL staal in het staalspoelbakje doen.



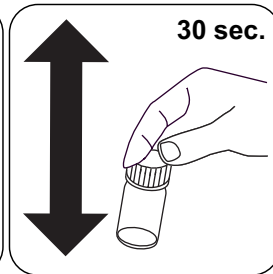
Blank

Sample

In elk spoelbakje **een Vario IRON TPTZ F10 poederpakje** doen.

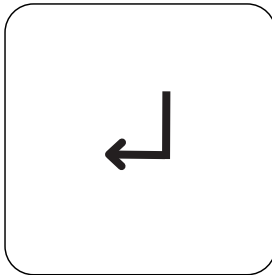


De spoelbakjes afsluiten.

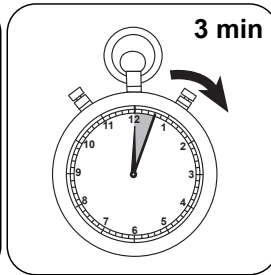


30 sec.

De inhoud mengen door te schudden (30 sec.).

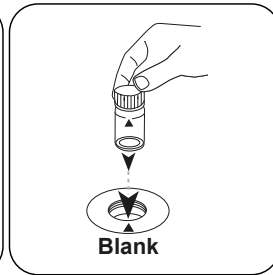


De toets **ENTER** indrukken.



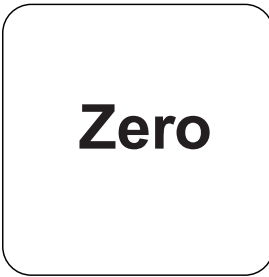
3 min

De reactietijd van **3 minuten** afwachten.

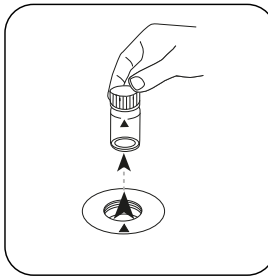


Blank

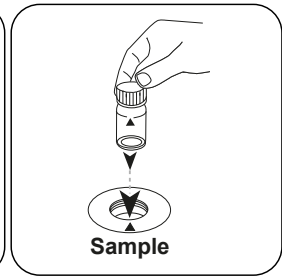
Het **nulspoelbakje** in de meetschacht plaatsen. Op de positionering letteren.



De toets **NUL** indrukken.

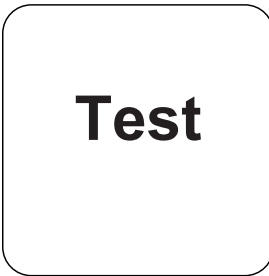


Het spoelbakje uit de meetschacht nemen.



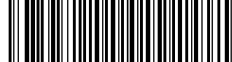
Het **staalspoelbakje** in de meetschacht plaatsen. Op de positionering letteren.

NL



De toets **TEST** (XD: **START**) indrukken.

De display toont het resultaat in mg/L IJzer.



Chemische methode

TPTZ

Aanhangsel

NL

Verstoringen

Permanente verstoringen

Als er storingen optreden, wordt de kleurvorming geremd of worden er neerslagvormen gevormd. De gegevens hebben betrekking op een norm met een ijzerconcentratie van 0,5 mg/L.

Verstoringen	verstoort vanaf
Cd	4
Cr ³⁺	0.25
Cr ⁶⁺	1.2
Co	0.05
Cu	0.6
CN ⁻	2.8
Mn	50
Hg	0.4
Mo	4
Ni	1
NO ₂ ⁻	0.8

Literatuurverwijzing

G. Frederic Smith Chemical Co., The Iron Reagents, 3rd ed. (1980)

KS4.3 T / 20

方法名称

方法号

用于方法检测的条形码

测量范围

酸性 / 指示剂

屏幕显示: MD 100 / MD 110 / MD 200

化学方法

儀器的具體信息

測試可以在以下設備上執行。此外還指出了所需的比色杯和光度計的吸收範圍。

儀器類型	比色皿	λ	測量範圍
MD 200, MD 600, MD 610, MD 640, MultiDirect, PM 620, PM 630	\varnothing 24 mm	610 nm	0.1 - 4 mmol/l $K_{S4.3}$
SpectroDirect, XD 7000, XD 7500	\varnothing 24 mm	615 nm	0.1 - 4 mmol/l $K_{S4.3}$

