

Multiparameter Photometer for Boiler & Cooling Tower





Dear
Customer,Thank you for choosing a Hanna Instruments product.
Please read this instruction manual carefully before using this instrument.
This manual will provide you with the necessary information for correct use of this
instrument, as well as a precise idea of its versatility.
If you need additional technical information, do not hesitate to e-mail us at
tech@hannainst.com or view our worldwide contact list at www.hannainst.com.

All rights are reserved. Reproduction in whole or in part is prohibited without the written consent of the copyright owner, Hanna Instruments Inc., Woonsocket, Rhode Island, 02895, USA.

TABLE OF CONTENTS

TABLE OF CONTENTS

1.	PRELIMINARY EXAMINATION	6
2.	SAFETY MEASURES	6
3.	SPECIFICATIONS	7
4.	ABBREVIATIONS	8
5.	DESCRIPTION	9
	5.1. GENERAL DESCRIPTION & INTENDED USE	9
	5.2. PRECISION & ACCURACY	9
	5.3. FUNCTIONAL DESCRIPTION	10
	5.4. PRINCIPLE OF OPERATION	11
	5.5. OPTICAL SYSTEM	12
6.	GENERAL OPERATIONS	13
	6.1. POWER CONNECTION & BATTERY MANAGEMENT	13
	6.2. MODE SELECTION	
	6.3. GENERAL SETUP	14
	6.4. CONTEXTUAL HELP	17
7.	LOGGING DATA & DATA MANAGEMENT	18
	7.1. LOGGING DATA	18
	7.2. ADDING SAMPLE & USER NAMES TO LOG DATA	18
	7.3. DATA MANAGEMENT	19
8.	PHOTOMETER MODE	20
	8.1. METHOD SELECTION	20
	8.2. COLLECTING & MEASURING SAMPLES AND REAGENTS	20
	8.3. CUVETTE PREPARATION	21
	8.4. TIMERS & MEASUREMENT FUNCTIONS	23
	8.5. CHEMICAL FORMULA & UNIT CONVERSION	23
	8.6. METER VALIDATION & CAL CHECK	24
	8.7. ABSORBANCE MEASUREMENTS	25
9.	PROBE MODE	26
	9.1. pH MEASUREMENT	26
	9.2. pH CALIBRATION	27

	9.3. pH MESSAGES & WARNINGS	28
	9.4. pH GLP	29
	9.5. pH ELECTRODE CONDITIONING & MAINTENANCE	30
10.	METHOD PROCEDURES	32
	10.1. ALUMINUM	32
	10.2. AMMONIA LOW RANGE	36
	10.3. AMMONIA MEDIUM RANGE	39
	10.4. AMMONIA HIGH RANGE	42
	10.5. BROMINE	45
	10.6. CHLORINE DIOXIDE	47
	10.7. CHLORINE DIOXIDE, RAPID METHOD	51
	10.8. CHLORINE, FREE	55
	10.9. CHLORINE, TOTAL	58
	10.10. CHROMIUM(VI) LOW RANGE	62
	10.11. CHROMIUM(VI) HIGH RANGE	64
	10.12. COPPER LOW RANGE	66
	10.13. COPPER HIGH RANGE	68
	10.14. HYDRAZINE	70
	10.15. IRON LOW RANGE	72
	10.16. IRON HIGH RANGE	75
	10.17. IRON(II)	77
	10.18. MOLYBDENUM	80
	10.19. NITRATE	83
	10.20. NITRITE LOW RANGE	86
	10.21. NITRITE HIGH RANGE	88
	10.22. OXYGEN, DISSOLVED	90
	10.23. OXYGEN SCAVENGERS (CARBOHYDRAZIDE)	93
	10.24. OXYGEN SCAVENGERS (DIETHYLHYDROXYLAMINE) (DEHA)	96
	10.25. OXYGEN SCAVENGERS (HYDROQUINONE)	99
	10.26. OXYGEN SCAVENGERS (ISO-ASCORBIC ACID)	02
	10.27. pH1	05
	10.28. PHOSPHATE LOW RANGE 1	07

TABLE OF CONTENTS

10.29. PHOSPHATE HIGH RANGE	109
10.30. SILICA LOW RANGE	112
10.31. SILICA HIGH RANGE	115
10.32. ZINC	118
11. WARNINGS & ERRORS	120
12. STANDARD METHODS	123
13. ACCESSORIES	125
13.1. REAGENT SETS	125
13.2. pH ELECTRODES	128
13.3. pH SOLUTIONS	129
13.4. OTHER ACCESSORIES	131
CERTIFICATION	133
RECOMMENDATIONS FOR USERS	133
WARRANTY	134

1. PRELIMINARY EXAMINATION

Remove the instrument and accessories from the packaging and examine it carefully. For further assistance, please contact your local Hanna Instruments Office or email us at tech@hannainst.com. Each H183305 is delivered in a rugged carrying case and is supplied with:

- Sample cuvette (4 pcs.)
- Sample cuvette cap (4 pcs.)
- Cloth for wiping cuvettes
- Scissors
- USB cable
- 5 Vdc power adapter
- 60 mL glass bottle
- Instrument quality certificate
- Instruction manual

Note: Save all packing material until you are sure that the instrument works correctly. Any damaged or defective item must be returned in its original packing material with the supplied accessories.

2. SAFETY MEASURES

- The chemicals contained in the reagent kits may be hazardous if improperly handled.
- Read the Safety Data Sheets (SDS) before performing tests.
 - Safety equipment: Wear suitable eye protection and clothing when required and follow instructions carefully.
 - Reagent spills: If a reagent spill occurs, wipe up immediately and rinse with plenty of water. If reagent contacts skin, rinse the affected area thoroughly with water. Avoid breathing released vapors.
 - Waste disposal: For proper disposal of reagent kits and reacted samples, contact a licensed waste disposal provider.

SPECIFICATIONS

3. SPECIFICATIONS

Measurement Channels		5 x optical channels 1 x digital electrode channel (pH measurement)	
	Range	0.000 to 4.000 Abs	
	Resolution	0.001 Abs	
	Accuracy	±0.003 Abs @ 1.000 Abs	
	Light source	Light Emitting Diode	
Photometer	Bandpass filter bandwidth	8 nm	
	Bandpass filter wavelength accuracy	±1.0 nm	
	Light detector	Silicon photocell	
	Cuvette types	Round, 24.6 mm diameter	
	Number of methods	37	
	Range	-2.00 to 16.00 pH (\pm 1000.0 mV)*	
	Resolution	0.01 pH (0.1 mV)	
	Accuracy	±0.01 pH (±0.2 mV) @ 25 °C / 77 °F	
Probe	Temperature compensation	ATC, -5.0 to 100.0 °C (23.0 to 212.0 °F)*	
	Calibration	two-point, from five available buffers (4.01, 6.86, 7.01, 9.18, 10.01 pH)	
	Electrode	Intelligent pH / temperature electrode	
	Range	-20.0 to 120.0 °C (-4.0 to 248.0 °F)	
Temperature	Resolution	0.1 °C (0.1 °F)	
	Accuracy	\pm 0.5 °C @ 25 °C (\pm 0.9 °F @ 77 °F)	
	Logging	1000 readings (mixed photometer and electrode)	
	Display	128 x 64 pixel B/W LCD with backlight	
	USB-A (Host) functions	Mass-storage host	
	USB-B (Device) functions	Power input, mass-storage device	
	Battery life	> 500 photometer measurements or 50 hours of continuous pH measurement	
Additional Specifications	Power supply	5 Vdc USB 2.0 power adapter / type micro-B connector 3.7 Vdc Li-polymer rechargeable battery, non-serviceable	
	Environment	0 to 50 °C (32 to 122 °F) 0 to 95% RH, non-serviceable	
	Dimensions	206 x 177 x 97 mm (8.1 x 7.0 x 3.8")	
	Weight	1.0 kg (2.2 lbs.)	

*Limits will be reduced to actual probe / sensor limits.

4. ABBREVIATIONS

Abs	Absorbance
GLP	Good Laboratory Practice
NIST	National Institute of Standards and Technology
EPA	US Environmental Protection Agency
g/L	grams per liter (parts per thousand, ppt)
μ g/L	micrograms per liter (parts per billion, ppb)
mg/L	milligrams per liter (parts per million, ppm)
HR	High Range
LR	Low Range
MR	Medium Range

5. DESCRIPTION

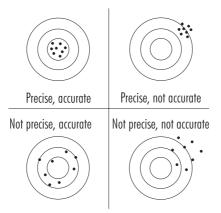
5.1. GENERAL DESCRIPTION & INTENDED USE

HI83305 multiparameter photometer is a compact and versatile meter with two measurement modes, Photometer and Probe. Photometer mode includes a CAL Check[™] feature and 37 different methods that cover a wide variety of applications, making it ideal for both benchtop and portable operations. With the CAL Check[™] feature users are able to validate the performance of the instrument and apply a user calibration (if necessary). Hanna Instruments CAL Check[™] cuvettes are made with NIST traceable standards. Probe mode uses a digital pH probe with a one or two-point calibration.

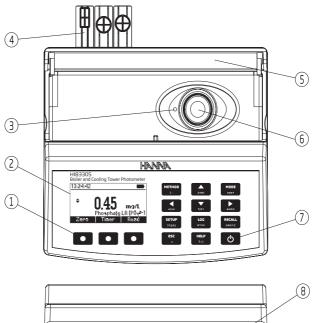
- Digital electrode input for pH measurements
- · Certified CAL Check cuvettes to confirm meter functionality
- Dual purpose micro-USB flash drive
- Lithium polymer rechargeable battery
- Auto-off
- Absorbance mode
- User and sample name entry
- GLP features

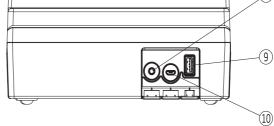
5.2. PRECISION & ACCURACY

Precision is how closely repeated measurements are to one another. Precision is usually expressed as standard deviation. Accuracy is defined as the closeness of a test result to the true value. Although good precision suggests good accuracy, precise results can be inaccurate. The figure explains these definitions. For each method, the accuracy is expressed in the related measurement section.



5.3. FUNCTIONAL DESCRIPTION





- 1. Splash-proof keypad
- 2. Liquid Crystal Display (LCD)
- 3. Indexing mark
- 4. Protective port covers
- 5. Light-blocking cover panel
- 6. Cuvette holder
- 7. ON/OFF power button
- 8. 3.5 mm TRRS (jack) input for digital electrodes
- 9. Standard USB host connector for data transfer to a USB flash drive
- 10. Micro-USB device connector for power or PC interface

Keypad Description

The keypad contains 12 direct keys and 3 functional keys with the following functions:



Press the functional key to perform the function displayed above it on the LCD.



Press to access the list of photometer methods.



Press to move up in a menu or a help screen, to increment a set value or to access second level functions.



Press to toggle between photometer and probe (pH electrode) mode.



Press to move left in a menu or to decrement a set value.



Press to move down in a menu or a help screen, to decrement a set value or to access second level functions.



Press to move right in a menu or to increment a set value.



Press to access the setup screen.



Press to log the current reading.



Press to review saved logs.



Press to exit the current screen.

Press to display the help screen.

ON/OFF power button

5.4. PRINCIPLE OF OPERATION

Absorption of light is a typical phenomenon of interaction between electromagnetic radiation and matter. When a light beam crosses a substance, some of the radiation may be absorbed by atoms, molecules or crystal lattices. Photometric chemical analysis is based on specific chemical reactions between a sample and reagent to produce a light-absorbing compound.

If pure absorption occurs, the fraction of light absorbed depends both on the optical path length through the matter and on the physical-chemical characteristics of the substance according to the Lambert-Beer Law. If all other factors are constant, the concentration "c" can be calculated from the absorbance of the substance.

DESCRIPTION

Lambert Beer Law:

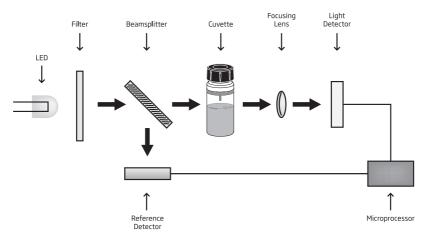
-log I/I
$$_{\rm o} = \epsilon_{\lambda}$$
 c d
or
A = ϵ_{λ} c d

 $I_o =$ intensity of incident light beam

- I = intensity of light beam after absorption
- $\epsilon_{\lambda} = molar$ extinction coefficient at wavelength λ
- c = molar concentration of the substance

d = optical path through the substance

5.5. OPTICAL SYSTEM



Instrument Block Diagram

The internal reference system (reference detector) of the HI83305 photometer compensates for any drifts due to power fluctuations or ambient temperature changes, providing a stable source of light for your blank (zero) measurement and sample measurement.

LED light sources offer superior performance compared to tungsten lamps. LEDs have a much higher luminous efficiency, providing more light while using less power. They also produce little heat, which could otherwise affect electronic stability. LEDs are available in a wide array of wavelengths, whereas tungsten lamps have poor blue / violet light output.

Improved optical filters ensure greater wavelength accuracy and allow a brighter, stronger signal to be received. The end result is higher measurement stability and less wavelength error.

A focusing lens collects all of the light that exits the cuvette, eliminating errors from cuvette imperfections and scratches, eliminating the need to index the cuvette.

6. GENERAL OPERATIONS

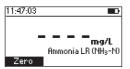
6.1. POWER CONNECTION & BATTERY MANAGEMENT

The meter can be powered from an AC / DC adapter (included) or from the built-in rechargeable battery.

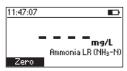
The meter will perform an auto-diagnostic test when it is first powered on. During this test, the Hanna Instruments logo will appear on the LCD. After 5 seconds, if the test was successful, the last method used will appear on the display.

The battery icon on the LCD will indicate the battery status:

• battery is charging from external adapter • battery fully charged (meter connected to AC / DC adapter)

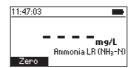


• battery capacity (no external adapter)

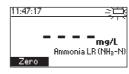


• battery exhausted (no external adapter)





• battery near 0% (no external adapter)



To conserve battery, the meter will turn off automatically after 15 minutes of inactivity (30 minutes after a Zero measurement). If a photometer measurement is on the screen, an auto-log is created before shutdown.

6.2. MODE SELECTION

The HI83305 has two operational modes: Photometer and Probe.

