

Operating manual

AmmoLyt[®] 700 IQ



IQ SENSOR NET ammonium sensor

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Accuracy when going to press

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Note

The latest version of the present operating manual can be found on the Internet under <u>www.WTW.com.</u>

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1 Overview

1.1 How to use this component operating manual



Fig. 1-1 Structure of the IQ SENSOR NET operating manual

The IQ SENSOR NET operating manual has a modular structure like the IQ SENSOR NET itself. It consists of a system operating manual and the operating manuals of all the components used.

Please file this component operating manual in the ring binder of the system operating manual.





1.2 Structure of the AmmoLyt[®] 700 IQ ammonium sensor

Fig. 1-2 Structure of the AmmoLyt[®] 700 IQ ammonium sensor

1	Protective hood
2	Temperature probe
3	AmmoLyt [®] combination electrode with AmmoLyt [®] NHA reference electrode and AmmoLyt [®] NHA/AT replacement electrode (electrode not contained in the scope of delivery).
4	Receptacle for the AmmoLyt [®] combination electrode
5	Sensor shaft

Note

The ammonium electrodes that can be used are available as accessories (see chapter 6 REPLACEMENT PARTS AND ACCESSORIES).

Screening of the ammonium sensor

The AmmoLyt[®] 700 IQ ammonium sensor and the AmmoLyt combination electrode together with the IQ SENSOR NET system form a measuring system that is protected to a high degree against low and high frequency interference as well as against the indirect effects of lightning strikes.

1.3 Recommended fields of application

The AmmoLyt[®] 700 IQ ammonium sensor forms a measuring system for the online determination of ammonium ions. It supplements the oxygen measurement in the aeration tank and enables a more efficient aeration control.

The AmmoLyt[®] 700 IQ ammonium sensor, in conjunction with the AmmoLyt ammonium combination electrode, is suitable for stationary ammonium measurements in water/wastewater applications.

Safety 2

This component operating manual contains special instructions that must be followed in the operation of the AmmoLyt[®] 700 IQ ammonium sensor. Thus, it is essential to read this component operating manual before carrying out any work using this sensor. In addition to this manual, the SAFETY chapter of the IQ SENSOR NET system operating manual must be followed.

Always keep this component operating manual together with the system operating manual and any other component operating manuals in the vicinity of the IQ SENSOR NET system.

Special user The ammonium sensor was developed for applications in online qualifications measurement - essentially in the field of wastewater treatment. Thus, we assume that the operators are familiar with the necessary precautions to take when dealing with chemicals as a result of their professional training and experience.

General safety The individual chapters of this operating manual use the following instructions safety labels to indicate different levels of danger:

Warning

indicates instructions that must be followed precisely in order to prevent serious dangers to persons.

Caution

indicates instructions that must be followed precisely in order to avoid slight injuries or damage to the instrument or the environment.

Other labels

Note

indicates notes that draw your attention to special features.

Note

indicates cross-references to other documents, e.g. operating manuals.

2.1 Authorized use

The authorized use of the AmmoLyt[®] 700 IQ comprises its use as an ammonium sensor together with an ammonium combination electrode in the IQ SENSOR NET.

The technical specifications according to chapter 8 TECHNICAL DATA must be observed. Only operation according to the instructions given in this operating manual is considered to be authorized.

Any other use is considered to be **unauthorized**. Unauthorized use invalidates any claims with regard to the guarantee.

Caution

Only connect and operate the sensor together with IQ SENSOR NET accessories.

2.2 General safety instructions

The sensor left the factory in a safe and secure technical condition.

Function and operational safety

The failure-free function and operational safety of the sensor is only guaranteed if the generally applicable safety measures and the special safety instructions in this operating manual are followed during its use.

The failure-free function and operational safety of the sensor is only guaranteed under the environmental conditions that are specified in chapter 8 TECHNICAL DATA.

The specified temperature (chapter 8 TECHNICAL DATA) must be maintained during the operation and transport of the sensor. Protect the sensor, particularly against frost or overheating.

Caution

The sensor may only be opened by specialists authorized by WTW.

Safe operation If safe operation is no longer possible, the sensor must be taken out of operation and secured against inadvertent operation.

Safe operation is no longer possible if the sensor:

- has been damaged in transport
- has been stored under adverse conditions for a lengthy period of time
- is visibly damaged
- no longer operates as described in this manual.

If you are in any doubt, contact the supplier of your sensor.

Obligations of the
operatorThe operator of the sensor must ensure that the following rules and
regulations are followed when dealing with hazardous substances:

- EEC directives for protective labor legislation
- National protective labor legislation
- Safety regulations
- Safety data sheets of the chemical manufacturer.

3 Commissioning

3.1 Scope of delivery

- AmmoLyt[®] 700 IQ
- The sensor is fitted with a protective hood and protective caps
- Operating manual.

3.2 Installation

Connection cable

The SACIQ sensor connection cable is required to connect the sensor. Information on this and other IQ SENSOR NET accessories is given in the WTW catalog and on the Internet.

Note

How to connect the SACIQ sensor connection cable to the terminal strip of an MIQ module is described in chapter 3 INSTALLATION of the IQ SENSOR NET system operating manual.

Caution

The AmmoLyt[®] 700 IQ ammonium sensor unit may only be immersed in conjunction with a mounted combination electrode. Moisture must be prevented from penetrating the ammonium sensor during the replacement of the electrode as, otherwise, the sensor could be destroyed.

Which electrodes can be used in conjunction with the AmmoLyt[®] 700 IQ ammonium sensor is given in section 6.1 SENSOR AND ELECTRODES.

Are the plug connections dry?

Before connecting the sensor and sensor connection cable, please make sure that the plug connections are dry. If moisture gets into the plug connections, first dry the plug connections (dab them dry or blow them dry using compressed air).

Note

Do not suspend the sensor on the sensor connection cable. Use an armature or electrode holder. Information on this and other IQ SENSOR NET accessories is given in the WTW catalog and on the Internet.

Connecting the sensor to the sensor connection cable

- Take the protective caps off the plug connections of the sensor and the SACIQ sensor connection cable, and keep them safe.
 Dlug the isok of the SACIQ concert connection cable onto the
- 2 Plug the jack of the SACIQ sensor connection cable onto the plug head connector of the sensor. At the same time, rotate the socket so that the pin in the plug head connector (1) clicks into one of the two holes in the jack.
- 3 Then, screw the coupling ring (2) of the sensor connection cable onto the sensor up to the stop.