材料

所需材料 (部分可選) :

標題	包裝單位	貨號
Alka-M-Photometer	片劑 / 100	513210BT
Alka-M-Photometer	片劑 / 250	513211BT

應用列表

- 污水處理
- 飲用水處理
- 原水處理

備註

1. 術語總度-m、m-值、總碱度和酸容量 $K_{S4.3}$ 是相同的。
2. 準確地遵守 10 ml 的樣本體積對分析結果的準確度至關重要。

語言代碼 ISO 639-1

修訂狀態

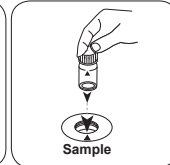
CN 方法手冊 01/20

开始测量

进行测定 $K_{s4.3}$ 片剂酸容量

选择设备中的方法。

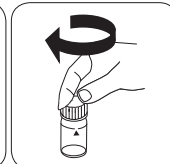
对于这种方法，在以下设备上不能进行 ZERO 测量：XD 7000, XD 7500

用 10 ml 样本填充 24 mm 比密封比色杯。
色杯。将样本比色杯放入测量轴
中。注意定位。

• • •

加入 ALKA-M-PHOTOME-
TER 片剂。

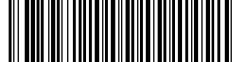
用轻微的扭转压碎片剂。



密封比色杯。

CN 方法手册 01/20

ZH



T 铁

M220

0.02 - 1 mg/L Fe

FE

Ferrozine/巯乙酸盐

材料

所需材料 (部分可选) :

ZH

试剂	包装单位	货号
铁 II LR (Fe ²⁺)	片剂 / 100	515420BT
铁 II LR (Fe ²⁺)	片剂 / 250	515421BT
铁 LR (Fe ²⁺ und Fe ³⁺)	片剂 / 100	515370BT
铁 LR (Fe ²⁺ und Fe ³⁺)	片剂 / 250	515371BT

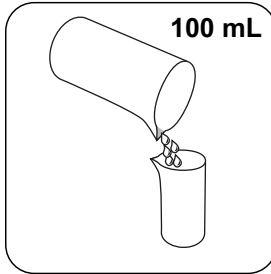
准备

1. 在必要情况下, 已用有机化合物作为腐蚀抑制剂处理的水须被氧化, 从而破坏铁复合物。为此, 将 100 ml 样本与 1 ml 浓硫酸和 1 ml 浓硝酸混合并蒸发至一半。冷却后进行消解。