Photometer mode enables on-demand measurement of a cuvette using the integrated optical system.

Probe mode enables continuous measurement using a Hanna digital electrode connected to the 3.5 mm port.

To switch between Photometer mode and Probe mode, use the MODE key.

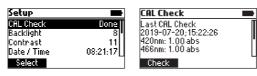
Note: The active mode cannot be switched while in Setup, Recall or Method menus.

6.3. GENERAL SETUP

Press the SETUP key to enter in Setup menu, highlight desired option using the $\blacktriangle \nabla$ keys and press Select.

CAL Check (Photometer Mode Only)

Press **Select** to enter the CAL Check screen. The date, time and values for the last CAL Check are displayed on the screen. To start a new CAL Check press **Check** and follow the prompts on the screen. See METER VALIDATION & CAL CHECK for additional information.



Temperature Unit (Probe Mode Only)

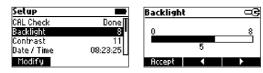
Option: °C or °F Press the functional key to select the desired temperature unit.

Setup	-
Temperature Unit	°C
Backlight	5 •
Contrast	11
Date / Time	15:01:33
°F	

Backlight

Values: 0 to 8

Press **Modify** to access the backlight intensity. Use the functional keys or the **I** keys to increase or decrease the value. Press **Accept** to confirm or press the **ESC** key to return to the **Setup** menu without saving the new value.



Contrast

Values: 0 to 20

Press **Modify** to change the display's contrast. Use the functional keys or the **I** keys to increase or decrease the value. Press **Accept** to confirm the value or the **ESC** key to return to the **Setup** menu without saving the new value.

Setup		Contrast	œ
CAL Check	Done		
Backlight	8 4	0	20
Contrast	11		
Date / Time	08:23:52	6	
Modify		Accept 🚽	

Date & Time

Press **Modify** to change the date and time. Press the functional keys or the **keys** to highlight the value to be modified (year, month, day, hour, minute or second). Use the **keys** to change the value. Press **Accept** to confirm or **ESC** key to return to the **Setup** without saving the new date or time.



Time Format

Option: AM/PM or 24-hour

Press the functional key to select the desired time format.

Setup	
Backlight	5
Contrast	11
Date / Time	13:35:59
Time Format	24-hour
AM/PM	

Date Format

Option: DD/MM/YYYY, MM/DD/YYYY, YYYY/MM/DD, YYYY-MM-DD, Mon DD, YYYY, DD-Mon-YYYY, YYYY-Mon-DD

Press **Modify** to change the date format. Use the **AV** keys to select the desired format. Press **Select** to confirm or the **ESC** key to return to the **Setup** menu without saving the new format.

Setup		Date Format	
Contrast	11	YYYY-MM-DD	Γ
Date / Time	13:36:10	Mon DD, YYYY	-
Time Format	24 hour	DD-Mon-YYYY	
Date Format	Mon DD, YYYY	YYYY-Mon-DD	
Modify		Select	

Decimal Separator

Option: Comma (,) or Period (.)

Press the functional key to select the desired decimal separator. The decimal separator is used on the measurement screen and CSV (Comma-Separated Values) files.

Setup	
Date / Time	13:36:27
Time Format	24 hour
Date Format	Mon DD, YYYY
Decimal Separ	ator •
,	

Language

Option: Português, Deutsch, English, Español, Français, Italiano, Dutch

Press **Modify** to change the language. Use the **A** keys to select the desired language. Press **Select** to change the language.

Setup		Language	⊂3
Decimal Separator	• []	English	
Language	English	Español	
Beeper		Français	
Instrument ID	000000	Italiano	
Modify		Select	

Beeper

Option: Enable or Disable

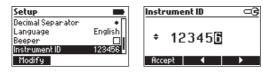
When enabled, a short beep is heard every time a key is pressed. A long beep alert sounds when the pressed key is not active or an error is detected. Press the functional key to enable or disable the beeper.

Setup	
Date Format	Mon DD, YYYY 🗍
Decimal Separ	ator •
Language	English
Beeper	
Enable	

Instrument ID

Option: 0 to 999999

This option is used to set the instrument's ID (identification number). Press **Modify** to access the instrument ID screen. Use the functional keys or the $\triangleleft \triangleright$ keys to highlight the digit to be modified. Press the $\triangleleft \lor$ keys in order to set the desired value. Press **Accept** to confirm the value or press the **ESC** key to return to the **Setup** menu without saving the new value.



Meter Information

Press **Select** to view the model, serial number, firmware version and selected language. Press the **ESC** key to return to the **Setup** menu.

etup		Meter Info	rmation
.anguage	English	Model	HI833
Beeper		Serial #	AAA00000
Instrument ID	000000	Firmware	1.
Meter Information		Language	Engli
Select		www.h	annainst.com

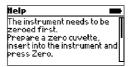
Probe Information (pH Mode Only)

Press **Select** to view model number, serial number and firmware version for the connected probe. Press the **ESC** key to return to the **Setup** menu.



6.4. CONTEXTUAL HELP

HI83305 offers an interactive contextual help mode that assists the user at any time. To access the help screen press the HELP key. The instrument will display additional information related to the current screen. To read all the available information, scroll the text using the **A**V keys. Press the **ESC** key to return to the previous screen.



7. LOGGING DATA & DATA MANAGEMENT

The instrument features a data log function to help you keep track of all your analysis. The data log can hold 1000 individual measurements. Storing, viewing and deleting the data is possible using the LOG and RECALL keys.

7.1. LOGGING DATA

Press the LOG key and the last valid measurement will be stored with a date and time stamp. Only valid measurements can be stored.



7.2. ADDING SAMPLE & USER NAMES TO LOG DATA

A sample ID and user ID can be added to the saved log. Use the **A v** keys to highlight the Sample ID or User ID then press **Modify**. Sample ID and user ID are entered using the alphanumeric multi-tapping keypad.

Log save	2/1000 🗅
2.44 mg/L	(NH ₃ -N)
May 10,2019	11:26:05 AM
Sample ID	
User ID	
Modify	Log

Enter one character at a time by pressing the key with the assigned character repeatedly until the desired character is highlighted. For reference, a list of the characters available for the current key will be shown under the text box.

The character will be entered after a two-second delay or after another key is pressed.

Sample ID	Sample ID
Sam	Sam
MN0 mno 6	
Accept 4 Clear	Accept 🛛 🔹 Clear

Press Accept to update the sample or user ID.

Press ◀ functional key to delete the last character.

Press Clear to delete all of the characters.

Press the ESC key to discard all changes and return to the previous screen.

7.3. DATA MANAGEMENT

Viewing & Deleting

Data can be viewed, deleted and exported to a USB drive or a PC by pressing the **RECALL** key. Use the \blacktriangle keys to scroll through the saved logs. Press **Info** to view additional information about the selected log.

Log Rea	all	1/5 🖿	Log Info	1/5 🗖
30/08 30/08 30/08 30/08	1.40 mg/L 2.00 mg/L 7.6 pH 8.7 mg/L	.NH₃-N	1.40 mg/L Cl0 Chlorine Dio> 30/08/2019 Sample ID:	ade
Info S	Export	Delete	Previous	

Use Delete to erase logged data. After pressing Delete the prompt on the display will confirm the action.

Delete Meter Log	Delete All Meter Logs
Do you want to delete the selected log?	Do you want to delete all logs?
Yes No Del All	Yes No

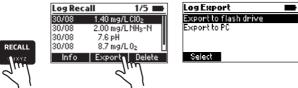
Press No or the ESC key to return to the previous screen.

Press Yes to delete the selected log.

Press **Del All** to erase all the logged data. If **Del All** is pressed the prompt on the display will confirm the action. Press **Yes** to delete all logged data, **No** or the **ESC** key to return to the log recall.

Data Export

Log data can be exported to a USB flash drive or to a PC. To access data export functions, press the **RECALL** key then **Export**.



Use the \blacktriangle keys to select the desired export location.

For export to flash drive, insert the USB flash drive into the dedicated port at the back of the meter labeled HOST USB, then follow the on-screen prompts.

For export to PC, connect the meter to a PC using the supplied micro-USB cable. Insert the cable into the port at the back of the meter labeled PC PWR. Follow the on-screen prompts. When the meter says PC connected, the meter will appear as a removable disk. Use a file manager (such as Windows Explorer or Mac Finder) to move the file from the meter to the PC.

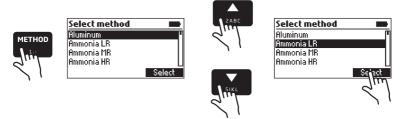
Log data is exported as a single file (H183305.csv) containing all logged photometer and probe data. The CSV file may be opened with a text editor or spreadsheet application.

8. PHOTOMETER MODE

8.1. METHOD SELECTION

In order to select the desired method press the **METHOD** key and a screen with the available methods will appear.

Press the \blacktriangle keys to highlight the desired method. Press Select.

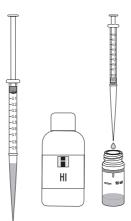


After the desired method is selected, follow the procedure described in the related section. Before performing a method, read all the instructions carefully.

8.2. COLLECTING & MEASURING SAMPLES AND REAGENTS

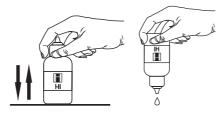
Proper Use of Syringe

- 1. Push the plunger completely into the syringe and insert the tip into the solution.
- 2. Pull the plunger up until the lower edge of the seal is exactly on the mark for the desired volume.
- 3. Take out the syringe and clean the outside of the syringe tip, be sure that no drops are hanging on the tip of the syringe. Then, keeping the syringe in a vertical position, push the plunger down into the syringe, the desired volume has been delivered.



Proper Use of Dropper Bottle

- 1. Tap the dropper on the table several times.
- 2. Remove the cap and wipe the outside of the tip with a cloth.
- 3. Keep the dropper bottle in a vertical position while dosing the reagent.



Proper Use of Powder Packet

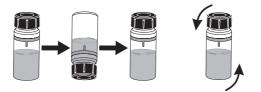
- 1. Use scissors to open the powder packet.
- 2. Push the edges of the packet to form a spout.
- 3. Pour out the content of the packet.



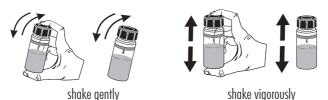
8.3. CUVETTE PREPARATION

Proper mixing is very important for reproducibility of the measurements. The proper mixing technique for each method is listed in the method procedure.

(a) Invert the cuvette a couple of times or for a specified time: hold the cuvette in the vertical position. Turn the cuvette upside-down and wait for all of the solution to flow to the cap end, then return the cuvette to the upright vertical position and wait for all of the solution to flow to the cuvette bottom. This is one inversion. The correct speed for this mixing technique is 10 to 15 complete inversions in 30 seconds. This mixing technique is indicated with "invert to mix" and one of the following icons:



(b) Shaking the cuvette, moving the cuvette up and down. The movement may be gentle or vigorous. This mixing technique is indicated with "shake gently" or "shake vigorously", and one of the following icons:



(c) Swirl the cuvette gently to mix the solution. This mixing technique is indicated with one of the following icons:



In order to avoid reagent leaking and to obtain more accurate measurements, close the cuvette first with the supplied High-Density Polyethylene (HDPE) plastic stopper _____ and then the black cap.

Whenever the cuvette is placed into the measurement holder, it must be dry outside and free of fingerprints, oil and dirt. Wipe it thoroughly with HI731318 microfiber cleaning cloth or a lint-free wipe prior to insertion. Shaking the cuvette can generate bubbles in the sample, causing higher readings. To obtain accurate measurements, remove such bubbles by swirling or by gently tapping the cuvette.

Do not let the reacted sample stand too long after reagent is added. For best accuracy, respect the timings described in each specific method.

It is possible to take multiple readings in a row, but it is recommended to take a new zero reading for each sample and to use the same cuvette for zeroing and measurement when possible.

Discard the sample immediately after the reading is taken, or the glass might become permanently stained.

All the reaction times reported in this manual are at 25 °C (77 °F). In general, the reaction time should be increased for temperatures lower than 20 °C (68 °F) and decreased for temperatures higher than 25 °C (77 °F).

Interferences

In the method measurement section the most common interferences that may be present in a typical water sample have been reported. It is possible that a particular application could introduce other compounds that will also interfere.

8.4. TIMERS & MEASUREMENT FUNCTIONS

Each method requires a different preparation procedure, reaction times and sample preparations. If a timer or timers are necessary for proper sample preparation, the **Timer** will be available.

To use a reaction timer, press **Timer**. The default timer will start immediately. To stop and reset the timer, press **Stop**.

If the selected method requires more than one timer, the meter will automatically select each timer in the appropriate order. To bypass the default order, you may press the desired key to activate a different timer (only while the current timer is stopped). Press **Continue** to start the active timer.

For some methods, the timer is only necessary after a Zero measurement has been performed. In this case, the timer key will only be available after the Zero measurement has been performed.

If the method requires a Zero or Read measurement after a timer has expired, the meter will automatically perform the appropriate action. Follow the instructions in the method procedure.

To perform a Zero or Read measurement, insert the prepared cuvette, then press Zero or Read. A Zero measurement must be conducted before a Read measurement.

8.5. CHEMICAL FORMULA & UNIT CONVERSION

Chemical formula and unit conversion factors are pre-programmed into the instrument and are method specific. In order to view the displayed result in the desired chemical formula press the \blacktriangle keys to access the second level function and then press **Chem Frm** to toggle between the available chemical formulas for the selected method.



8.6. METER VALIDATION & CAL CHECK

Warning: Do not validate the meter with standard solutions other than the Hanna Instruments[®] CAL Check Standards. For accurate validation results, please perform tests at room temperature, 18 to 25 °C (64.4 to 77.0 °F).

Validation of the HI83305 involves absorbance measurements of certified Hanna Instruments[®] CAL Check Standards (see ACCESSORIES section). The CAL Check screen guides the user through the measurement of each CAL Check Standard and applies the factory calibration corrections to each measurement. The HI83305 stores the results of the most recent CAL Check measurements which may be viewed on the CAL Check screen. Compare these results with the values printed on the Certificate provided with each Hanna Instruments[®] CAL Check Standards kit. To perform a validation:

1. Press the **SETUP** key.



2. Highlight CAL Check, then press Select.

Setup	
CAL Check	Done
Backlight	8
Contrast	11
Date / Time	08:21:17
elect	
Jun	
≤ 1	
1	

3. Follow the prompts on the screen. The meter will prompt to measure each cuvette provided in the Hanna Instruments[®] CAL Check Standards kit. To exit the process at any time, press **ESC** key.