Fig. 3-1 Connecting the sensor

3.3 Commissioning / Getting the sensor ready for measuring

Mounting the combination electrode

1 Unscrew the protective hood from the sensor.

Pull off the blind plug from the plug head socket of the sensor.

3 Prepare the combination electrode to be mounted in the sensor.

To do so, screw the AmmoLyt[®] NHA/AT exchange electrode onto the AmmoLyt[®] NHA reference electrode without leaving any gap (see AmmoLyt[®] NHA/AT operating manual).

Note

2

The protection hood of the AmmoLyt $^{\ensuremath{\mathbb{R}}}$ 700 IQ sensor can be used as a wrench.

4 Grease the two sealing rings (1) of the AmmoLyt[®] electrode using the grease from the tube supplied.

5 Unscrew the protection cap from the plug head of the reference electrode.

6 Screw the reference electrode into the plug head socket of the sensor.

Caution

Push the electrode into the sensor right up to the stop so that the connection is watertight. Leaks could lead to the destruction of the sensor.

8 Screw the protective hood onto the sensor.

9	Carry out the settings for the sensor on the terminal of the measuring system (see section 3.4).
10	In the case of initial commissioning, condition the sensor toge- ther with the mounted combination electrode in a diluted stan- dard solution, e.g. ES/NH4_ISA-10 10 mg/l NH4-N for about 2 hours (see section 6.2).
11	Calibrate the sensor and electrode with the measuring system (see section 4.1).

3.4 Carrying out the settings for the sensor on the terminal of the IQ SENSOR NET system

The following settings can be carried out for the sensor:

Menu item	Possible settings	Explanations	
Measuring mode	 NH4 NH4-N mV 	The citation form of the mass concentrati- on or the voltage of the electrode.	
Measuring range (NH4)	 AutoRange 1 1290 mg/l 0.1 129.0 mg/l 	2 measuring ranges can be selected. With <i>AutoRange</i> , the instrument automati- cally switches to the suitable measuring range.	
Measuring range (NH4-N)	 AutoRange 1 1000 mg/l 0.1 100.0 mg/l 	2 measuring ranges can be selected. With <i>AutoRange</i> , the instrument automati- cally switches to the suitable measuring range.	
Measuring range (mV)	• -2000 2000 mV	Fixed range	
Temperaturmodus	● °C ● °F	Unit of the measured temperature value (Celsius, Fahrenheit).	

Menu item	Possible settings	Explanations
<i>Cal. procedure</i> (only in the <i>NH4</i> and <i>NH4-N</i> measuring	• 1 point standard (1)	• 1-point calibration with a standard solu- tion. The concentration of the standard solution must be entered.
mode)	• 1 point ref. (2)	 1-point calibration in the test sample with independent determination of the NH₄⁺ concentration by a reference measurement. The concentration of the test sample determined in a reference procedure must be entered.
	● 2 point stand. (3)	• 2-point calibration using any two WTW standard solutions. The concentration of the standard solutions must be entered.
	• Simple std. add. (4)	 A known concentration of standard solution is added to the sample. The NH₄⁺ concentration in the test sample is determined from the change in potential.
	● Double std. add. (5)	 A known quantity of a standard solution is added to the test sample in two steps. The NH₄⁺ concentration in the test sample is determined from the change in potential during the first and second standard addition.
		Note: The calibration procedures are described in detail in section 4.1 CALIBRATION.
<i>ORP offset</i> (only in <i>mV</i> measu- ring mode)	-100 mV +100 mV	You can set the voltage zero point here.
Initial calibration	On Off	An <i>Initial calibration</i> is required when the sensor is calibrated for the first time or when an electrode or the entire combination electrode has been replaced. During initial calibration, the basis for the evaluation of the drift potential is determined. Here you can select whether the next calibration should be an <i>Initial calibration</i> . After the initial calibration has been carried out, the setting for <i>Initial calibration</i> automatically switches to <i>Off.</i>

Menu item	Possible settings	Explanations	
Temp. adjustment	-1.5 °C +1.5 °C	The temperature compensation function enables the temperature sensor to be ba- lanced against a reference temperature measurement (displacement of the zero point by ± 1.5 °C).	
		 Notes: Due to the thermal capacity of the sensor, it is necessary to place it in a container with at least 2 liters of water. 	
		• Leave the sensor in this container for at least 15 minutes while stirring occasio- nally, then carry out the adjustment. If the temperature difference of the water and sensor is > 10°C, leave the sensor in the container for at least one hour while stirring occasionally.	
Potassium compens.	On Off	Potassium ions in the test sample interfere with the measurement and lead to increa-	
Potassium conc. (only with Potassium compens.: On)	0 100 mg/l	sed values (see section 4.2.2). After determining the potassium content of the test sample by a reference measure- ment you can input the determined potas- sium content here. The potassium compensation corrects the measured value accordingly.	
Save and quit		The system confirms the saving of the set- tings and the display switches to the next higher level.	
Quit		The display switches to the next higher level without saving the new settings.	

Carrying out settings

1	Switch to the measured value display with (M) .
2	Open the <i>Settings</i> menu with S.
3	Select and confirm the menu item <i>Settings of sensors and diff. sensors -> Measuring range</i> column with ③ and ④.
4	Select an entry with ③.

Teri	Terminal PC 🛛 01 Jan 2001 00:01 🖨 🛕 🛈						\odot
Set	Settings of sensors a			l diff.	sensors	:	140
&	No.	Sensor na	ame	Mea	suring (range	
	S01	99160001	NH	4-N Aut	oRange		
Select 🛠 edit sensor settings 🖞							

Fig. 3-2 140 - Settings of sensors and diff. sensors

5 Confirm the selection with K. The settings of the sensor are displayed.

Terminal PC	01 Jan	2001 00	10 2 🛆 🛈			
SO1 AmmoLy†7001Q 99160001						
Measuring mode		NH4-N				
Measuring range	•	AutoRange				
Temperature mod	le	°C				
Cal. procedure		1 point	standard (1)			
Temp. adjustmer	nt	0.0 K				
Potassium compe	ens.	Off				
Initial calibra	ation	Off				
Save and quit						
Quit						
Select setting 😘						

Fig. 3-3 140 - Settings of sensors and diff. sensors

- 6 Make the sensor settings with 0 and confirm each of them with 0.
- 7 Select the *Save and quit* menu item with O and confirm with OK. The new settings are stored in the sensor.