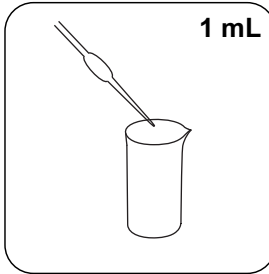
备注

1. 用这种方法测定总溶解的 Fe²⁺ 和 Fe³⁺。
2. 为了测定 Fe²⁺, 使用 IRON (II) LR 片剂代替 IRON LR 片剂。

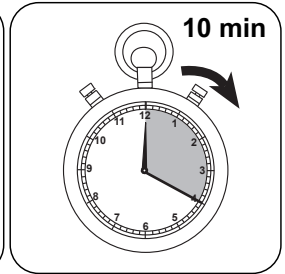
消解



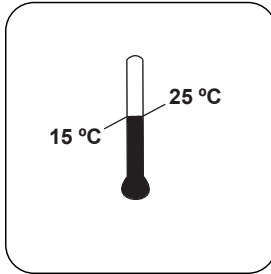
用 100 mL 样本填充合适的样本容器。



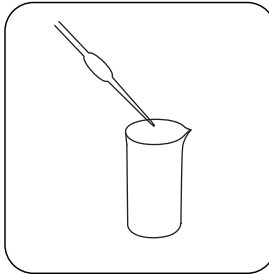
加入 1 mL 浓硫酸 ($\geq 95\%$)。



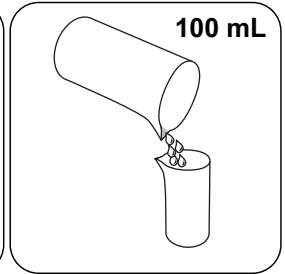
将样本加热 10 分钟，或一直加热直到其完全溶解。



将样本冷却到室温。



将样本的 pH 值从氨溶液 (10-25%) 调节到 3-5。



将样本用去离子水填充至 100 mL。

使用该样本来分析总溶解铁。

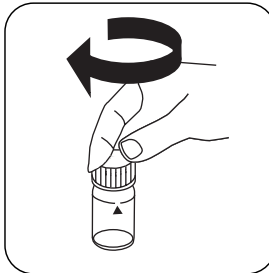
进行测定 铁 (II,III) ，用片剂溶解

选择设备中的方法。

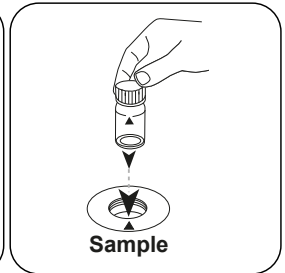
为了测定溶解和未溶解铁，进行中所述的消解。



用 10 mL 样本填充 24 mm 比色杯。



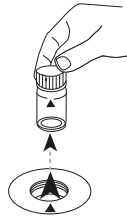
密封比色杯。



将样本比色杯放入测量轴中。注意定位。



Zero

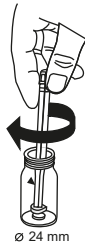


按下 **ZERO** 按钮。

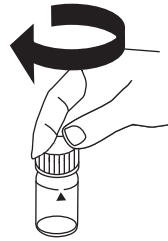
从测量轴上取下比色杯。



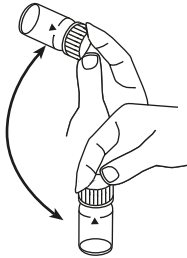
加入 **IRON LR** 片剂。



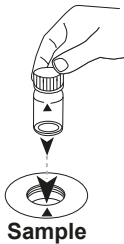
用轻微的扭转压碎片剂。



密封比色杯。



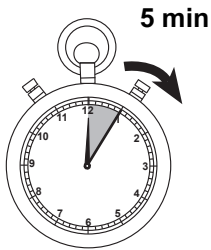
通过旋转溶解片剂。



将样本比色杯放入测量轴中。注意定位。

Test

按下 **TEST (XD: START)** 按钮。



等待 **5 分钟** 反应时间。

反应时间结束后，自动进行测量。

结果在显示屏上显示为 mg / l 铁。

化学方法

Ferrozine/巯乙酸盐

附錄

干扰说明

可消除干扰

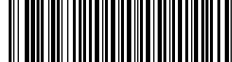
1. 铜的存在使测量结果增加了 10 %。样本中铜浓度为 10 mg/L 时，测量结果增加 1 mg/L 铁。
干扰可以通过添加硫脲来消除

方法验证

检出限	0.01 mg/L
测定下限	0.016 mg/L
测量上限	1 mg/L
灵敏度	0.92 mg/L / Abs
置信范围	0.013 mg/L
标准偏差	0.005 mg/L
变异系数	1.23 %

参考文献

Photometrische Analyse, Lange/ Vjedelek, Verlag Chemie 1980, S. 102



PP 铁

M222

0.02 - 3 mg/L Fe⁹⁾

FE1

1,10-邻菲罗啉

材料

所需材料 (部分可选) :

ZH

试剂	包装单位	货号
VARIO 铁 F10	粉剂 / 100 片	530560
VARIO 铁 F10	粉剂 / 1000 片	530563

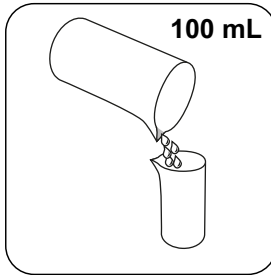
准备

1. 在分析前氧化铁需要一个弱、强或 Digesdahl 消解 (酸消解方法) 。
2. 在分析前应将强碱性或酸性水的 pH 值调节到 3 和 5 之间。
3. 对于含有可见锈的样本, 应保持至少 5 分钟的反应时间。
4. 在必要情况下, 已用有机化合物作为腐蚀抑制剂处理的水须被氧化, 从而破坏铁复合物。为此, 将 100 ml 样本与 1 ml 浓硫酸和 1 ml 浓硝酸混合并蒸发至一半。冷却后进行消解。