4. Press the ESC key to return in Setup menu.



8.7. ABSORBANCE MEASUREMENTS

Raw absorbance measurements may be performed on the HI83305 for personal or diagnostic purposes. For example, you may monitor the stability of a reagent blank by occasionally measuring its absorbance versus deionized water.

To measure the raw absorbance of a prepared sample:

1. Press the **METHOD** key.



- 2. Highlight the appropriate Absorbance method (according to the wavelength to be used), then press **Select**. Absorbance methods are located at the bottom of the method list.
- 3. Prepare the sample cuvette according to the method.
- 4. Insert a cuvette filled with deionized water, then press Zero.
- 5. Insert the prepared sample cuvette, then press Read.

Warning: Never use absorbance methods for validation using Hanna Instruments[®] CAL Check cuvettes. The factory calibration corrections for CAL Check cuvettes are applied while in CAL Check mode only!

9. PROBE MODE

9.1. pH MEASUREMENT

The HI83305 can be used to perform direct pH measurements by connecting a Hanna Instruments[®] digital pH electrode with a 3.5 mm TRRS connector. Connect the electrode to the 3.5 mm port marked with EXT PROBE located at the rear of the meter. If the meter is in Photometer mode, set the meter to Probe mode by pressing the **MODE** key.



Press Calibrate to open the calibration window.

Press **GLP** to review the calibration information.

Press Range to switch between pH and mV.

For high accuracy it is recommended to calibrate your electrode often. pH electrodes should be recalibrated at least once per week, but daily calibration is recommended. Always recalibrate after cleaning an electrode, see pH CALIBRATION section for more information.

To take pH measurements:

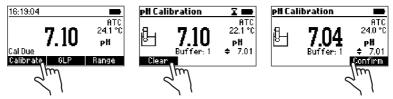
- Remove the protective cap and rinse the electrode with water.
- Collect some sample in a clean, dry beaker.
- Preferably, rinse the electrode with a small amount of sample.
- Submerse the electrode tip approximately 3 cm (11/4") into the sample to be tested and stir the sample gently. Make sure the electrode junction is completely submersed.
- \bullet Allow time for the electrode to stabilize in the sample. When the \blacksquare symbol disappears, your reading is stable.

If measurements are taken successively in different samples, it is recommended to rinse the electrodes thoroughly with deionized or distilled water and then with some of the next sample to prevent cross-contamination.

pH measurements are affected by temperature. Hanna Instruments[®] digital pH electrodes include a built-in temperature sensor and automatically calculate corrected pH values. The measured temperature is displayed on the screen with the pH measurements.

9.2. pH CALIBRATION

From the probe measurement screen, press **Calibrate** to begin the calibration process. During pH calibration, the display will show the current pH reading, temperature reading, selected buffer type and the buffer number ("Buffer: 1" for the first buffer, "Buffer: 2" for the second buffer).



Press Clear to clear the current calibration.

Press **Confirm** to accept the current calibration point (only available if the reading is stable and within the limits for the selected buffer).

Press the **A** keys to cycle through the list of available buffers: pH 4.01, 6.86, 7.01, 9.18, 10.01. Press the **ESC** key to exit calibration and return to pH measurement mode.

Preparation

Pour small quantities of the buffer solutions into clean beakers. If possible, use plastic beakers to minimize any EMC interferences. For accurate calibrations and to minimize cross-contamination, use two beakers for each buffer solution: one for rinsing the electrode and one for calibration. If you are measuring in the acidic range, use pH 7.01 or 6.86 as the first buffer and pH 4.01 as the second buffer. If you are measuring in the alkaline range, use pH 7.01 or 6.86 as the first buffer and pH 10.01 or 9.18 as the second buffer.

Procedure

Calibration can be performed using one or two calibration buffers. For more accurate measurements, a two-point calibration is recommended.

Submerse the pH electrode approximately 3 cm $(1\frac{1}{4})$ into a buffer solution and stir gently.

When the reading is stable and close to the selected buffer, press **Confirm** to accept and store the calibration point. The meter will prompt for the second buffer (Buffer: 2). To use only a one-point calibration, press the **ESC** key to exit calibration mode. The meter will store the calibration information to the probe and return to measurement mode. To continue calibrating with a second buffer, rinse and submerse the pH electrode approximately 3 cm $(1\frac{1}{4})$ into the second buffer solution and stir gently. If necessary, use the \mathbf{A} keys to select a different buffer value.

When the reading is stable and close to the selected buffer, press **Confirm** to accept and store the second calibration point.

The meter will store the two-point calibration information to the probe and return to Measurement mode. The list of calibrated buffers will appear at the bottom of the screen.

9.3. pH MESSAGES & WARNINGS

No Probe

No probe is connected or the probe is broken.

Connecting

The meter has detected a probe and is reading the probe configuration and calibration information.

Incompatible Probe

The connected probe is not compatible with this device.

Incompatible Calibration

The probe's current calibration is not compatible with this meter. The calibration must be cleared to use this probe.

Exceeded Probe Range

The pH and / or temperature measurement exceed the specifications of the probe. The measurement value(s) will be blinking.

Broken Temperature Sensor

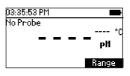
The temperature sensor inside the probe is broken. Temperature compensation will revert to a fixed value of 25 °C (77 °C).

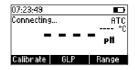
Cal Due

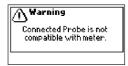
The probe has no calibration. See pH CALIBRATION section for details.

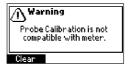
Clean Probe

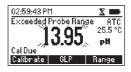
The offset is outside the accepted window or the slope is under the accepted lower limit. Cleaning the probe will improve the pH electrode's response, repeat the calibration after cleaning. See pH ELECTRODE CONDITIONING & MAINTENANCE section for details.

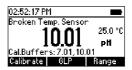




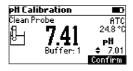












Check Probe & Buffer

There is a large difference between the pH measurement and the selected buffer value or the electrode slope is outside of the accepted slope limit. Clean the probe and confirm the correct buffer selection.

Wrong Temperature

The buffer temperature is outside of the acceptable window for the selected buffer value.



pH Calibration	
Wrong Temperature	ATC
18. 701	112.3 °C
변 /.UI	рH
Buffer: 1	
Clear	Confirm

9.4. pH GLP

Good Laboratory Practice (GLP) refers to a quality control function used to ensure uniformity and consistency of sensor calibrations and measurements. To view the GLP information, press the **GLP** key from the probe measurement screen.

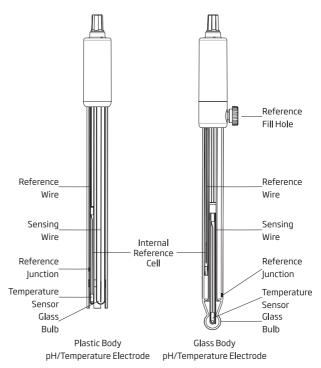


The pH GLP screen displays the date and time, buffers, slope and offset for the last calibration. If the probe has not been calibrated, "No User Calibration" is displayed. Press the **ESC** key to return to the measurement mode.

Last pH Cal
Feb 14,2019 07:27:16
Cal.Buffers: 4.01 , 7.01 Offset: 0.7mV
Slope: 100.1%

Last pH	Cal
Nollsen	Calibration
10050	odilor delori

9.5. pH ELECTRODE CONDITIONING & MAINTENANCE



Remove the protective cap. Do not be alarmed if salt deposits are present, this is normal. Rinse the probe with water.

Shake the electrode down as you would do with a clinical thermometer to eliminate any air bubbles inside the glass bulb. If the bulb and / or junction are dry, soak the electrode in H170300 or H180300 Storage solution for a minimum of 30 minutes. Rinse with water. Calibrate before using.

For refillable electrodes if the filling solution (electrolyte) is more than 2½ cm (1") below the fill hole, add H17082 or H18082 3.5M KCI Electrolyte solution. Unscrew the fill hole cover during measurements so the liquid reference junction maintains an outward flow of electrolyte.

Storage Procedure

To minimize clogging and ensure a quick response time, the glass bulb and the junction should be kept moist and not allowed to dry out.

Replace the solution in the protective cap with a few drops of H170300 or H180300 Storage solution or Filling solution (H17082 or H18082 3.5M KCl Electrolyte solution). pH 4.01 or 7.01 buffer can also be used.

Note: Never store the electrode in distilled or deionized water.

Periodic Maintenance

Inspect the electrode and the cable. The cable used for connection to the instrument must be intact and there must be no points of broken insulation on the cable, connectors must be perfectly clean and dry. If there are any scratches or cracks on the electrode stem or bulb, replace the electrode.

For refillable electrodes, refill the reference chamber with fresh electrolyte (HI7082 or HI8082 3.5M KCI Electrolyte solution). Allow the electrode to stand upright for 1 hour.

Cleaning Procedure

Several cleaning solutions are available:

- General Soak in Hanna H17061 or H18061 General cleaning solution for approximately 30 minutes.
- Protein —Soak in Hanna H17073 or H18073 Protein cleaning solution for 15 minutes.
- Inorganic Soak in Hanna H17074 Inorganic cleaning solution for 15 minutes.
- Oil and grease Rinse with Hanna H17077 or H18077 Oil and Fat cleaning solution.

After performing any of the cleaning procedures, rinse the electrode thoroughly with distilled water, refill the reference chamber with fresh electrolyte (refillable electrodes only) and soak the electrode in H170300 or H180300 Storage solution for at least 1 hour before taking measurements.

Temperature Correlation for pH Sensitive Glass

Verify the temperature range by reading the limits on electrode's cap. The pH electrode's life is temperature dependent. If constantly cycled between two temperatures, the life of the electrode is drastically reduced.

10. METHOD PROCEDURES

10.1. ALUMINUM

SPECIFICATIONS

Range	0.00 to 1.00 mg/L (as Al ³⁺)
Resolution	0.01 mg/L
Accuracy	\pm 0.04 mg/L \pm 4% of reading at 25 °C
Light Source	LED with narrow band interference filter \textcircled{O} 525 nm
Method	Adaptation of the Aluminon Method

REQUIRED REAGENTS

Code	Description	Quantity
HI93712A-0	Aluminum Reagent A	1 packet
HI93712B-0	Aluminum Reagent B	1 packet
HI93712C-0	Aluminum Reagent C	1 packet

REAGENT SETS

HI93712-01	Reagents for 100 tests
HI93712-03	Reagents for 300 tests
For other accessories	see ACCESSORIES section.

MEASUREMENT PROCEDURE

- Select the Aluminum method using the procedure described in the METHOD SELECTION section.
- Fill a graduated beaker with 50 mL of sample.
- Add one packet of H193712A-0 Aluminum Reagent A and mix until completely dissolved.
- Add one packet of HI93712B-0 Aluminum Reagent B and mix until completely dissolved.



- Fill two cuvettes with 10 mL of sample (up to the mark).
- Add one packet of HI93712C-0 Aluminum Reagent C to one cuvette (#1). Replace the plastic stopper and the cap. Shake gently until completely dissolved. This is the blank.
- Insert the first cuvette (#1) into the holder and close the lid.

mg/L Aluminum (AI3+)

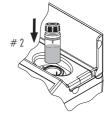
Bead

16:42:12

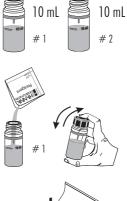
• Press Timer and the display will show the countdown prior to the zero or wait 15 minutes and then press Zero. The display will show "-0.0-" when the meter is zeroed and ready for measurement.

- Remove the blank and insert the second cuvette (#2) into the holder and close the lid.
- 10:02:38 AM mg/L inum (Al≥

15min





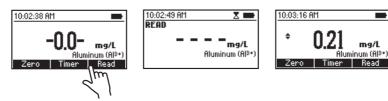




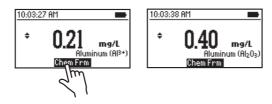
mg/L

Aluminum (Al³+

• Press Read to start the reading. The instrument displays the results in mg/L of aluminum (Al³⁺).



- Press the \blacktriangle or \blacktriangledown key to access the second level functions.
- Press Chem Frm to convert the result to mg/L of aluminum oxide (Al₂O₃).

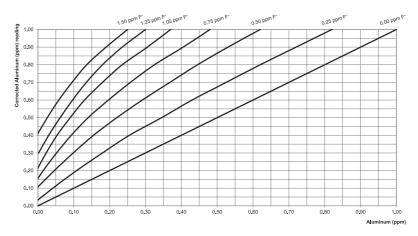


• Press the \blacktriangle or \blacktriangledown key to return to the measurement screen.

INTERFERENCES

Interference may be caused by:

- Alkalinity above 1000 mg/L
- Phosphate above 50 mg/L
- Iron above 20 mg/L
- Fluoride must be absent. If the fluoride concentration is known, the aluminum concentration can be determined using the graph below:



To use the fluoride interference graph:

- 1. Follow the measurement procedure to obtain the aluminum concentration.
- 2. Locate the aluminum reading on x-axis.
- 3. Follow the line up, until it intersects the fluoride curve corresponding to the fluoride concentration in the sample.
- 4. From the intersection of the fluoride and aluminum line, follow the line to the left until it intersects the y-axis. This point corresponds to the corrected aluminum concentration in the sample.

E.g. Aluminum reading on meter 0.40 ppm and fluoride content in sample 0.50 ppm, corrected aluminum concentration in sample is 0.75 ppm.

10.2. AMMONIA LOW RANGE

SPECIFICATIONS

Range	0.00 to 3.00 mg/L (as NH ₃ -N)
Resolution	0.01 mg/L
Accuracy	\pm 0.04 mg/L \pm 4% of reading at 25 °C
Light Source	LED with narrow band interference filter @ 420 nm
Method	Adaptation of the ASTM Manual of Water and Environmental Technology,
	D1426 Nessler Method

REQUIRED REAGENTS

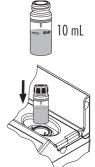
Code	Description	Quantity
HI93700A-0	Ammonia Low Range Reagent A	4 drops
HI93700B-0	Ammonia Low Range Reagent B	4 drops

REAGENT SETS

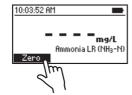
HI93700-01	Reagents for 100 tests
HI93700-03	Reagents for 300 tests
For other accessories	see ACCESSORIES section.