	4	Calibration and measuring		
	4.1	Calibration		
	4.1.1	General information		
Why calibrate?	When change charac curve i voltage	an ammonium electrode is operated its characteristic curve es with the course of time. The characteristic curve is generally terized by the slope and the axis intercept. The characteristic s the base for calculating the measured value from the electrode e.		
	With calibration, the current characteristic curve parameters are deter- mined as follows:			
	 With all calibration procedures, i.e. 1-point and 2-point procedures, the change of the axis intercept is determined ("drift potential"). 			
	 All t slop 	wo-point calibration procedures additionally determine the be.		
Why calibrate?	Calibra trode a	ate during the initial commissioning, after exchanging an elec- and at regular intervals (depending on the application).		
Calibration log and calibration history	The calibration history contains the calibration log of the initial and last of the following calibrations. You can call up the calibration history via the <i>Calibration history of selected sensor</i> display option.			
Citation form	There nium c nium c ammoi the set	are two common citation forms for the specification of ammo- ontents in a solution. The citation form NH4 specifies the ammo- ontent (NH_4^+). The citation form NH4-N specifies the amount of nium nitrogen only. You can select the required citation form in tings for the sensor.		
	If the c yourse	itation forms do not correspond, you can carry out the conversion If.		
	1 mg/l	NH4 = 0.777 mg/l NH4-N.		
	1 mg/l	NH4-N = 1.287 mg/l NH4.		

Note

Ordering information on ammonium standard solutions can be found in section 6.2.

4.1.2 Overview of the calibration procedures

For ammonium measurements with the AmmoLyt[®] 700 IQ sensor, the following calibration procedures can be selected:

1 point standard (1) 1-point calibration in a standard solution.

1 point ref. (2) 1-point calibration in the test sample. Calibration is carried out determining the ammonium concentration by an independent reference procedure, e.g. photometry.

- *2 point stand. (3)* 2-point calibration in two standard solutions.
- *Simple std. add. (4)* Calibration in the test sample with a one-time addition of standard solution.
- **Double std. add. (5)** Calibration in the test sample with a double addition of standard solution.

Determined calibration data

Depending on the calibration procedure (single-point or two-point), the following data are determined during calibrating:

Calibration procedu- res	Drift potential:	Slope
1 point standard (1)	*	
1 point ref. (2)	*	
2 point stand. (3)	*	*
Simple std. add. (4)	*	
Double std. add. (5)	*	*

In the calibration log and calibration history (section 4.1.10), the slope and drift potential are output. The drift potential and slope inform about the age condition of the electrode.

Note

Single-point calibration procedures take over the slope of the last twopoint calibration. If no valid data from a two-point calibration are available, the default setting (59.16 mV) is used. In both cases the value is marked with * in the calibration history.

4.1.3 Calibration in practice

Initial calibration The first calibration (initial calibration) is especially important as it is the reference point for all other calibrations (following calibrations).

An initial calibration is required each time an electrode is commissioned. The initial calibration is switched on and off in the setting menu of the sensor (see section 3.4).

With the initial calibration, the zero point for the drift potential is determined. It serves as the reference value for the drift potential, which is determined and recorded in the calibration log with every following calibration. In addition to the slope, the drift potential informs about the age condition of the electrode (see section 4.1.10).

Note

An optimum initial calibration is achieved by a calibration procedure that determines the current slope of the electrode (*2 point stand. (3)* or *Double std. add. (5)*). If the slope is not determined, the default setting (59.16 mV) is taken over. A value that is taken over is marked with * in the calibration history (see section 4.1.10).

Following the initial calibration, we recommend to carry out a calibration with the *1 point ref. (2)* procedure in order to compensate for matrix effects of the real test sample. Apart from that, the potassium content of the test sample should be determined and checked for whether a potassium compensation is required (see section 4.2.2).

Following calibrations Basically, any calibration procedure can be used for the following calibrations. The slope of the electrode should be determined at regular intervals in any case in order to be able to evaluate the aging of the electrode. If the electrode cannot be calibrated any more because the slope is too low it is blocked for measurement and has to be exchanged.

"Emergency operation" with invalid slope with invalid slope with a subsequent valid single-point calibration as a stopgap solution until the electrode is exchanged. The last valid slope is used in the measuring operation. With the single-point calibration, a corresponding note appears quoting the slope that is used.

Fig. 4-1 Sequence of calibrations

Preparatory activities

4.1.4 General course of a calibration

An optimum calibration result is possible by,

- before calibrating,
 - conditioning the sensor in the ES/NH4_ISA-10 standard solution with 10 mg/I NH4-N for approx. 10 minutes
 - setting the required calibration procedure (see section 3.4).
- providing a similar temperature of standard solution and test sample for calibration

Note

When exchanging an electrode or the entire combination electrode an initial calibration (see section 4.1.3) always has to be carried out.

Course

1	Switch to the measured value display with \bigcirc .
2	Select the measured value display of the required sensor with ③.
3	Call up the calibration with ①. The <i>Maintenance condition: Linked outputs are frozen.</i> window appears.

Termina	al 1	01	Sept	2001 11	1 18	3	\mathbb{A}	\odot
Calibra	ate sen	sor						200
01 M	⊑ ainter	<mark>со</mark> nance	cond	1 ition:	7.3	°C		
	inked	outpu	uts a	re fro	zen.			
03	ontinu	le						
	21	- . U	Turt	> 01	3440	000		
Cancel	calibr	ation	using	Esc.				

Fig. 4-2 Maintenance condition

Note

When the maintenance condition is activated, linked outputs remain in their current condition. In the measured value display, the measured value or the condition indicator of the sensor flashes.