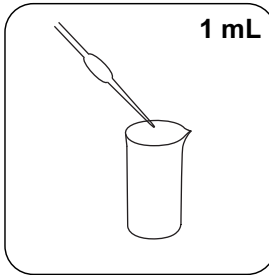
备注

1. 用这种方法测定所有形式的溶解铁和大部分形式的未溶解铁。
2. 准确度不会因未溶解的粉末而降低。

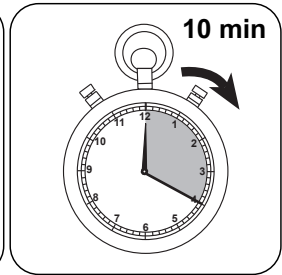
消解



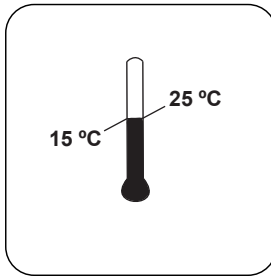
用 100 mL 样本填充合适的样本容器。



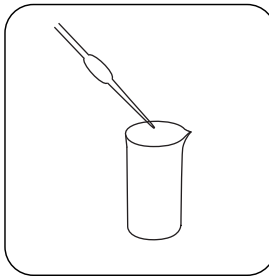
加入 1 mL 浓硫酸 ($\geq 95\%$)。



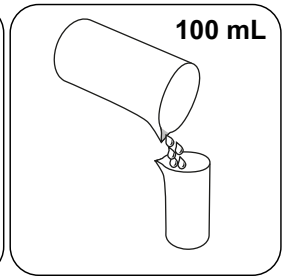
将样本加热 10 分钟，或一直加热直到其完全溶解。



将样本冷却到室温。



将样本的 pH 值从氨溶液 (10-25%) 调节到 3-5。



将样本用去离子水填充至 100 mL。

使用该样本来分析 总溶解铁。

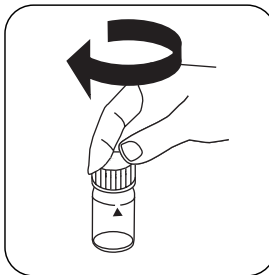
进行测定 铁 (II,III) ，用 Vario 粉包溶解

选择设备中的方法。

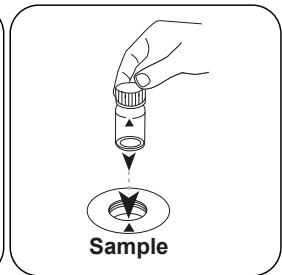
为了测定 铁 片剂法，进行 中所述的消解。



用 10 mL 样本填充 24 mm 比色杯。



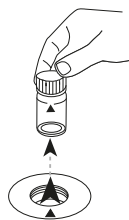
密封比色杯。



将样本比色杯放入测量轴中。注意定位。



Zero

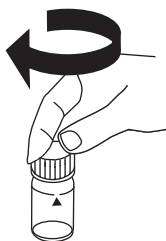


按下 **ZERO** 按钮。

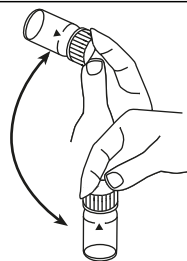
从测量轴上取下比色杯。



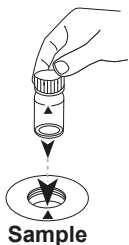
加入 **Vario FERRO F10** 粉包。



密封比色杯。



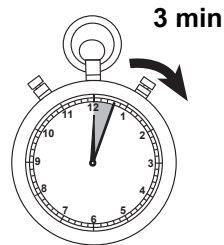
通过旋转混合内容物。



将样本比色杯放入测量轴中。注意定位。

Test

按下 **TEST (XD: START)** 按钮。



等待 **3 分钟** 反应时间。

反应时间结束后，自动进行测量。

结果在显示屏上显示为 **mg / l 铁**。



化学方法

1,10-邻菲罗啉

附录

干扰说明

持续干扰

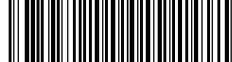
1. 铍干扰测量。

参照

DIN 38406-E1
标准方法 3500-Fe-1997
US EPA 40 CFR 136

⁹⁾ 试剂可测定大部分未溶解铁氧化物，无消解

ZH



铁 (TPTZ) PP

M223

0.02 - 1.8 mg/L Fe

FE2

TPTZ

材料

所需材料 (部分可选) :