MEASUREMENT PROCEDURE

- Select the Ammonia LR method using the procedure described in the METHOD SELECTION section.
- Fill the cuvette with 10 mL of unreacted sample (up to the mark). Replace the plastic stopper and the cap.



- Insert the cuvette into the holder and close the lid.
- Press Zero. The display will show "-0.0-" when the meter is zeroed and ready for measurement.



10:04:21 AM	X 🖛
ZERO	
	mg/L
A	mmonia LR (NH₃-N)



- Remove the cuvette.
- Add 4 drops of H193700A-0 Ammonia Low Range Reagent A. Replace the plastic stopper and the cap. Swirl to mix the solution.

- Add 4 drops of H193700B-0 Ammonia Low Range Reagent B. Replace the plastic stopper and the cap. Swirl to mix the solution.
- Insert the cuvette into the holder and close the lid.

ma/l

LB (NH»-N)

Bead

B (NH₅-N

10:04:30 AM

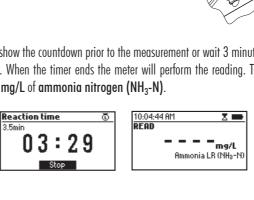
Zero

10:05:16 AM 4

• Press Timer and the display will show the countdown prior to the measurement or wait 3 minutes and 30 seconds and press Read. When the timer ends the meter will perform the reading. The instrument displays the results in mg/L of ammonia nitrogen (NH₃-N).







• Press Chem Frm to convert the result to mg/L of ammonia (NH₃) and ammonium (NH₄⁺).



- Press the \blacktriangle or \blacktriangledown key to return to the measurement screen.

INTERFERENCES

- Hardness above 1 g/L
- Iron
- Sulfide may cause turbidity
- Organic compounds like acetone above 0.1%, alcohols, aldehydes, aliphatic and aromatic amines, chloramines, glycine, or urea above 10 mg/L, to remove the interference distillation is required

10.3. AMMONIA MEDIUM RANGE

SPECIFICATIONS

Range	0.00 to 10.00 mg/L (as NH ₃ -N)
Resolution	0.01 mg/L
Accuracy	\pm 0.05 mg/L \pm 5% of reading at 25 °C
Light Source	LED with narrow band interference filter @ 420 nm
Method	Adaptation of the ASTM Manual of Water and Environmental Technology,
	D1426, Nessler Method

REQUIRED REAGENTS

Code	Description	Quantity
HI93715A-0	Ammonia Medium Range Reagent A	4 drops
HI93715B-0	Ammonia Medium Range Reagent B	4 drops

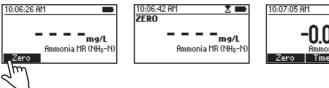
REAGENT SETS

HI93715-01	Reagents for 100 tests
HI93715-03	Reagents for 300 tests
For other accessor	ies see ACCESSORIES section.

MEASUREMENT PROCEDURE

- Select the Ammonia MR method using the procedure described in the METHOD SELECTION section.
- Fill the cuvette with 10 mL of unreacted sample (up to the mark). Replace the plastic stopper and the cap.
- 10 mL ĨT

- Insert the cuvette into the holder and close the lid.
- Press Zero. The display will show "-0.0-" when the meter is zeroed and ready for measurement.





• Remove the cuvette.

10:07:05 AM

Zero

10:07:41 AM

Zero

- Add 4 drops of H193715A-0 Ammonia Medium Range Reagent A. Replace the plastic stopper and the cap. Swirl to mix the solution.
- Add 4 drops of H193715B-0 Ammonia Medium Range Reagent B. Replace the plastic stopper and the cap. Swirl to mix the solution.

• Insert the cuvette into the holder and close the lid.

mg/L

Read

mg/L MR (NH₃-N)

Bead

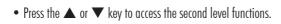
Ammonia MR (NH₂-N)

 Press Timer and the display will show the countdown prior to the measurement or wait 3 minutes and 30 seconds and press Read. When the timer ends the meter will perform the reading. The instrument displays the results in mg/L of ammonia nitrogen (NH₃-N).

03:29

Reaction time

3.5min









Χ.

mg/L

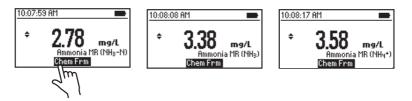
Ammonia MR (NH₂-N)

10:06:53 AM

READ

Ō

- **AMMONIA MEDIUM RANGE**
- Press Chem Frm to convert the result to mg/L of ammonia (NH₃) and ammonium (NH₄⁺).



- Press the \blacktriangle or \blacktriangledown key to return to the measurement screen.

INTERFERENCES

- Hardness above 1 g/L
- Iron
- Sulfide may cause turbidity
- Organic compounds like acetone above 0.1%, alcohols, aldehydes, aliphatic and aromatic amines, chloramines, glycine, or urea above 10 mg/L, to remove the interference distillation is required

10.4. AMMONIA HIGH RANGE

SPECIFICATIONS

Range	0.0 to 100.0 mg/L (as NH ₃ -N)
Resolution	0.1 mg/L
Accuracy	± 0.5 mg/L $\pm 5\%$ of reading at 25 °C
Light Source	LED with narrow band interference filter @ 420 nm
Method	Adaptation of the ASTM Manual of Water and Environmental Technology,
	D1426, Nessler Method

REQUIRED REAGENTS

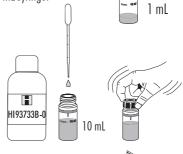
Code	Description	Quantity
HI93733A-0	Ammonia High Range Reagent A	4 drops
HI93733B-0	Ammonia High Range Reagent B	9 mL

REAGENT SETS

HI93733-01	Reagents for 100 tests
HI93733-03	Reagents for 300 tests
For other accessorie	es see ACCESSORIES section.

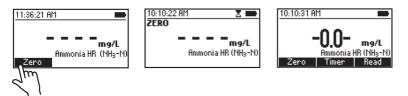
MEASUREMENT PROCEDURE

- Select the Ammonia HR method using the procedure described in the METHOD SELECTION section.
- Add 1 mL of unreacted sample to the cuvette using a 1 mL syringe.
- Use the pipette to fill the cuvette up to the 10 mL mark with HI93733B-0 Ammonia High Range Reagent B. Replace the plastic stopper and the cap. Swirl to mix the solution.





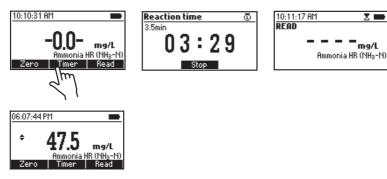
• Press Zero. The display will show "-0.0-" when the meter is zeroed and ready for measurement.



∀~())))

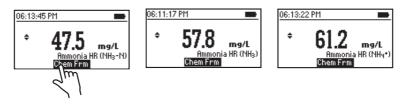
×4

- Remove the cuvette.
- Add 4 drops of H193733A-O Ammonia High Range Reagent A. Replace the plastic stopper and the cap. Swirl to mix the solution.
- Insert the cuvette into the holder and close the lid.
- Press Timer and the display will show the countdown prior to the measurement or wait 3 minutes and 30 seconds and press Read. When the timer ends the meter will perform the reading. The instrument displays the results in mg/L of ammonia nitrogen (NH₃-N).



• Press the \blacktriangle or \blacktriangledown key to access the second level functions.

• Press Chem Frm to convert the result to mg/L of ammonia (NH₃) and ammonium (NH₄⁺).



- Press the \blacktriangle or \blacktriangledown key to return to the measurement screen.

INTERFERENCES

- Hardness above 1 g/L
- Iron
- Sulfide may cause turbidity
- Organic compounds like acetone above 0.1%, alcohols, aldehydes, aliphatic and aromatic amines, chloramines, glycine, or urea above 10 mg/L, to remove the interference distillation is required

10.5. BROMINE

SPECIFICATIONS

Range	0.00 to 8.00 mg/L (as Br ₂)
Resolution	0.01 mg/L
Accuracy	\pm 0.08 mg/L \pm 3% of reading at 25 °C
Light Source	LED with narrow band interference filter @ 525 nm
Method	Adaptation of Standard Methods for the Examination of Water and
	Wastewater, 18 th Edition, DPD Method

REQUIRED REAGENTS

Code	Description	Quantity
HI93716-0	Bromine Reagent	1 packet

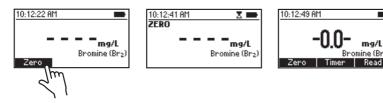
REAGENT SETS

HI93716-01	Reagents for 100 tests
HI93716-03	Reagents for 300 tests
For other accessori	es see ACCESSORIES section.

MEASUREMENT PROCEDURE

- Select the Bromine method using the procedure described in the METHOD SELECTION section.
- Fill the cuvette with 10 mL of unreacted sample (up to the mark). Replace the plastic stopper and the cap.
- 10 mL

- Insert the cuvette into the holder and close the lid.
- Press Zero. The display will show "-0.0-" when the meter is zeroed and ready for measurement.

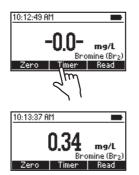


- Remove the cuvette.
- Add one packet of HI93716-0 Bromine Reagent. Replace the plastic stopper and the cap. Shake gently for about 20 seconds to dissolve most of the reagent.
- Insert the cuvette into the holder and close the lid.





 Press Timer and the display will show the countdown prior to the measurement or wait 2 minutes and 30 seconds and press Read. When the timer ends the meter will perform the reading. The instrument displays the results in mg/L of bromine (Br₂).





	Ζ 🖛
_	
_	— _ mg/L
	Bromine (Br ₂)
	-

R

INTERFERENCES

- Chlorine, Iodine, Ozone, Oxidized forms of Chromium and Manganese
- Hardness greater than 500 mg/L CaCO₃, to remove the interference shake the sample for approximately 1 minute after adding the reagent
- Alkalinity greater than 300 mg/L CaCO₃ or acidity greater than 150 mg/L CaCO₃, the color of the sample may develop only partially or rapidly fade, to remove the interference neutralize the sample with diluted HCl or NaOH

10.6. CHLORINE DIOXIDE

SPECIFICATIONS

Range	0.00 to 2.00 mg/L (as ClO_2)
Resolution	0.01 mg/L
Accuracy	\pm 0.10 mg/L \pm 5% of reading at 25 °C
Light Source	LED with narrow band interference filter $@$ 575 nm
Method	Adaptation of the Chlorophenol Red Method

REQUIRED REAGENT

Code	Description	Quantity
HI93738A-0	Chlorine Dioxide Reagent A	1 mL
HI93738B-0	Chlorine Dioxide Reagent B	1 packet
HI93738C-0	Chlorine Dioxide Reagent C	1 mL
HI93738D-0	Chlorine Dioxide Reagent D	1 mL

REAGENT SETS

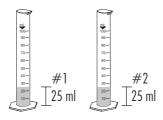
HI93738-01	Reagents for 100 tests
HI93738-03	Reagents for 300 tests
For other accessorie	es see ACCESSORIES section.

SAMPLING PROCEDURE

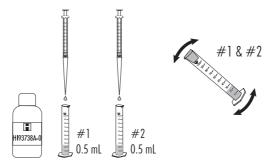
It is recommended to analyze Chlorine Dioxide samples immediately after collection. Chlorine Dioxide samples must be stored in sealed dark glass bottles, with minimal head space. Excessive heat (above 25 °C / 77 °F), agitation and exposure to light must be avoided.

MEASUREMENT PROCEDURE

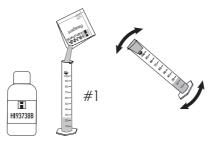
- Select the Chlorine Dioxide method using the procedure described in the METHOD SELECTION section.
- Fill two graduated mixing cylinders (#1 & #2) up to the 25 mL mark with the sample.



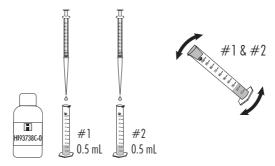
• Add 0.5 mL of H193738A-0 Chlorine Dioxide Reagent A to each cylinder (#1 & #2), using a 1 mL syringe, cap them and invert several times to mix.



• Add one packet of H193738B-O Chlorine Dioxide Reagent B to one of the two cylinders (#1), cap and invert it several times until it is totally dissolved. This is the blank.



• Add 0.5 mL of H193738C-0 Chlorine Dioxide Reagent C to each cylinder (#1 & #2), using a 1 mL syringe, cap them and invert several times to mix.



• Add 0.5 mL of H193738D-0 Chlorine Dioxide Reagent to each cylinder (#1 & #2), using a 1 mL syringe, cap them and invert several times to mix. Cylinder #2 is the reacted sample.

#1

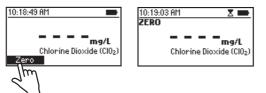
#1

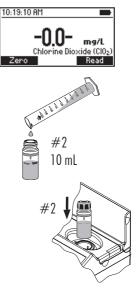
#1

10 mL

Η #1 #2 HI93738D-0 0.5 mL 0.5 mL

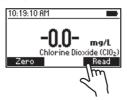
- Fill cuvette (#1) with 10 mL of the blank (up to the mark). Replace the plastic stopper and the cap.
- Insert the blank (#1) into the holder and close the lid.
- Press Zero. The display will show "-0.0-" when the meter is zeroed and ready for measurement.

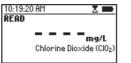




- Fill second cuvette (#2) with 10 mL of the reacted sample (up to the mark). Replace the plastic stopper and the cap.
- Insert the sample into the holder and close the lid.

• Press Read to start the reading. The instrument displays the results in mg/L of chlorine dioxide (CIO₂).