Starting the calibration

4 Confirm with OK. The display appears with a note that an initial calibration may be necessary.

Term	inal PC 🛛 01 Jan 2001 00 11 🖨 🛕	0
Cali	brate sensor 2	200
01	∧∟ mg/L 25.0 °C Note: In case of electrode change perform initial cali-	
	bration. Before each calibra-	
03	tion, measure and enter potas- sium content (see settings)	
	Continue	
Canc	el calibration using Esc.	

Fig. 4-3 200 - Calibrate sensor

5 Confirm with OK. The following display (or similar, depending on the selected calibration procedure) appears:

Terminal PC 🛛 🛛 🛛 🛛 🛛 🖓 🖉 🚺 🕕			
Cal	brate sensor 200		
01	Cal.: 1 POINT STANDARD (1)		
02	Have standard ready for calibration		
03			
Cano	el calibration using Esc.		

Fig. 4-4 200 - Calibrate sensor

6 The further course depends on the respective calibration procedure. The individual operating steps are given in the sections 4.1.5 to 4.1.9.

Note

You can break off the calibration procedure at any time with the key. The system continues to work with the old calibration data. However, you have to switch off the maintenance condition in any case.

Note

The sensor determines a stable measured value with each measurement during the course of a calibration. The display shows a continuation display and the current electrode voltage in mV.

After calibrating, the following display appears:

Fig. 4-5 After calibrating

Completing the calibration	7	Confirm with 🔍. The measured value display appears again (the measured value flashes because the sensor is still in the maintenance condition).
	8	If the calibration was successful, immerse the sensor in the sample.
	9	Wait for a stable measured value.
	10	Switch off the maintenance condition (press 🕅 and, in the <i>Display/Options</i> menu, make the setting).
	11	Switch to the measured value display with (M) . The measured value no longer flashes.

Note

If the calibration was not successful, "----" appears on the display and a corresponding entry with instructions appears in the log book. Follow the instructions and repeat the calibration.

4.1.5 Calibration procedure, 1 point standard (1)

The *1 point standard (1)* 1-point calibration is carried out using one standard solution.

Note

Determine the potassium content of the sample (see section 4.2.2). Activate the *Potassium compens*. function and adjust the potassium content (see section 3.4).

The potassium content of the sample does not effect calibration but the measurement following.

Operating steps during calibration

Display	Explanation		
<i>Cal.: 1 POINT STANDARD (1)</i> <i>Have standard ready for calibra-</i> <i>tion</i>	You can use any standard solu- tion for this. The ammonium value should be as close to the value expected for the test sample as possible. Confirm with OK.		
<i>Select standard concentration</i> (1 / 10 / 100) mg/l <i>NH4-N</i>	If necessary select a standard with ②. Confirm with ④.		
* Rinse electrode	Follow the instructions on the display. Confirm with 🕅.		
* Immerse electrode in standard. * Wait for a stable measured value.	Confirm with 🕅. As soon as a stable measured value is reached, the next display appears.		
Calibration successful Conc. (NH4-N) x mg/l Slope y mV* Drift voltage z mV End of 1 POINT STAND. (1) cal.	The values for <i>Conc. (NH4-N)</i> , <i>Slope</i> and <i>Drift voltage</i> are dis- played. The slope is taken over from the last calibration that determined the slope. The calibration is complete. Con- firm with (K). The display returns to the measu- red value display.		

4.1.6 Calibration procedure, 1 point ref. (2)

the ammonium content in this calibration procedure.

The *1 point ref. (2)* 1-point calibration is carried out with the sample and in two main steps, each of which is started with ⓒ.

Note

Note

Determine the calibration value and take the sample before the start of the aeration in the aeration tank. The ammonium concentration is highest before the aeration. After the aeration the ammonium content can decrease to a value that lies within the range of the detection limit. It does not make sense to calibrate in this range.

Determine the potassium content together with the reference value for

Operating steps during step 1

Display	Explanation
<i>Cal.: 1 POINT REFERENCE (2)</i> <i>Step 1: Determining the refe-</i> <i>rence voltage</i>	Confirm with 🕅.
 * Rinse electrode. * Immerse electrode in sample. * Observe conditioning time of 15 minutes. 	Before starting the calibration the sensor should be immersed in the test sample for conditioning for at least 15 min. Start the measurement with OK. This step determines and stores the reference voltage. As soon as a stable measured value is recognized, the next dis- play appears.
Step 1 finished. Switch to the measured value display with 2x 'OK'. After det. the reference concentration, start calib. step 2 with 'C'.	Step 1 of the calibration is fini- shed. Press \textcircled{K} <u>twice</u> to switch to the measured value display. The sensor is in the maintenance con- dition.

Note

During the subsequent determination of the reference concentration in the laboratory you can use the sensor for measuring again by simply abolishing the maintenance condition.

The sensor continues to use the old calibration data. The reference voltage determined in step 1 of the calibration is <u>not</u> lost. It remains stored until step 2 of the calibration is completed. It does not have to be noted and entered again.

Sampling and determining the reference concentration

You have to be in the measured value display to continue the calibration. Continue with sampling and determining the reference concentration as follows.

1 Take a sample.

Note

The ammonium content has to be determined immediately after taking the sample as the ammonium content changes very quickly due to the micro organisms that are present. It is best to take the sample using a syringe filter for transport to the laboratory or to stabilize it otherwise. When adding stabilizing solutions, the dilution factor has to be taken into account.

2	Determine the concentration of ammonium and potassium in the laboratory. With this procedure, the potassium content of the sample affects calibration as well as measurement.
3	 In the Settings of sensors and diff. sensors menu (see section 3.4) Switch on the Potassium compens. function and enter the potassium content.
4	Continue the calibration as follows with \bigcirc .

Operating steps during	Display	Explanation
step 2	<i>Cal.: 1 POINT REFERENCE (2)</i> <i>Step 2: Enter reference concen-</i> <i>tration</i> <i>Reference voltage already deter-</i> <i>mined</i>	Confirm with 🔍.
	Continue with Input ref. conc. New calibration	Here you can select whether you want to repeat step 1 of the calib- ration (New calibration), or enter the reference concentration (Input ref. conc.). If necessary select the step with (Confirm with (K).
	Input reference concentration Citation form/Value range	Confirm with 🔍.