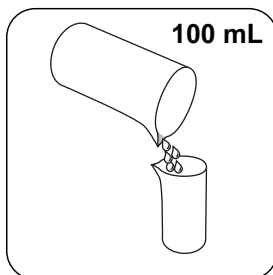
ZH

试剂	包装单位	货号
VARIO 铁 TPTZ F10	粉剂 / 100 片	530550

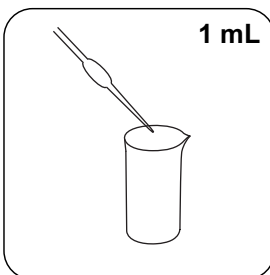
准备

1. 总铁的测定需要消解。TPTZ 试剂检测到大多数氧化铁不消解。
2. 在分析前用稀盐酸溶液 (1:1) 冲洗所有的实验室玻璃器皿, 然后用去离子水冲洗除去铁沉积物, 这可能导致结果轻微较高。
3. 在分析前 (用 0.5 mol/l 硫酸或 1 mol/l 氢氧化钠溶液) 应将强碱性或酸性水的 pH 范围调节到 3 和 8 之间。
4. 在必要情况下, 已用有机化合物作为腐蚀抑制剂处理的水须被氧化, 从而破坏铁复合物。为此, 将 100 ml 样本与 1 ml 浓硫酸和 1 ml 浓硝酸混合并蒸发至一半。冷却后进行消解。