INTERFERENCES Interference may be caused by: • Strong oxidants

10.7. CHLORINE DIOXIDE, RAPID METHOD

SPECIFICATIONS

Range	0.00 to 2.00 mg/L (as ClO ₂)
Resolution	0.01 mg/L
Accuracy	\pm 0.10 mg/L \pm 5% of reading at 25 °C
Light Source	LED with narrow band interference filter $@$ 525 nm
Method	Adaptation of Standard Methods for the Examination of Water and Wastewater, $18^{\rm th}$ Edition, 4500 ClO $_2$ D

REQUIRED REAGENT

Code	Description	Quantity
HI96779A-0	Chlorine Dioxide Reagent A	5 drops
HI96779B-0	Chlorine Dioxide Reagent B	1 packet

REAGENT SETS

HI96779-01	Reagents for 100 tests
HI96779-03	Reagents for 300 tests
For other accessori	es see ACCESSORIES section

PRINCIPLE

The reaction between the Chlorine Dioxide and DPD indicator causes a pink tint in the sample, the addition of glycine as a masking agent inhibits the response of free chlorine.

APPLICATION

Drinking water, tap water, treated water

SAMPLING PROCEDURE

Collect the sample in a clean glass bottle and analyze it immediately. Chlorine dioxide is a strong oxidizing agent and is unstable in water.

SIGNIFICANCE & USE

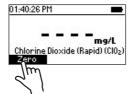
Chlorine Dioxide is a commonly-used alternative to chlorine (Cl_2) as a water disinfectant. The Chlorophenol Red method (non-rapid method) reacts specifically with chlorine dioxide with little interference from free chlorine or chloramines, but the method procedure is cumbersome. The Chlorine Dioxide Rapid Method based on the DPD (N,N-diethyl-p-phenylenediamine) indicator is a much simpler method by comparison, but it is susceptible to interference from other oxidizers. Glycine (Reagent A) is able to convert free chlorine to chloroaminoacetic acid without affecting the analysis of chlorine dioxide content.

MEASUREMENT PROCEDURE

- Select the Chlorine Dioxide (Rapid) method using the procedure described in the METHOD SELECTION section.
- Fill the cuvette with 10 mL of unreacted sample (up to the mark).

• Add 5 drops of HI96779A-0 Chlorine Dioxide Reagent A.

- Replace the plastic stopper and the cap. Shake gently for 30 seconds.
- Wait 30 seconds.
- Insert the cuvette into the holder and close the lid.
- Press Zero. The display will show "-0.0-" when the meter is zeroed and ready for measurement.



01:41:19 PM	Z 🖛
ZERO	1006603 - 670
	mg/L
Chlorine Dioxide	e (Rapid) (ClO ₂)







01:43:17 PM

- Remove the cuvette.
- Add one packet of HI96779B-0 Chlorine Dioxide Reagent B.
- Replace the plastic stopper and the cap. Shake gently for 20 seconds.
- Insert the cuvette into the holder and close the lid.

Chlorine Dioxide (R Zero

Time

• Press Timer and the display will show the countdown prior to the measurement or wait 1 minute and press Read. When the timer ends the meter will perform the reading. The instrument displays the results in mg/L of CIO₂.









INTERFERENCES

- Acidity, Alkalinity, Flocculating agents, Hardness, Inorganic and Organic Chloramines, Manganese, Metals, Monochloramine, Oxidized forms of Chromium and Manganese, Ozone and Peroxides
- Chlorine above 5 mg/L
- Bromine above 0.1 mg/L
- Highly buffered samples or extreme sample pH

10.8. CHLORINE, FREE

SPECIFICATIONS

Range	0.00 to 5.00 mg/L (as Cl_2)
Resolution	0.01 mg/L
Accuracy	\pm 0.03 mg/L \pm 3% of reading at 25 °C
Light Source	LED with narrow band interference filter $@$ 525 nm
Method	Adaptation of the EPA DPD Method 330.5

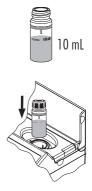
REQUIRED REAGENTS

TOWDER		
Code	Description	Quantity
HI93701-0	Free Chlorine Reagent	1 packet
LIQUID	-	
Code	Description	Quantity
HI93701A-F	Free Chlorine Reagent A	3 drops
HI93701B-F	Free Chlorine Reagent B	3 drops
REAGENT SETS		

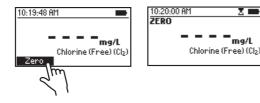
HI93701-F	Reagents for 300 tests (liquid)	
HI93701-01	Reagents for 100 tests (powder)	
HI93701-03	Reagents for 300 tests (powder)	
For other accessories see ACCESSORIES section.		

MEASUREMENT PROCEDURE

- Select the Chlorine (Free) method using the procedure described in the METHOD SELECTION section.
- Fill the cuvette with 10 mL of unreacted sample (up to the mark). Replace the plastic stopper and the cap.
- Insert the cuvette into the holder and close the lid.



• Press Zero. The display will show "-0.0-" when the meter is zeroed and ready for measurement.





• Remove the cuvette.

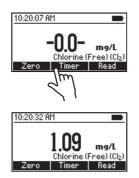
POWDER REAGENT PROCEDURE

- Add the content of one packet of HI93701-0 Free Chlorine Reagent. Replace the plastic stopper and the cap. Shake gently for 20 seconds.
- Insert the cuvette into the holder and close the lid.





• Press **Timer** and the display will show the countdown prior to the measurement or wait 1 minute and press **Read**. When the timer ends the meter will perform the reading. The instrument displays the results in **mg/L** of **chlorine (Cl₂)**.







CHLORINE, FREE

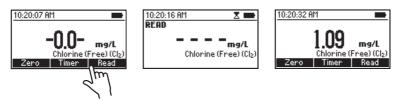
×3

10 ml

×3

LIQUID REAGENT PROCEDURE

- To an empty cuvette add 3 drops of HI93701A-F Free Chlorine Reagent A and 3 drops of HI93701B-F Free Chlorine Reagent B.
- Replace the plastic stopper and the cap. Swirl gently to mix.
- Add 10 mL of unreacted sample (up to the mark). Replace the plastic stopper and the cap. Shake gently.
- Insert the cuvette into the holder and close the lid.
- Press Read to start the reading. The instrument displays the results in mg/L of chlorine (Cl₂).



Note: Free and Total Chlorine have to be measured separately with fresh sample following the related procedure if both values are desired.

INTERFERENCES

- Bromine, Iodine, Oxidized forms of Chromium and Manganese, Ozone
- Hardness greater than 500 mg/L CaCO₃, to remove the interference shake the sample for approximately 2 minutes after adding the powder reagent
- Alkalinity greater than 250 mg/L CaCO₃ or acidity value greater than 150 mg/L CaCO₃, the color of the sample may develop only partially or rapidly fade, to remove the interference neutralize the sample with diluted HCl or NaOH

10.9. CHLORINE, TOTAL

SPECIFICATIONS

Range	0.00 to 5.00 mg/L (as Cl_2)
Resolution	0.01 mg/L
Accuracy	\pm 0.03 mg/L \pm 3% of reading at 25 °C
Light Source	LED with narrow band interference filter $@$ 525 nm
Method	Adaptation of the EPA DPD Method 330.5

REQUIRED REAGENTS

TOWDER		
Code	Description	Quantity
HI93711-0	Total Chlorine Reagent	1 packet
LIQUID		
Code	Description	Quantity
HI93701A-T	Total Chlorine Reagent A	3 drops
HI93701B-T	Total Chlorine Reagent B	3 drops
HI93701C-T	Total Chlorine Reagent C	1 drop
	· · · · · · · · · · · · · · · · · · ·	·

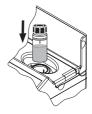
REAGENT SETS

HI93701-T	Reagents for 300 tests (liquid)
HI93711-01	Reagents for 100 total tests (powder)
HI93711-03	Reagents for 300 total tests (powder)
For other accessories s	ee ACCESSORIES section.

MEASUREMENT PROCEDURE

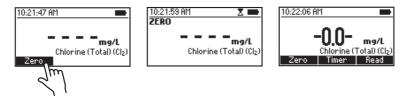
- Select the Chlorine (Total) method using the procedure described in the METHOD SELECTION section.
- Fill the cuvette with 10 mL of unreacted sample (up to the mark). Replace the plastic stopper and the cap.





• Insert the cuvette into the holder and close the lid.

• Press Zero. The display will show "-0.0-" when the meter is zeroed and ready for measurement.

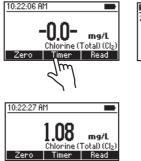


• Remove the cuvette.

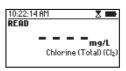
POWDER REAGENT PROCEDURE

 Add 1 packet of HI93711-0 Total Chlorine Reagent. Replace the plastic stopper and the cap. Shake gently for 20 seconds.

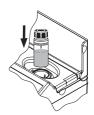
- Insert the cuvette into the holder and close the lid.
- Press Timer and the display will show the countdown prior to the measurement or wait 2 minutes and 30 seconds and press Read. When the timer ends the meter will perform the reading. The instrument displays the results in mg/L of chlorine (Cl₂).





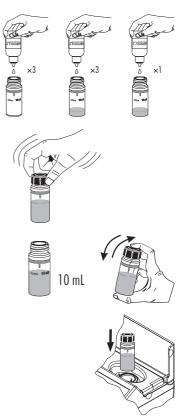




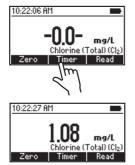


LIQUID REAGENT PROCEDURE

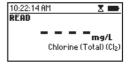
- To an empty cuvette add 3 drops of H193701A-T Total Chlorine Reagent A, 3 drops of H193701B-T Total Chlorine Reagent B and 1 drop of H193701C-T Total Chlorine Reagent C.
- Replace the plastic stopper and the cap. Swirl gently to mix.
- Add 10 mL of unreacted sample (up to the mark). Replace the plastic stopper and the cap. Shake gently.
- Insert the cuvette into the holder and close the lid.



• Press **Timer** and the display will show the countdown prior to the measurement or wait 2 minutes and 30 seconds and press **Read**. When the timer ends the meter will perform the reading. The instrument displays the results in **mg/L** of **chlorine** (Cl₂).







Note: Free and Total Chlorine have to be measured separately with fresh unreacted samples following the related procedure if both values are desired.

INTERFERENCES

- Bromine, Iodine, Oxidized forms of Chromium and Manganese, Ozone
- Hardness greater than 500 mg/L CaCO₃, to remove the interference shake the sample for approximately 2 minutes after adding the powder reagent
- Alkalinity greater than 250 mg/L CaCO₃ or acidity greater than 150 mg/L CaCO₃, the color of the sample may develop only partially or may rapidly fade, to remove the interference neutralize the sample with diluted HCl or NaOH

10.10. CHROMIUM(VI) LOW RANGE

SPECIFICATIONS

Range	0 to 300 µg/L (as Cr (VI))
Resolution	1 µg/L
Accuracy	\pm 10 μ g/L \pm 4% of reading at 25 °C
Light Source	LED with narrow band interference filter @ 525 nm
Method	Adaptation of the ASTM Manual of Water and Environmental Technology,
	D1687 Diphenylcarbohydrazide Method

REQUIRED REAGENTS

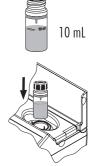
Code	Description	Quantity
HI93749-0	Chromium(VI) Low Range Reagent	1 packet

REAGENT SETS

HI93749-01	Reagents for 100 tests
HI93749-03	Reagents for 300 tests
For other accessori	es see ACCESSORIES section.

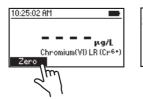
MEASUREMENT PROCEDURE

- Select the Chromium(VI) LR method using the procedure described in the METHOD SELECTION section.
- Fill the cuvette with 10 mL of unreacted sample (up to the mark). Replace the plastic stopper and the cap.
- Insert the cuvette into the holder and close the lid.



• Press Zero. The display will show "-0.0-" when the meter is zeroed and ready for measurement.

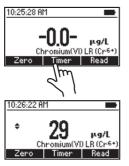
10 OF 00 OM



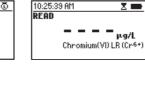
10:20:20) HE				<u> </u>	_
ZERO						
		-	-	-		
					μg	L
	Chr	omi	umC\	/0 LI	R (Ci	r6+)



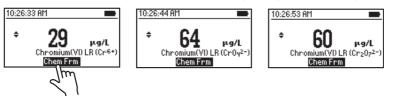
- Remove the cuvette.
- Add one packet of H193749-0 Chromium(VI) Low Range Reagent. Replace the plastic stopper and the cap. Shake vigorously for about 10 seconds.
- Insert the cuvette into the holder and close the lid.
- Press Timer and the display will show the countdown prior to the measurement or wait 6 minutes and press Read. When the timer ends the meter will perform the reading. The instrument displays concentration in µg/L of chromium (Cr⁶⁺).







- Press the \blacktriangle or \blacktriangledown key to access the second level functions.
- Press Chem Frm to convert the result to μ g/L of chromate (Cr0₄²⁻) and dichromate (Cr₂0₇²⁻).



• Press the \blacktriangle or \blacktriangledown key to return to the measurement screen.

INTERFERENCES

- Vanadium above 1 mg/L, wait 10 minutes before reading to remove the interference
- Iron above 1 mg/L
- Mercurous and mercuric ions slight inhibition of the reaction

10.11. CHROMIUM(VI) HIGH RANGE

SPECIFICATIONS

Range	0 to 1000 µg/L (as Cr(VI))
Resolution	1 µg/L
Accuracy	$\pm5\mu$ g/L $\pm4\%$ of reading at 25 °C
Light Source	LED with narrow band interference filter @ 525 nm
Method	Adaptation of the ASTM Manual of Water and Environmental Technology,
	D1687-92, Diphenylcarbohydrazide Method

REQUIRED REAGENTS

Code	Description	Quantity
HI93723-0	Chromium(VI) High Range Reagent	1 packet

REAGENT SETS

HI93723-01	Reagents for 100 tests
HI93723-03	Reagents for 300 tests
For other accessorie	es see ACCESSORIES section.

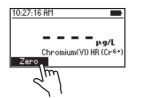
MEASUREMENT PROCEDURE

- Select the Chromium(VI) HR method using the procedure described in the METHOD SELECTION section.
- Fill the cuvette with 10 mL of unreacted sample (up to the mark). Replace the plastic stopper and the cap.





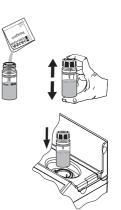
- Insert the cuvette into the holder and close the lid.
- Press Zero. The display will show "-0.0-" when the meter is zeroed and ready for measurement.