Citation form of ref. conc. NH4(0.1129.0mg/l) NH4(11290mg/l) NH4N(0.1100.0mg/l) NH4N(11000mg/l)	If necessary select the citation form with ③. Confirm with [®] .	
Input reference concentration Value determined	Confirm with 🛞.	
Value of ref. concentration x mg/l NH4-N	If necessary set the reference concentration determined with ③. Confirm with ⓒ.	
Calibration successful Conc. (NH4-N) x mg/l Slope y mV* Drift voltage z mV End of 1 POINT REF. (2) cal.	The values for <i>Conc. (NH4-N)</i> , <i>Slope</i> and <i>Drift voltage</i> are dis- played. The slope is taken over from the last calibration that determined the slope. The calibration is complete. Con- firm with OK. The display returns to the measu- red value display.	

4.1.7 Calibration procedure, 2 point stand. (3)

The 2 point stand. (3) two-point calibration is carried out with two standard solutions of different concentrations. The procedure includes two conditioning processes. Each conditioning process takes 15 min and is important so the temperatures of standard solution and electrode can adjust to each other and a stable concentration can be reached at the electrode membrane. During the conditioning process the remaining time is displayed on the screen. The process cannot be shortened. By pressing (13) or (13) you cancel the entire calibration and the old calibration data is still used. The time required for the entire calibration is approx. 40 min.

Note

Determine the potassium content of the sample (see section 4.2.2). Activate the *Potassium compens*. function and adjust the potassium content (see section 3.4).

The potassium content of the sample does not effect calibration but the measurement following.

If you carry out a *1 point ref. (2)* calibration directly after the *2 point stand. (3)* calibration it is sufficient to determine and adjust the potassium content during the *1 point ref. (2)* calibration (matrix alignment) (see section 4.1.6).

Operating steps during	Displays	Explanation
canbration	Unscrew protective hood. Clean and rinse sensor incl. elec- trode and protective hood, then reassemble them.	Prepare the sensor as described. After completing the operating steps confirm with 🕅.
	<i>Cal.: 2 POINT STANDARD (3)</i> <i>Have standard 1 ready for calibra-</i> <i>tion</i>	Confirm with 🕅.
	Immerse sensor in standard 1. Observe minimum immersion depth (70mm).	Confirm with 🕅.
	Select standard concentration (1 / 10 / 100) mg/l NH4-N	If necessary select another con- centration of the standard with (.). Confirm with (K). Subsequently wait for the condi- tioning time of 15 minutes to expire. The remaining time is shown on the display.
	Discard used standard. Immerse sensor in new standard with simi- lar concentration. Start calibration.	Confirm with 🕅.
	Cal.: 2 POINT STANDARD (3) Calibration values for standard 1 determined Have standard 2 ready	Confirm with 🕅.
	Rinse sensor with standard 2. Immerse sensor in standard 2. Observe minimum immersion depth (70mm).	Confirm with 🕅.
	Select standard concentration (1 / 10 / 100) mg/l NH4-N	If necessary select the concen- tration of the standard with (). Confirm with (). Subsequently wait for the condi- tioning time of 15 minutes to expire. The remaining time is shown on the display.
	Discard used standard. Immerse sensor in new standard with simi- lar concentration. Start calibration.	Confirm with 🛞.

Displays	Explanation	
<i>Cal.: 2 POINT STANDARD (3)</i> <i>Calibration values for standard 2</i> <i>determined</i>	Confirm with 🛞.	
Calibration successful Conc. (NH4-N) x mg/l Slope y mV Drift voltage z mV End of 2 POINT STAND. (3) cal.	The values for <i>Conc. (NH4-N)</i> , <i>Slope</i> and <i>Drift voltage</i> are dis- played. The calibration is complete. Con- firm with OK. The display returns to the measu- red value display.	

4.1.8 Calibration procedure, Simple std. add. (4)

This calibration with a simple standard addition is carried out in the test sample while adding standard. The volumes of the test sample and standard have to be dosed exactly.

The following appliances are suitable for dosing:

- Measuring cylinder to determine the volume of the sample
- Pipette (microliter pipette as necessary) for accurate dosing of the standard. The higher the concentration of the standard, the more important it is to measure the amount of calibration standard very carefully.

Note

Before calibrating, determine the potassium content of the sample (see section 4.2.2).

Before calibrating, activate the *Potassium compens*. function and adjust the potassium content (see section 3.4).

The potassium content of the test sample affects the calibration result.

Observe the following points for the simple standard addition:

- The volume of the standard added to the test sample changes the test sample. Therefore, the volume of the standard should not be more than approx. 1 % of the volume of the sample.
 <u>Example:</u> With 100 ml sample volume, the added volume should not be more than 1 ml.
- Through the addition of standard solution, the ammonium content of the test sample should be increased at least twofold but not more than tenfold.

Based on these conditions, the range that can be calibrated is from 0.17 to 76 mg/l NH4-N (or 0.22 to 99 mg/l NH4). If the expected measured values are outside this range, a different calibration procedure has to be selected.

Conditions and calibration range for simple standard addition

Concentration and quantity of standard solution

The concentration and quantity of the standard solution to be added depend on the measuring range expected. They can be determined by means of the following diagram. The requirements for the simple standard addition are automatically met while doing so.

 $V_P:V_S$ is the volume ratio between the test sample and standard to be added. If several combinations for concentration/volume ratio (bars) are possible for the determination, select the bar where the expected NH4-N concentration is more in the center.

Example:

The expected average concentration is approx. 3.5 mg/l NH4-N. The vertical line in the diagram crosses three ranges:

- Concentration 1 g/l and volume ratio 200:1
- Concentration 1 g/l and volume ratio 100:1
- Concentration 10 g/l and volume ratio 500:1

Selection of the range: Basically, all of the three ranges are suitable. The lower bar is crossed by the line on the right edge only. Therefore, the other two ranges are better suited. For further selection, decide which bar covers the expected measuring range better, i.e. whether the expected values are rather lower or higher than the average value. Besides, the decision can be made according to practical points of view (availability of standard solution and suitable dosing equipment). Entry of volumes

The volume of standard to be added in this calibration procedure has to be entered on the terminal of the IQ SENSOR NET with an accuracy of 1/10 ml. Therefore, the volume has to be rounded up or down accordingly.