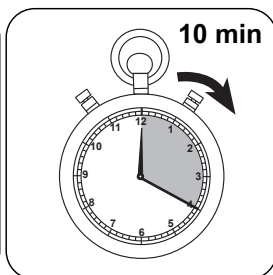
消解



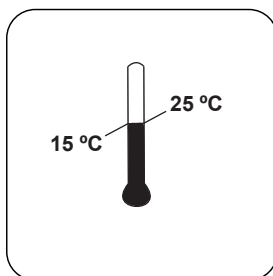
用 100 mL 样本填充合适的样本容器。



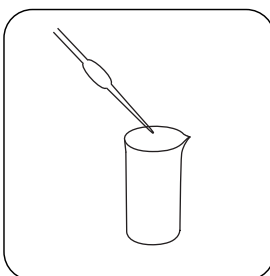
加入 1 mL 浓硫酸 ($\geq 95\%$)。



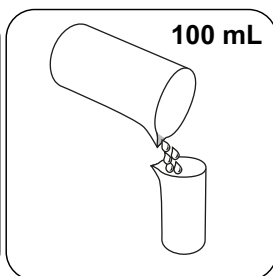
将样本加热 10 分钟，或一直加热直到其完全溶解。



将样本冷却到室温。



将样本的 pH 值从氨溶液 (10-25%) 调节到 3-5。



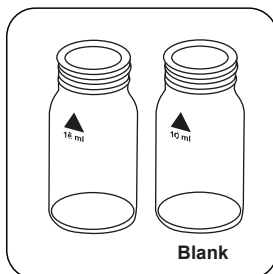
将样本用去离子水填充至 100 mL。

使用该样本来分析 总溶解铁。

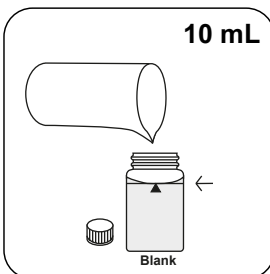
进行测定 铁，总 Vario 粉包

选择设备中的方法。

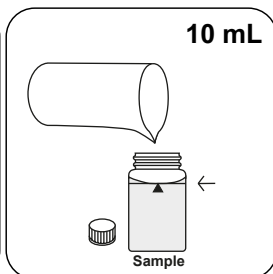
为了测定 总铁，进行 中所述的消解。



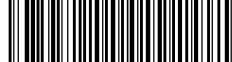
准备两个干净的 24 mm 比色杯。将一个比色杯标记为空白比色杯。



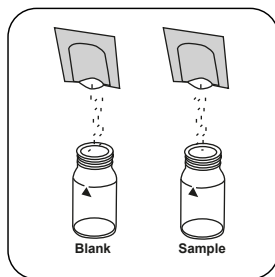
加入 10 mL 去离子水到比色杯中。



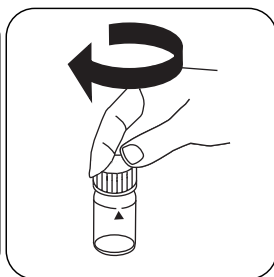
加入 10 mL 样本到样本比色杯中。



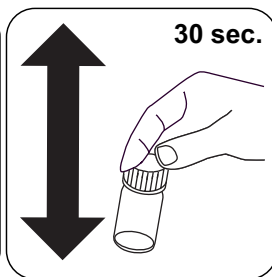
ZH



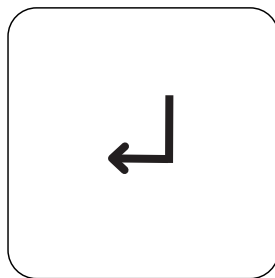
在每个比色杯中加入一个 Vario IRON TPTZ F10 粉包。



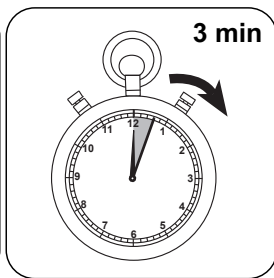
密封比色杯。



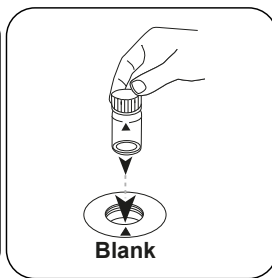
通过摇晃混合内容物 (30 sec.)。



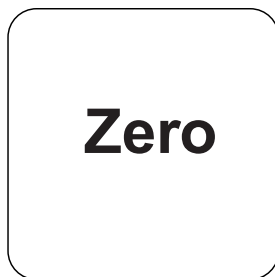
按下 ENTER 按钮。



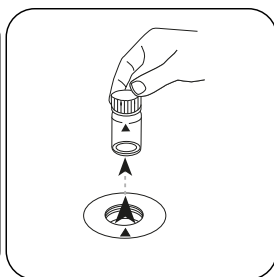
等待 3 分钟反应时间。



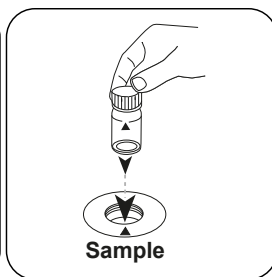
将空白比色杯放入测量轴中。注意定位。



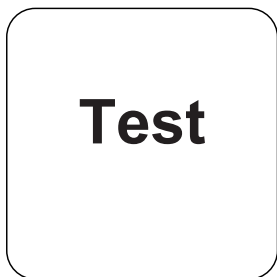
按下 ZERO 按钮。



从测量轴上取下比色杯。



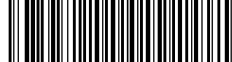
将样本比色杯放入测量轴中。注意定位。



按下 **TEST** (XD: **START**)
按钮。

结果在显示屏上显示为 mg / l 铁。

ZH



化学方法

TPTZ

附录

ZH

干扰说明

持续干扰

出现干扰时，颜色形成被抑制或形成沉淀。说明是指铁浓度为 0.5 mg/L 的标准。

干扰	限 / [mg/l]
Cd	4
Cr ³⁺	0.25
Cr ⁶⁺	1.2
Co	0.05
Cu	0.6
CN ⁻	2.8
Mn	50
Hg	0.4
Mo	4
Ni	1
NO ₂ ⁻	0.8

参考文献

G. Frederic Smith Chemical Co., The Iron Reagents, 3rd ed.(1980)

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