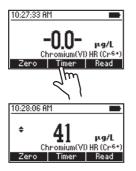
10:27:20 P	111			
ZERO				
-		-	-	
			- μο	
C	hromi	ium(V	1) HR ()	(r6+)



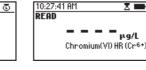
- Remove the cuvette.
- Add one packet of H193723-0 Chromium(VI) High Range Reagent. Replace the plastic stopper and the cap. Shake vigorously for about 10 seconds.
- Insert the cuvette into the holder and close the lid.



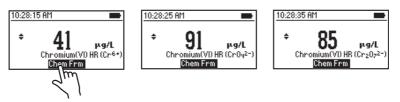
• Press **Timer** and the display will show the countdown prior to the measurement or wait 6 minutes and press **Read**. When the timer ends the meter will perform the reading. The instrument displays concentration in $\mu g/L$ of **chromium** (Cr⁶⁺).







- Press the \blacktriangle or \blacktriangledown key to access the second level functions.
- Press Chem Frm to convert the result to $\mu g/L$ of chromate (Cr0₄^{2⁻}) and dichromate (Cr₂0₇^{2⁻}).



• Press the \blacktriangle or \blacktriangledown key to return to the measurement screen.

INTERFERENCES

- Vanadium above 1 mg/L, wait 10 minutes before reading to remove the interference
- Iron above 1 mg/L
- Mercurous and mercuric ions slight inhibition of the reaction

10.12. COPPER LOW RANGE

SPECIFICATIONS

Range	0.000 to 1.500 mg/L (as Cu ²⁺)
Resolution	0.001 mg/L
Accuracy	\pm 0.010 mg/L \pm 5% of reading at 25 °C
Light Source	LED with narrow band interference filter @ 575 nm
Method	Adaptation of the EPA Method

REQUIRED REAGENTS

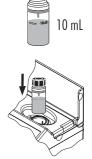
Code	Description	Quantity
HI95747-0	Copper Low Range Reagent	1 packet

REAGENT SETS

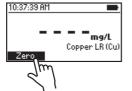
HI95747-01	Reagents for 100 tests
HI95747-03	Reagents for 300 tests
For other accessories s	ee ACCESSORIES section.

MEASUREMENT PROCEDURE

- Select the Copper LR method using the procedure described in the METHOD SELECTION section.
- Fill the cuvette with 10 mL of unreacted sample (up to the mark). Replace the plastic stopper and the cap.



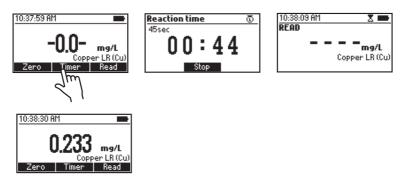
- Insert the cuvette into the holder and close the lid.
- Press Zero. The display will show "-0.0-" when the meter is zeroed and ready for measurement.



10:37:51 AM	2 🖬
ZERO	
	mg/L Copper LR (Cu)
	Copper LK (CU)



- Remove the cuvette.
- Add one packet of H195747-0 Copper Low Range Reagent. Replace the plastic stopper and the cap. Shake gently for about 15 seconds.
- Insert the cuvette into the holder and close the lid.
- Press **Timer** and the display will show the countdown prior to the measurement or wait 45 seconds and press **Read**. When the timer ends the meter will perform the reading. The instrument displays the results in **mg/L** of **copper (Cu)**.



INTERFERENCES

- Cyanide, Silver
- For samples overcoming buffering capacity of reagent around pH 6.8, pH should be adjusted between 6 and 8

10.13. COPPER HIGH RANGE

SPECIFICATIONS

Range	0.00 to 5.00 mg/L (as Cu ²⁺)
Resolution	0.01 mg/L
Accuracy	\pm 0.02 mg/L \pm 4% of reading at 25 °C
Light Source	LED with narrow band interference filter @ 575 nm
Method	Adaptation of the EPA Method

REQUIRED REAGENTS

Code	Description	Quantity
HI93702-0	Copper High Range Reagent	1 packet

REAGENT SETS

HI93702-01	Reagents for 100 tests
HI93702-03	Reagents for 300 tests
For other accessories	see ACCESSORIES section.

MEASUREMENT PROCEDURE

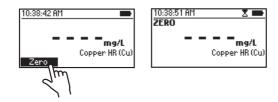
- Select the Copper HR method using the procedure described in the METHOD SELECTION section.
- Fill the cuvette with 10 mL of unreacted sample (up to the mark). Replace the plastic stopper and the cap.

• Insert the cuvette into the holder and close the lid.





• Press Zero. The display will show "-0.0-" when the meter is zeroed and ready for measurement.

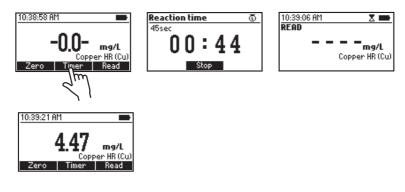




- Remove the cuvette.
- Add one packet of H193702-0 Copper High Range Reagent. Replace the plastic stopper and the cap. Shake gently for about 15 seconds.
- Insert the cuvette into the holder and close the lid.



• Press **Timer** and the display will show the countdown prior to the measurement or wait 45 seconds and press **Read**. When the timer ends the meter will perform the reading. The instrument displays the results in **mg/L** of **copper (Cu)**.



INTERFERENCES

- Cyanide, Silver
- For samples overcoming buffering capacity of reagent around pH 6.8, pH should be adjusted between 6 and 8

10.14. HYDRAZINE

SPECIFICATIONS

Range	0 to 400 μ g/L (as N ₂ H ₄)
Resolution	1 μg/L
Accuracy	\pm 4% of full scale reading at 25 °C
Light Source	LED with narrow band interference filter @ 466 nm
Method	Adaptation of the ASTM Manual of Water and Environmental Technology,
	Method D1385, p-Dimethylaminobenzaldehyde Method

REQUIRED REAGENT

Code	Description	Quantity
HI93704-0	Hydrazine Reagent	24 drops

REAGENT SETS

HI93704-01	Reagents for 100 tests
HI93704-03	Reagents for 300 tests
For other accessorie	es see ACCESSORIES section.

MEASUREMENT PROCEDURE

- Select the Hydrazine method using the procedure described in the METHOD SELECTION section.
- Fill one cuvette (#1) with 10 mL of deionized water (up to the mark).



10 mL

#2

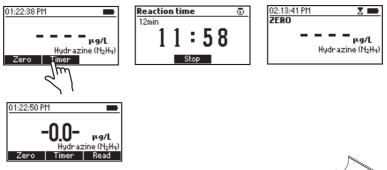
- Fill a second cuvette (#2) with 10 mL of unreacted sample (up to the mark).
- Add 12 drops of the HI93704-0 Hydrazine Reagent to each cuvette. Replace the plastic stoppers and the caps. Shake gently to mix (about 30 seconds).







- Insert the cuvette (#1) into the holder and close the lid.
- Press Timer and the display will show the countdown prior to zeroing the blank or wait 12 minutes and press Zero. The display will show "-0.0-" when the meter is zeroed and ready for measurement.

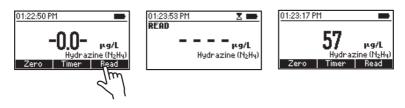


- Remove the blank.
- Insert the cuvette with the reacted sample (#2) into the holder and close the lid.



#

• Press Read to start the reading. The instrument displays concentration in μ g/L of hydrazine (N₂H₄).



INTERFERENCES

- Highly colored samples
- Highly turbid samples
- Aromatic amines

10.15. IRON LOW RANGE

SPECIFICATIONS

Range	0.000 to 1.600 mg/L (as Fe)
Resolution	0.001 mg/L
Accuracy	\pm 0.010 mg/L \pm 8% of reading at 25 °C
Light Source	LED with narrow band interference filter @ 575 nm
Method	Adaptation of the TPTZ Method

REQUIRED REAGENTS

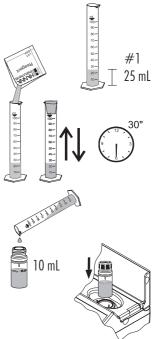
Code	Description	Quantity
HI93746-0	Iron Low Range Reagent	2 packets

REAGENT SETS

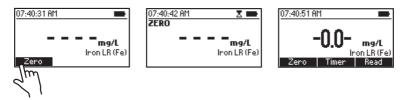
HI93746-01	Reagents for 50 tests
HI93746-03	Reagents for 150 tests
For other accessories	see ACCESSORIES section.

MEASUREMENT PROCEDURE

- Select the Iron LR method using the procedure described in the METHOD SELECTION section.
- Fill one graduated mixing cylinder up to the 25 mL mark with deionized water.
- Add one packet of H193746-0 Iron Low Range Reagent, close the cylinder and shake vigorously for 30 seconds. This is the blank.
- Fill a cuvette with 10 mL of the blank (up to the mark). Replace the rubber stopper.
- Insert the cuvette into the holder and close the lid.



• Press Zero. The display will show "-0.0-" when the meter is zeroed and ready for measurement.

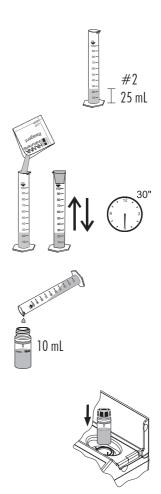


- Remove the cuvette.
- Fill another graduated mixing cylinder up to the 25 mL mark with the sample.

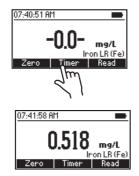
• Add one packet of H193746-0 Iron Low Range Reagent, close the cylinder and shake vigorously for 30 seconds. This is the reacted sample.

• Fill a cuvette with 10 mL of the reacted sample (up to the mark). Replace the rubber stopper.

• Insert the sample into the holder and close the lid.



• Press **Timer** and the display will show the countdown prior to the measurement or wait for 30 seconds and press **Read**. When the timer ends the meter will perform the reading. The instrument displays concentration in **mg/L** of **iron (Fe)**.





2 🗪
mg/L
Iron LR (Fe)

INTERFERENCES

- Manganese above 50.0 mg/L
- Cadmium, Molybdenum above 4.0 mg/L
- Cyanide above 2.8 mg/L
- Chromium(VI) above 1.2 mg/L
- Nickel above 1.0 mg/L
- Nitrite ion above 0.8 mg/L
- Copper above 0.6 mg/L
- Mercury above 0.4 mg/L
- Chromium(III) above 0.25 mg/L
- Cobalt above 0.05 mg/L
- Sample pH should be between 3 and 4 to avoid fading or turbidity formation

10.16. IRON HIGH RANGE

SPECIFICATIONS

Range	0.00 to 5.00 mg/L (as Fe)
Resolution	0.01 mg/L
Accuracy	\pm 0.04 mg/L \pm 2% of reading at 25 °C
Light Source	LED with narrow band interference filter @ 525 nm
Method	Adaptation of Standard Methods for the Examination of Water and
	Wastewater, 23 rd Edition, 3500-Fe B, Phenanthroline Method

REQUIRED REAGENTS

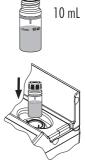
Code	Description	Quantity
HI93721-0	Iron High Range Reagent	1 packet

REAGENT SETS

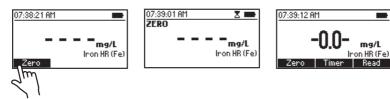
HI93721-01	Reagents for 100 tests
HI93721-03	Reagents for 300 tests
For other accessorie	es see ACCESSORIES section

MEASUREMENT PROCEDURE

- Select the Iron HR method using the procedure described in the METHOD SELECTION section.
- Fill the cuvette with 10 mL of unreacted sample (up to the mark). Replace the plastic stopper and the cap.



- Insert the cuvette into the holder and close the lid.
- Press Zero. The display will show "-0.0-" the meter is zeroed and ready for measurement.



- Remove the cuvette and add the content of one packet of H193721-0 Iron High Range Reagent. Replace the plastic stopper and the cap. Shake until powder is completely dissolved.
- Insert the cuvette into the holder and close the lid.





• Press **Timer** and the display will show the countdown prior to the measurement or wait 3 minutes and press **Read**. When the timer ends the meter will perform the reading. The instrument displays the result in **mg/L** of **iron (Fe)**.





07:39:46 AM		2 🗪
READ		
-	-	mg/L Iron HR (Fe)

INTERFERENCES

- Chloride above 185000 mg/L
- Magnesium above 100000 mg/L CaCO₃
- Calcium above 10000 mg/L CaCO₃
- Molybdate Molybdenum above 50 mg/L

10.17. IRON(II)

SPECIFICATIONS

Range	0.00 to 6.00 mg/L (as Fe ²⁺)
Resolution	0.01 mg/L
Accuracy	\pm 0.10 mg/L \pm 2% of reading at 25 °C
Light Source	LED with narrow band interference filter @ 525 nm
Method	Adaptation of Standard Methods for the Examination of Water and
	Wastewater, 23 rd Edition, 3500-Fe B, Phenanthroline Method

REQUIRED REAGENTS

Code	Description	Quantity
HI96776-0	Iron(II) Reagent	1 packet

REAGENTS SETS

HI96776-01	Reagents for 100 tests
HI96776-03	Reagents for 300 tests
For other accessories	see ACCESSORIES section

PRINCIPLE

In aqueous solution, reactive ferrous iron (Fe^{2+}) reacts with 1,10-phenanthroline to form an orange-red complex.

APPLICATION

Surface water, drinking water, mineral and groundwater, process control

SIGNIFICANCE & USE

Surface water typically contains up to 0.7 mg/L of iron. Drinking water typically contains up to 0.3 mg/L of iron, but this level may increase significantly if plumbing fixtures contain iron. In well-oxygenated, non-acidic waters, iron exists mainly in the ferric form (Fe^{3+}) and will precipitate as iron oxide hydroxide (FeO(OH)). However, anoxic water may have high levels of dissolved ferrous iron (Fe^{2+}) which could precipitate in heating/cooling systems or other equipment after exposure to air. The Iron(II) method measures the ferrous (Fe^{2+}) form of iron.

MEASUREMENT PROCEDURE

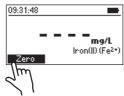
Warning: Method is temperature-dependent. Sample temperature must be between 18 °C and 22 °C.

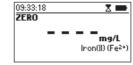
- Select the Iron(II) method using the procedure described in the METHOD SELECTION section.
- Fill the cuvette with 10 mL of unreacted sample (up to the mark). Replace the plastic stopper and the cap.
- Insert the cuvette into the holder and close the lid.