Note

Make sure the citation form is correct for all specifications of the concentration.

ring	Display	Explanation
tion	<i>Cal.: SIMPLE STD. ADD. (4)</i> <i>Have test sample ready for calibra-</i> <i>tion</i>	Confirm with 📧.
	Input sample volume (100 1000) ml	If necessary select the volume of the sample with ③. Confirm with ⓒ.
	* Wait for a stable measured value.	Confirm with 🔍.
	<i>Cal.: SIMPLE STD. ADD. (4)</i> <i>Reference voltage of test sample</i> <i>determined</i> <i>Have standard ready</i>	Confirm with K. The measurement of the test sample begins.
	Select standard concentration (1 / 10) g/l NH4	If necessary select the concen- tration of the standard with ④. Confirm with ⓒ.
	Input standard volume (0.1 20.) ml	If necessary select the volume of the standard with ③. Confirm with 0K.
	<i>Cal.: SIMPLE STD. ADD. (4)</i> <i>Add standard to test sample</i>	Add the quantity of standard that was entered to the test sample. Subsequently confirm with K.
	* Wait for a stable measured value.	Confirm with 🔍.
	Cal.: SIMPLE STD. ADD. (4) Reference voltage determined after standard addition	Confirm with 📧.
	Calibration successful Conc. (NH4-N) x mg/l Slope y mV* Drift voltage z mV End of SIMPLE STD ADD (4) cal.	The values for <i>Conc. (NH4-N)</i> , <i>Slope</i> and <i>Drift voltage</i> are dis- played. The calibration is complete. Confirm with ^(K) . The display returns to the measured value display.

Operating steps during calibration

4.1.9 Calibration procedure, *Double std. add. (5)*

This calibration with double standard addition is carried out in the test sample while adding standard with a certain concentration in two steps. The volumes of test sample and standard have to be dosed exactly.

The following appliances are suitable for dosing:

- Measuring cylinder to determine the volume of the sample
- Pipette to exactly dose the standard

Note

Before calibrating, determine the potassium content of the sample (see section 4.2.2).

Before calibrating, activate the *Potassium compens.* function and adjust the potassium content (see section 3.4).

The potassium content of the test sample affects the calibration result.

Concentration and quantity of standard solution

The quantity of standard solution to be added is permanently fixed for the double standard addition:

- First addition: 1 % of the sample volume
- Second addition: 2 % of the (original) sample volume

After both additions, the original ammonium content of the test sample should be increased at least twofold but not more than tenfold.

Based on these conditions, the range that can be calibrated is from 2.5 to 220 mg/l NH4-N (or 3.2 to 283 mg/l NH4). If the expected measured values are outside this range, a different calibration procedure has to be selected.

The concentration of the standard solution to be added depends on the measuring range expected. It can be determined by means of the following diagram. The requirements for the double standard addition are automatically met while doing so.

Note

Operating steps during calibration

Make sure the citation form is correct for all specifications of the concentration.

Display	Explanation
Cal.: DOUBLE STD. ADD. (5) Have test sample ready for cali- bration	Confirm with 🕅.
<i>Input sample volume</i> (100 1000) ml	If necessary select the volume of the sample with \textcircled{O} . Confirm with \textcircled{OK} .
* Rinse electrode. * Immerse electrode in test sam- ple. * Wait for a stable measured value.	Confirm with (). The measure- ment of the test sample begins.
<i>Cal.: DOUBLE STD. ADD. (5) Reference voltage of test sample determined Have standard ready</i>	Confirm with 🕅.
Select standard concentration (1 / 10) g/l NH4	If necessary select the concen- tration of the standard with ①. Confirm with ④.
Add standard to sample	Confirm with 🛞.
* Wait for a stable measured value.	Confirm with 🕅. The measurement begins.

Display	Explanation
Cal.: DOUBLE STD. ADD. (5) Reference voltage determined after first standard addition	Confirm with 📧.
Add standard to sample	Confirm with 🛞.
* Wait for a stable measured value.	Confirm with 🔍. The measurement begins.
Cal.: DOUBLE STD. ADD. (5) Reference voltage determined after second standard addition	Confirm with 📧.
Calibration successful Conc. (NH4-N) x mg/l Slope y mV Drift voltage z mV End of DOUBLE STD ADD (5) cal.	The values for <i>Conc. (NH4-N)</i> , <i>Slope</i> and <i>Drift voltage</i> are dis- played. The calibration is complete. Con- firm with K. The display returns to the measured value display.

4.1.10 Calibration result

Calibration evaluation After calibrating the system automatically evaluates the calibration data and current state of the sensor. The drift potential and slope are evaluated separately. For a valid calibration, the values have to be within the following ranges:

Drift potential: -45 ... +45 mV

Possible results of the

calibration

Display after the calib- ration	Log book entries (meaning/actions)
Measured value display	Sensor was successfully calibrated. Slope and drift potential are within the valid range. For calibration data, see calibration history.
""	Sensor could not be calibrated. Slope and/or drift potential are outside the valid range. Sensor blocked for measurement.
	 Service the sensor immediately (see chapter 5).
	 View the calibration history.
	 Check the calibration conditions and calibration standard.
	Note: After determining an invalid slope, the sen- sor can be further operated with a subse- quent valid single-point calibration as a stopgap solution until the electrode is exchanged. The last valid slope is used in the measuring operation. With the single- point calibration, a corresponding note appears quoting the slope that is used.

A calibration can have the following results:

Note

Viewing the calibration data

Information on the contents and structure of the log book and how you can call it up is given in the LOG BOOK chapter of the IQ SENSOR NET system operating manual.

The calibration data can be viewed as follows via the *Calibration history* display option.

1	Select the relevant sensor in the measured value display on the terminal.
2	Press the 🛞 key. The <i>Display/Options</i> menu appears.
3	Select the <i>Calibration history of selected sensor</i> menu item. The data of the last calibrations of the sensor appear on the display.