• Press Zero. The display will show "-0.0-"; the meter is zeroed and ready for measurement.







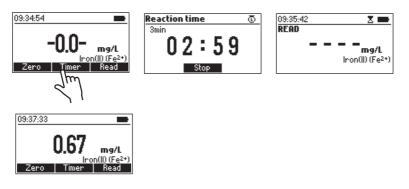
• Remove the cuvette and add the content of one packet of H196776-0 Iron(II) Reagent. Replace the plastic stopper and the cap. Shake gently for 30 seconds.





• Insert the cuvette into the holder and close the lid.

 Press Timer and the display will show the countdown prior to the measurement or wait 3 minutes and press Read. The instrument displays the result in mg/L of Iron (Fe²⁺).



Warning: Timing is critical for accurate measurement. Reaction times beyond 3 minutes may cause some ferric iron (Fe^{3+}) to also react, producing false high measurements.

INTERFERENCES

- Chloride, Sulfate above 1000 mg/L
- Ammonium, Calcium, Potassium, Sodium above 500 mg/L
- Silver above 100 mg/L
- Carbonate, Chromium(III) and (VI), Cobalt, Lead, Mercury, Nitrate, Zinc above 50 mg/L
- Nickel above 25 mg/L
- Copper above 10 mg/L
- Tin above 5 mg/L
- Extreme pH or highly buffered samples, the pH of the sample must be between 3.8 and 5.5 after addition of the reagent

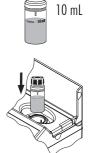
10.18. MOLYBDENUM

SPECIFICATIONS

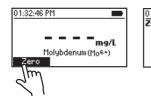
JILCHICAHONS				
Range	0.0 to 40.0 mg/L (as Mo^{6+})			
Resolution	0.1 mg/L	0.1 mg/L		
Accuracy	\pm 0.3 mg/L \pm 5% of reading	∣at 25 °C		
Light Source	LED with narrow band interfere	ence filter @ 420 nm		
Method	Adaptation of the Mercaptoace	Adaptation of the Mercaptoacetic Acid Method		
REQUIRED REAG	ENTS			
Code	Description	Quantity		
HI93730A-0	Molybdenum Reagent A	1 packet		
HI93730B-0	Molybdenum Reagent B	1 packet		
HI93730C-0	Molybdenum Reagent C	1 packet		
REAGENT SETS				
HI93730-01	Reagents for 100 tests			
HI93730-03	Reagents for 300 tests			
For other accessorie	es see ACCESSORIES section.			

MEASUREMENT PROCEDURE

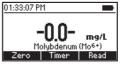
- Select the Molybdenum method using the procedure described in the METHOD SELECTION section.
- Fill the cuvette with 10 mL of unreacted sample (up to the mark). Replace the plastic stopper and the cap.



- Insert the cuvette into the holder and close the lid.
- Press Zero. The display will show "-0.0-" when the meter is zeroed and ready for measurement.



1:32:5	/ Pfi			
ERO				
				mg/L
	Mol	ybdi	enur	n(Mo6+)

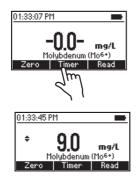


• Fill one graduated mixing cylinder up to the 25 mL mark with the sample.

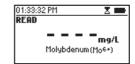
- Add one packet of HI93730A-0 Molybdenum Reagent A to the cylinder, close and invert several times until completely dissolved.
- Add one packet of H193730B-0 Molybdenum Reagent B to the cylinder, close and invert several times until completely dissolved.
- Add one packet of H193730C-0 Molybdenum Reagent C to the cylinder, close and shake vigorously.
- Fill an empty cuvette with 10 mL of reacted sample (up to the mark). Replace the plastic stopper and the cap.
- Insert the cuvette into the holder and close the lid.



 Press Timer and the display will show the countdown prior to the measurement or wait 5 minutes and press Read. When the timer ends the meter will perform the reading. The instrument displays concentration in mg/L of molybdenum (Mo⁶⁺).

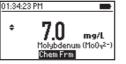






- Press the \blacktriangle or \blacktriangledown key to access the second level functions.
- Press Chem Frm to convert the result to mg/L of molybdate (MoO₄²⁻) and sodium molybdate (Na₂MoO₄).







• Press the \blacktriangle or \blacktriangledown key to return to the measurement screen.

INTERFERENCES

- Chromium above 1000 mg/L
- Sulfate above 200 mg/L
- Aluminum, Iron, Nickel above 50 mg/L
- Copper above 10 mg/L
- Nitrite must be absent
- Highly buffered samples or samples with extreme pH may exceed the buffering capacity of the reagents

10.19. NITRATE

SPECIFICATIONS

Range	0.0 to 30.0 mg/L (as NO ₃ ⁻ - N)
Resolution	0.1 mg/L
Accuracy	\pm 0.5 mg/L \pm 10% of reading at 25 °C
Light Source	LED with narrow band interference filter $@$ 525 nm
Method	Adaptation of the Cadmium Reduction Method

REQUIRED REAGENTS

Code	Description	Quantity
HI93728-0	Nitrate Reagent	1 packet

REAGENT SETS

HI93728-01	Reagents for 100 tests
HI93728-03	Reagents for 300 tests
For other accessori	es see ACCESSORIES section.

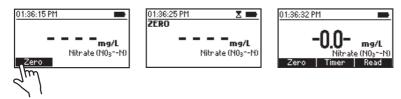
MEASUREMENT PROCEDURE

- Select the Nitrate method using the procedure described in the METHOD SELECTION section.
- Fill the cuvette with 10 mL of sample (up to the mark). Replace the plastic stopper and the cap.
- Insert the cuvette into the holder and close the lid.





• Press Zero. The display will show "-0.0-" when the meter is zeroed and ready for measurement.



- Remove the cuvette and add one packet of H193728-0 Nitrate Reagent.
- Replace the plastic stopper and the cap. Shake vigorously up and down for exactly 10 seconds. Continue to mix by inverting the cuvette gently for 50 seconds, while taking care not to induce air bubbles. Powder will not completely dissolve.

Note: The method is technique sensitive. See procedure described in CUVETTE PREPARATION section for proper mixing technique.

• Insert the cuvette into the holder and close the lid.

mg/L

Bead

Nitrate (N0s7-N

Nitnate (N0s=-N)

01:36:32 PM

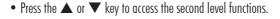
Zero

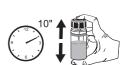
01:38:32 PM

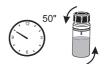
 Press Timer and the display will show the countdown prior to the measurement or wait 4 minutes and 30 seconds and press Read. When the timer ends the meter will perform the reading. The instrument displays the results in mg/L of nitrate-nitrogen (NO₃-N).

Reaction time

4.5min







01:38:23 PM

READ

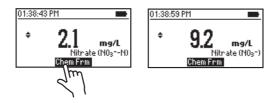
ō

28



≍ =

mg/L Nitrate (N0₃⁻-N) • Press Chem Frm to convert the result to mg/L of nitrate (NO₃⁻).



- Press the \blacktriangle or \blacktriangledown key to return to the measurement screen.

INTERFERENCES

- Ammonia and amines, as urea and primary aliphatic amines
- Chloride above 100 mg/L
- Chlorine above 2 mg/L
- Copper, Iron (Ferric), Strong oxidizing and reducing substances
- Sulfide must be absent

10.20. NITRITE LOW RANGE

SPECIFICATIONS

Range	0 to 600 μ g/L (as NO ₂ ⁻ -N)
Resolution	1 µg/L
Accuracy	\pm 20 μ g/L \pm 4% of reading at 25 °C
Light Source	LED with narrow band interference filter $@$ 466 nm
Method	Adaptation of the EPA Diazotization Method 354.1

REQUIRED REAGENTS

Code	Description	Quantity
HI93707-0	Nitrite Low Range Reagent	1 packet

REAGENT SETS

HI93707-01	Reagents for 100 tests
HI93707-03	Reagents for 300 tests
For other accessories	see ACCESSORIES section.

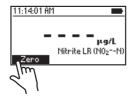
MEASUREMENT PROCEDURE

- Select the Nitrite LR method using the procedure described in the METHOD SELECTION section.
- Fill the cuvette with 10 mL of unreacted sample (up to the mark). Replace the plastic stopper and the cap.
- Insert the cuvette into the holder and close the lid.





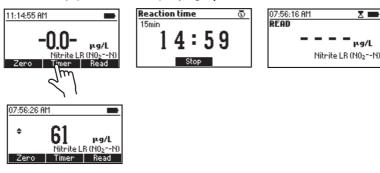
• Press Zero. The display will show "-0.0-" when the meter is zeroed and ready for measurement.



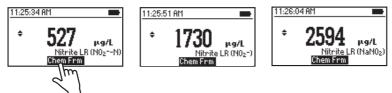
11:14:22 AM	2 📟
ZERO	
-	
	μg/L
	Nitrite LR (NO ₂ - N)

11:14:55 A	М	-
	-0 0-	
	0.0	µg/L
	Nitrite L	R (N02N)
Zero	Timer	Read

- Remove the cuvette.
- Add one packet of HI93707-0 Nitrite Low Range Reagent.
- Replace the plastic stopper and the cap. Shake gently for about 15 seconds.
- Insert the cuvette into the holder and close the lid.
- Press Timer and the display will show the countdown prior to the measurement or wait 15 minutes
- and press **Read**. When the timer ends the meter will perform the reading. The instrument displays concentration in μ g/L of nitrite-nitrogen (NO₂⁻-N).



- Press the \blacktriangle or \blacktriangledown key to access the second level functions.
- Press Chem Frm to convert the result to μ g/L of nitrite (NO₂⁻) and sodium nitrite (NaNO₂).



• Press the \blacktriangle or \blacktriangledown key to return to the measurement screen.

INTERFERENCES

- Antimonious, Auric, Bismuth, Chloroplatinate ions, Cupric, Iron (Ferric), Iron (Ferrous), Lead, Mercurous, Silver, Strong reducing or oxidating agents
- Nitrate above 100 mg/L could yield falsely high readings

10.21. NITRITE HIGH RANGE

SPECIFICATIONS

Range	0 to 150 mg/L (as NO_2^{-})
Resolution	1 mg/L
Accuracy	\pm 4 mg/L \pm 4% of reading at 25 °C
Light Source	LED with narrow band interference filter @ 575 nm
Method	Adaptation of the Ferrous Sulfate Method

REQUIRED REAGENTS

Code	Description	Quantity
HI93708-0	Nitrite High Range Reagent	1 packet

REAGENT SETS

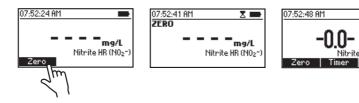
HI93708-01	Reagents for 100 tests
HI93708-03	Reagents for 300 tests
For other accessories	see ACCESSORIES section.

MEASUREMENT PROCEDURE

- Select the Nitrite HR method using the procedure described in the METHOD SELECTION section.
- Fill the cuvette with 10 mL of unreacted sample (up to the mark). Replace the plastic stopper and the cap.

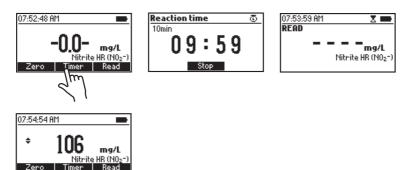
10 mL

- Insert the cuvette into the holder and close the lid.
- Press Zero. The display will show "-0.0-" when the meter is zeroed and ready for measurement.

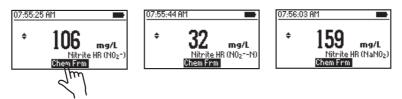


- Remove the cuvette.
- Add one packet of H193708-0 Nitrite High Range Reagent. Replace the plastic stopper and the cap. Shake gently until completely dissolved.

- Insert the cuvette into the holder and close the lid.
- Press Timer and the display will show the countdown prior to the measurement or wait 10 minutes and press Read. When the timer ends the meter will perform the reading. The instrument displays concentration in mg/L of nitrite (NO₂⁻).



- Press the \blacktriangle or \blacktriangledown key to access the second level functions.
- Press Chem Frm to convert the result to mg/L of nitrite-nitrogen (NO₂⁻-N) and sodium nitrite (NaNO₂).



• Press the \blacktriangle or \blacktriangledown key to return to the measurement screen.

10.22. OXYGEN, DISSOLVED

SPECIFICATIONS

Range	0.0 to 10.0 mg/L (as 0 ₂)
Resolution	0.1 mg/L
Accuracy	\pm 0.4 mg/L \pm 3% of reading at 25 °C
Light Source	LED with narrow band interference filter @ 420 nm
Method	Adaptation of Standard Methods for the Examination of Water and
	Wastewater, 18 th Edition, Azide Modified Winkler Method

REQUIRED REAGENTS

Code	Description	Quantity
HI93732A-0	Dissolved Oxygen Reagent A	5 drops
HI93732B-0	Dissolved Oxygen Reagent B	5 drops
HI93732C-0	Dissolved Oxygen Reagent C	10 drops

REAGENT SET

HI93732-01	Reagents for 100 tests
HI93732-03	Reagents for 300 tests
For other accessori	es see ACCESSORIES section

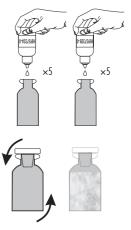
MEASUREMENT PROCEDURE

- Select the Oxygen (dissolved) method using the procedure described in the METHOD SELECTION section.
- Fill one 60 mL glass bottle completely with the unreacted sample.
- Replace the cap and ensure that a small part of the sample spills over.
- Remove the cap and add 5 drops of H193732A-0 and 5 drops of H193732B-0.
- Add more sample to fill the bottle completely. Replace the cap and ensure that a part of the sample spills over.

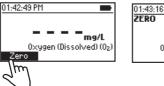
Note: This ensures no air bubbles have been trapped inside the bottle. Trapped air bubbles could alter readings.

• Invert the bottle several times until the sample turns orangeyellow and a flocculating agent appears.

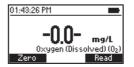




- Let the sample stand for approximately 2 minutes to allow flocculating agent to settle.
- When the upper half of the bottle is clear, add 10 drops of H193732C-0 Dissolved Oxygen Reagent C.
- Replace the cap and invert the bottle until the settled flocculating agent dissolves completely. The sample is ready for measurement when it is yellow and completely clear.
- Fill the first cuvette (#1) with 10 mL of the unreacted sample (up to the mark). Replace the plastic stopper and the cap.
- Insert the cuvette into the holder and close the lid.
- Press Zero. The display will show "-0.0-" when the meter is zeroed and ready for measurement.