Calibration history

Fig. 4-6 330 - Calibration history of selected sensor

The calibration history contains the following information:

Date	Date of the calibration
S(*)	Slope [mV] of the electrode The value is marked by * Values unchanged if it is not possible to determine the slope. Instead, the last determined value is taken over for the slope. This applies to all 1 point calibration procedures, 1 point standard (1), 1 point ref. (2) and Simple std. add. (4).
DV	Drift potential [mV]
Ref1/Ref2	 Concentration [mg/l NH4-N] of the standard with 1 point standard (1) of the test sample (reference measurement) with 1 point ref. (2) of the two standards with 2 point stand. (3) of the test sample (calculated) with Simple std. add. (4) [Ref1/-] of the test sample (calculated) with Double std. add. (5) [Ref1/- Ref2]
К+	Potassium content [mg/l]
Р	Calibration procedure, number 1 5
Τ	Temperature [°C]
R	Result of the calibration + : Calibration successful - : Calibration unsuccessful

Note

Only calibration data produced by the same calibration procedure can be compared.

4.2 Measuring

- 1 Submerse the sensor with the mounted combination electrode in the sample.
- 2 Read the measured value on the terminal of the IQ SENSOR NET system.

Note

Please pay attention to:

- the minimum immersion depth of the sensor (> 70 mm)
- the measuring range of the electrode used (see operating manual of the electrode).

Note

To keep the sensor clean, we strongly recommend to use the CH cleaning head (see chapter 6 REPLACEMENT PARTS AND ACCESSORIES).

4.2.1 Influence of the pH value

pH value > 7 The pH value of the test sample affects the chemical balance between NH_4^+ and NH_3 . From a pH value of 7, the NH_4^+ part of the test sample decreases and the NH_3 part increases with a rising pH value. NH_3 is not measured by the AmmoLyt[®].

Fig. 4-7 Influence of pH value and temperature (T) on the NH₄⁺ part in the test sample

For measurements in test samples with a pH value > 7, a compensation for the influence of the pH value may be necessary. The compensation can, e. g. be performed using the superior process control engineering.

Note

For details, please refer to WTW GmbH.

4.2.2 Potassium compensation

Measuring ammonium in the presence of potassium leads to increased ammonium values:

Potassium con- tents	Ammonium value increased by approx.
10 mg/l K ⁺	0.7 mg/l
50 mg/l K ⁺	3.4 mg/l

Higher measured values caused by potassium can be compensated for by switching on the *Potassium compens.* function and entering the potassium content in the *Settings of sensors and diff. sensors* menu (see section 3.4).

Potassium compensation for calibration The presence of potassium affects the calibration results of the following calibration procedures:

- 1 point ref. (2)
- Simple std. add. (4))
- Double std. add. (5)

For these calibration procedures, the current potassium content must always be determined and entered before or during calibration. The entered potassium compensation has an effect on the calibration result and thus on the subsequent measurements.

The calibration procedures, *1 point standard (1)* and *2 point stand. (3)* are carried out in standard solutions. Potassium does not interfere here. An incorrectly set potassium compensation affects the following measurement.

Potassium compensation for measurements

The function, *Potassium compens.*, should always be used if the concentration of the interfering ion causes the measurement signal to leave the range of the required accuracy. To avoid calibration and measuring errors, we recommend to

- routinely determine the potassium contents
- switch on the *Potassium compens.* function
- enter the potassium content.

The more the actual potassium content deviates from the value entered for the potassium compensation, the more the displayed measured value deviates from the current ammonium content.

Fig. 4-8 shows the interrelationship of real and displayed NH4-N value for different potassium contents. The characteristic curve with the optimum potassium compensation corresponds to the characteristic curve without any potassium content.

c (NH4-N) [mg/l] displayed by AmmoLyt 700 IQ

Fig. 4-8 Influence of potassium on the measured ammonium value

4.2.3 Further influences on the measured value

Greases, oils, certain tensides and similar substances can shorten the operational lifetime of the AmmoLyt[®] NHA/AT exchange electrode. Therefore, they should not be present in the test sample.

5 Maintenance and changing the electrode

The AmmoLyt[®] 700 IQ ammonium sensor is maintenance-free.

Warning

Contact with the sample can be dangerous for the user! Depending on the type of sample, suitable protective measures must be taken (protective clothing, protective goggles, etc.).

Note

To keep the sensor clean, we strongly recommend to use the CH cleaning head (see chapter 6 REPLACEMENT PARTS AND ACCESSORIES).

Note

Please read the maintenance of the combination electrode in the relevant operating manual of the electrode.

5.1 Exchanging the electrode

Note

We do not recommend unscrewing the measuring electrode from the sensor connection cable when changing the electrode. Otherwise, moisture and/or dirt can get into the plug connection where it can cause contact problems.

If you need to disconnect the sensor from the sensor connection cable, please note the following points:

- Before disconnecting the sensor from the SACIQ sensor connection cable, remove any larger pieces of contamination from the sensor, particularly in the area of the plug connection (brush it off in a bucket of tap water, wash it off with a hose or wipe it off with a cloth).
- Unscrew the sensor from the SACIQ sensor connection cable.
- Always place a protective cap on the plug head of the sensor and on the SACIQ sensor connection cable so that no moisture or dirt can get into the contacting surfaces.

Replacing the electrode

If it is necessary to replace an electrode, proceed as follows:

1 Unscrew the protective hood from the sensor.

2 Use the protective hood as a tool to lever out the electrode.

3 Carefully pull out the electrode until the plug head screwed fitting can be seen.

4 Unscrew the electrode from the plug head socket (for disposal, see section 5.2).

5 Screw in a new electrode.

- 6 Pu
- Push the unit into the sensor up to the stop.

7 Screw the protective hood onto the sensor.

8 Calibrate the sensor and the electrode with the measuring system (see section 4.1 CALIBRATION).

5.2 Disposal

Sensor We recommend disposing of the sensor as electronic refuse.

Electrodes If no official regulations apply to the contrary, used and defective electrodes can be treated as household waste.

6 Replacement parts and accessories

6.1 Sensor and electrodes

Ammonium sensor	Model	Order no.
	AmmoLyt [®] 700 IQ	107002

Ammonium electrodes	Model	Order no.
	Reference electrode AmmoLyt [®] NHA	107004
	Exchange electrode AmmoLyt [®] NHA/AT	107006

6.2 General accessories

Protective hood	Designation	Order no.
	AmmoLyt [®] 700 IQ-SK	107016

For the calibration procedures, *1 point standard (1)* and *2 point stand. (3)*:

Designation		Order no.
ES/NH4_ISA-1	1 mg/l NH4-N; 1000 ml	107010
ES/NH4_ISA-10	10 mg/l NH4-N; 1000 ml	107012
ES/NH4_ISA-100	100 mg/l NH4-N; 1000 ml	107014

For the calibration procedures, *Simple std. add. (4)* and *Double std. add. (5)*:

Designation		Order no.
SL NH4 19812	1000 mg/l NH4; 500 ml	250461
ES/NH4	1000 ml; 10 g/L NH4	120240

Standard solutions for

ammonium calibration

Cleaning system	Model	Order no.
	CH Cleaning Head	900107
	MIQ/CHV Valve Module	900109

Note

Information on other IQ SENSOR NET accessories is given in the WTW catalog and on the Internet.

7 What to do if...

Cause	Remedy
 Sensor not connected 	 Connecting the sensor
– Unknown	 Look in the log book

Measurement does not function

Cause	Remedy
 Electrode not connected 	 Connect electrode
 Liquid has penetrated the sensor 	 Sensor defective, send it back
 Sensor not connected 	 Connecting the sensor
 Instrument setting incorrect 	 Correct instrument setting

Measurement provides implausible measured values

Cause	Remedy	
 No calibration performed 	 Calibrate 	
 Electrode not connected or defective 	 Check electrode and electrode connection 	
 Electrode contaminated 	 Clean electrode 	
 Liquid has penetrated the sensor 	 Sensor defective, send it back 	
 Instrument setting incorrect 	 Correct the instrument setting (<i>Measuring mode</i> mg/l or mV) 	
 Potassium compensation is switched off 	 Switch on potassium compen- sation 	
 Potassium compensation with unsuitable value for potassium content 	 Determine and enter potassium content Recalibrate (see also section 4.2.2) 	

System cannot be	Cause	Remedy
Calibrated	 Slope of the electrode not within tolerance (see section 4.1.10) 	 Condition the electrode If the slope is still outside the tolerance: Replace the electrode
	 Drift of the electrode too high 	 Replace the electrode

8 Technical data

8.1 Measuring properties

Measuring principlePotentiometric measurement with an ammonium-selective combination electrode;
Integrated microprocessor electronics, shielded 2-wire connection for power and data transmission.

Measuring ranges and resolutions	Measuring mode	Measuring rar	nge	Resolution	
	NH4 ⁺	0.1 129.0 mg/l 1 1290 mg/l		0.1 mg/l 1 mg/l	
	NH4-N	0.1 100.0 mg/l 1 1000 mg/l		0.1 mg/l 1 mg/l	
	Voltage	-2000 2000	mV	1 mV	
Temperature	Probe type inte		integrated NTC	integrated NTC	
measurement	Measuring range		- 5 °C + 40 °C	- 5 °C + 40 °C (23 104 °F)	
	Accuracy		± 0.5 K		
	Resolution		0.1 K		
	Response time t ₉₉		< 15 s		
Temperature compensation	in the range of 0 °C + 40 °C (32 104 °F)				
	8.2 Operating characteristics				
Allowed temperature range	Measuring medium		0 °C + 40 °C (32 104 °F)		
	Storage/transport		With electrodes mounted: 0 °C + 40 °C (32 104 °F)		
			With electrodes -5 + 65 °C (2	s removed: 3 149 °F)	
Allowed pH range of the measuring medium	4 12 (pH range for measurement: 4 8.5)				
Pressure resistance	Sensor with electr	ode and SACIQ	sensor connection	n cable connected:	
	Max. allowed over	rpressure	2 x 10 ⁵ Pa (0.2	bar)	

Depth of immersion	min. 70 mm; max. 2 m depth
Operating position	pendulous to horizontal
Field of application	Controlling / monitoring in the aeration tank of waste water treatment plants, water and waste water monitoring

8.3 General features

8.4 Electrical data

Nominal voltage	Max. 24 VDC via the IQ SENSOR NET (for more details, see TECHNICAL DATA chapter of the IQ SENSOR NET system operating manual)
Power consumption	0.2 W
Protective class	111

9 Indexes

9.1 Explanation of the messages

This chapter contains a list of all the message codes and related message texts that can occur in the log book of the IQ SENSOR NET system for the AmmoLyt[®] 700 IQ sensor.

Note

Information on

- the contents and structure of the log book and
- the structure of the message code

is given in the LOG BOOK chapter of the IQ SENSOR NET system operating manual.

Note

All message codes of the AmmoLyt $^{\mbox{\scriptsize R}}$ 700 IQ end with the number "351".

9.1.1 Error messages

Message code	Message text
EA1351	<i>Meas. range exceeded or undercut * Check process * Select other meas. range</i>
EA2351	Sensor temperature too high! * Check process and application
EA3351	Sensor temperature too low! * Check process and application
EC1351	Sensor could not be calibrated, Sensor blocked for measurement * Check calibration conditions and calibration standard * View calibration history * Service sensor immediately (see operating manual)
ES1351	Component hardware defective * Contact WTW

9.1.2 Info messages

Message code	Message text
IC1351	Sensor has be * For calibration

Sensor has been successfully calibrated * For calibration data, see calibration history

9.2 Status info

The status info is a coded piece of information on the current status of a sensor. Each sensor sends this status info to the controller. The status info of sensors consists of 32 bits, each of which can have the value 0 or 1.

0 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15

Status info, general structure

		_
1 0 0 0 0 0 0 0 0	000000000	(general)
0 0 0 0 0 0 0 0 0	000000000	(internal
16 17 18 19 20 21 22 23	24 25 26 27 28 29 30 31	

The bits 0 - 15 are reserved for general information. The bits 16 - 21 are reserved for internal service information.

You obtain the status info:

- by a manual query in the *Settings/Service/List of all components* menu (see system operating manual)
- by an automated query

1

- of a superordinate process control (e. g. when connected to the Profibus)
- of the IQ Data Server (see IQ SENSOR NET Software Pack operating manual)

Note

The evaluation of the status info, e.g. in the case of an automated query, has to be made individually for each bit.

Status info AmmoLyt[®] 700 IQ

Status bit	Explanation
Bit 0	Component hardware defective
Bit 1-31	-