01:43:16 PM	Σ 🗪
ZERO	
	_
	mg/L
0xygen (Dis	solved) (0 ₂)



- Remove the cuvette.
- Fill second cuvette (#2) with 10 mL of the reacted sample (up to the mark). Replace the plastic stopper and the cap.





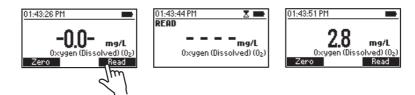
×10



• Insert the cuvette into the holder and close the lid.



• Press Read to start the reading. The instrument will display the results in mg/L of oxygen (0₂).



INTERFERENCES

Interference may be caused by:

• Reducing and oxidizing materials

10.23. OXYGEN SCAVENGERS (CARBOHYDRAZIDE)

SPECIFICATIONS

Range	0.00 to 1.50 mg/L (as Carbohydrazide)
Resolution	0.01 mg/L
Accuracy	\pm 0.02 mg/L \pm 3% of reading at 25 °C
Light Source	LED with narrow band interference filter @ 575 nm
Method	Adaptation of the Iron Reduction Method

REQUIRED REAGENTS

Code	Description	Quantity
HI96773A-0	Oxygen Scavengers Reagent A	2 packets
HI96773B-0	Oxygen Scavengers Reagent B	1 mL

REAGENT SET

HI96773-01	Reagents for 50 tests
HI96773-03	Reagents for 150 tests
For other accessor	ies see ACCESSORIES section

MEASUREMENT PROCEDURE

- Select the Oxy. Scavengers (Carbohy) method using the procedure described in the METHOD SELECTION section.
- Fill first cuvette (#1) with 10 mL of deionized water (up to the mark).

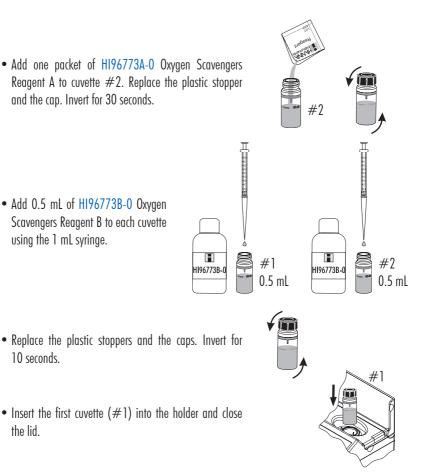


 Add one packet of H196773A-0 Oxygen Scavengers Reagent A to cuvette #1. Replace the plastic stopper and the cap. Invert for 30 seconds.

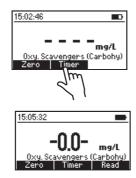








• Press **Timer** and the display will show countdown prior to the measurement or wait 10 minutes and press **Zero**. The display will show "-0.0-" when the meter is zeroed and ready for measurement.



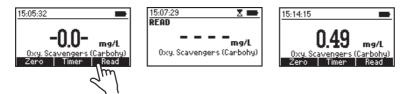


15:04:06
ZERO
— — — — _{mg/L}
0xy. Scavengers (Carbohy)

- Remove the cuvette.
- Insert the second cuvette (#2) into the holder and close the lid.



• Press Read to start reading. The instrument displays the results in mg/L of carbohydrazide.



INTERFERENCES

Interference may be caused by:

• Borate (as Na₂B₄O₇), Cobalt, Copper, Iron (Ferrous), Hardness (as CaCO₃), Light, Lignosulfonates, Manganese, Molybdenum, Nickel, Phosphate, Phosphonates, Sulfate, Temperature and Zinc

10.24. OXYGEN SCAVENGERS (DIETHYLHYDROXYLAMINE) (DEHA)

SPECIFICATIONS

Range	0 to 1000 µg/L (as DEHA)	
Resolution	1 µg/L	
Accuracy	\pm 5 μ g/L \pm 5% of reading at 25 $^\circ$	C
Light Source	LED with narrow band interference filter @ 575 nm	
Method	Adaptation of the Iron Reduction Method	
REQUIRED REAGENTS		
Code	Description	Quantity
HI96773A-0	Oxygen Scavengers Reagent A	2 packets
HI96773B-0	Oxygen Scavengers Reagent B	1 mL
REAGENT SET		
HI96773-01	Reagents for 50 tests	
HI96773-03	Reagents for 150 tests	
For other accessories see ACCESSORIES section.		

MEASUREMENT PROCEDURE

- Select the Oxy. Scavengers (DEHA) method using the procedure described in the METHOD SELECTION section.
- Fill first cuvette (#1) with 10 mL of deionized water (up to the mark).



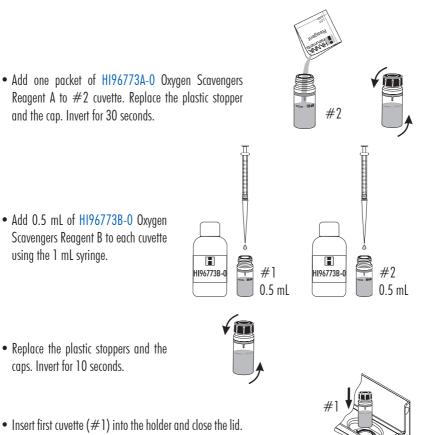
10 mL

#2

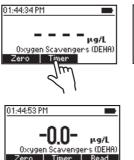
• Fill second cuvette (#2) with 10 mL of sample (up to the mark).



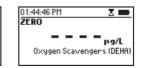




- Press Timer and the display will show countdown prior to the measurement or wait 10 minutes and press Zero. The display will show "-0.0-" when the meter is zeroed and ready for measurement.



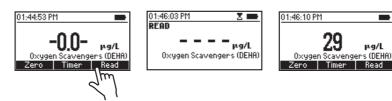




- Remove the cuvette.
- Insert the second cuvette (#2) into the holder and close the lid.



• Press **Read** to start reading. The instrument displays the results in μ g/L of DEHA.



INTERFERENCES

Interference may be caused by:

 Borate (as Na₂B₄O₇), Cobalt, Copper, Iron (Ferrous), Hardness (as CaCO₃), Light, Lignosulfonates, Manganese, Molybdenum, Nickel, Phosphate, Phosphonates, Sulfate, Temperature and Zinc

10.25. OXYGEN SCAVENGERS (HYDROQUINONE)

SPECIFICATIONS

Range	0.00 to 2.50 mg/L (as Hydroquinone)
Resolution	0.01 mg/L
Accuracy	\pm 0.04 mg/L \pm 3% of reading at 25 °C
Light Source	LED with narrow band interference filter $@$ 575 nm
Method	Adaptation of the Iron Reduction Method

REQUIRED REAGENTS

Code	Description	Quantity
HI96773A-0	Oxygen Scavengers Reagent A	2 packets
HI96773B-0	Oxygen Scavengers Reagent B	1 mL

REAGENT SET

HI96773-01	Reagents for 50 tests
HI96773-03	Reagents for 150 tests
For other accessor	ies see ACCESSORIES section

MEASUREMENT PROCEDURE

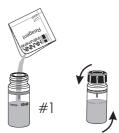
- Select the Oxy. Scavengers (Hydro) method using the procedure described in the METHOD SELECTION section.
- Fill first cuvette (#1) with 10 mL of deionized water (up to the mark).

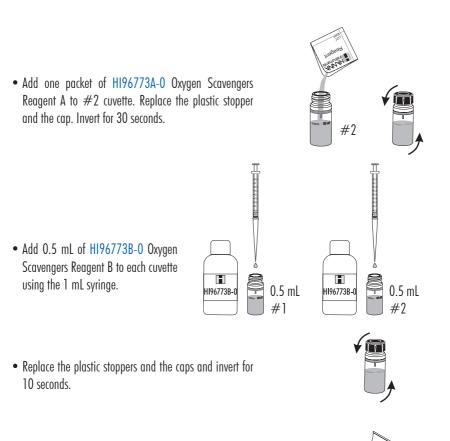


• Add one packet of H196773A-0 Oxygen Scavengers Reagent A to #1 cuvette. Replace the plastic stopper and the cap. Invert for 30 seconds.

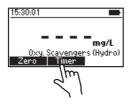








- Insert first cuvette (#1) into the holder and close the lid.
- Press **Timer** and the display will show countdown prior to the measurement or wait 2 minutes and press **Zero**. The display will show "-0.0-" when the meter is zeroed and ready for measurement.

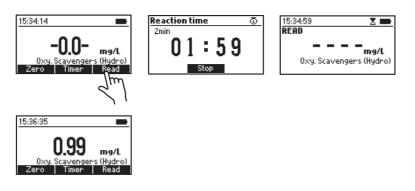


15:31:55	Z 🖚
ZERO	
• Oxy.	mg/L Scavengers (Hydro)



#

- Remove the cuvette.
- Insert the second cuvette (# 2) into the holder and close the lid.
- Press **Read** to start reading. The instrument displays the results in **mg/L** of **hydroquinone**.



INTERFERENCES

Interference may be caused by:

 Borate (as Na₂B₄O₇), Cobalt, Copper, Iron (Ferrous), Hardness (as CaCO₃), Light, Lignosulfonates, Manganese, Molybdenum, Nickel, Phosphate, Phosphonates, Sulfate, Temperature and Zinc

10.26. OXYGEN SCAVENGERS (ISO-ASCORBIC ACID)

SPECIFICATIONS

Range	0.00 to 4.50 mg/L (as Iso-Ascorbic Acid)
Resolution	0.01 mg/L
Accuracy	\pm 0.03 mg/L \pm 3 % of reading at 25 °C
Light Source	LED with narrow band interference filter @ 575 nm
Method	Adaptation of the Iron Reduction Method

REQUIRED REAGENTS

Code	Description	Quantity
HI96773A-0	Oxygen Scavengers Reagent A	2 packets
HI96773B-0	Oxygen Scavengers Reagent B	1 mL

REAGENT SET

HI96773-01	Reagents for 50 tests	
HI96773-03	Reagents for 150 tests	
For other accessories see ACCESSORIES section		

MEASUREMENT PROCEDURE

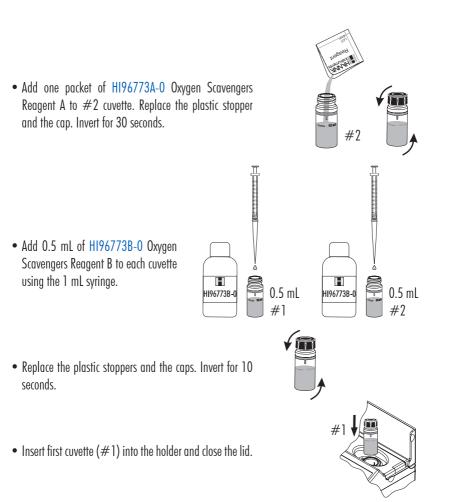
- Select the Oxygen Scavengers (ISA) method using the procedure described in the METHOD SELECTION section.
- Fill first cuvette (#1) with 10 mL of deionized water (up to the mark).
- Fill second cuvette (#2) with 10 mL of sample (up to the mark).

 Add one packet of H196773A-0 Oxygen Scavengers Reagent A to #1 cuvette. Replace the plastic stopper and the cap. Invert for 30 seconds.

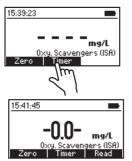




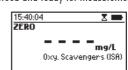




• Press **Timer** and the display will show countdown prior to the measurement or wait 10 minutes and press **Zero**. The display will show "-0.0-" when the meter is zeroed and ready for measurement.



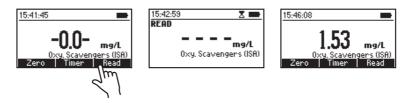
Reaction time () 10min 09:57 Stop



- Remove the cuvette.
- Insert the second cuvette (#2) into the holder and close the lid.



• Press Read to start reading. The instrument displays the results in mg/L of Iso-ascorbic acid.



INTERFERENCES

Interference may be caused by:

 Borate (as Na₂B₄O₇), Cobalt, Copper, Iron (Ferrous), Hardness (as CaCO₃), Light, Lignosulfonates, Manganese, Molybdenum, Nickel, Phosphate, Phosphonates, Sulfate, Temperature and Zinc

F

10.27. pH

SPECIFICATIONS

Range	6.5 to 8.5 pH
Resolution	0.1 pH
Accuracy	\pm 0.1 pH at 25 °C
Light Source	LED with narrow band interference filter @ 525 nm
Method	Adaptation of the Phenol Red Method

REQUIRED REAGENTS

Code	Description	Quantity
HI93710-0	pH Reagent	5 drops

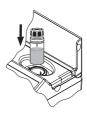
REAGENT SETS

HI93710-01	Reagents for 100 tests
HI93710-03	Reagents for 300 tests
For other accessorie	es see ACCESSORIES section.

MEASUREMENT PROCEDURE

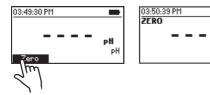
- Select the pH method using the procedure described in the METHOD SELECTION section.
- Fill the cuvette with 10 mL of unreacted sample (up to the mark). Replace the plastic stopper and the cap.
- Insert the cuvette into the holder and close the lid.





nН

• Press Zero. The display will show "-0.0-" when the meter is zeroed and ready for measurement.





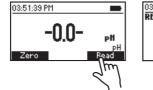
• Remove the cuvette and add 5 drops of H193710-0 pH Reagent Indicator. Replace the plastic stopper and the cap and mix the solution.

• Insert the cuvette into the holder and close the lid.

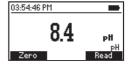


×5

• Press **Read** to start the reading. The instrument displays the result in **pH**.



03:53:4	5 PM				Σ 🖿
READ					
	-	-	-	-	рĦ
					PH



PHOSPHATE LOW RANGE

10.28. PHOSPHATE LOW RANGE

SPECIFICATIONS

Range	0.00 to 2.50 mg/L (as PO_4^{3-})
Resolution	0.01 mg/L
Accuracy	\pm 0.04 mg/L \pm 4% of reading at 25 °C
Light Source	LED with narrow band interference filter $\textcircled{0}$ 610 nm
Method	Adaptation of the Ascorbic Acid Method

REQUIRED REAGENTS

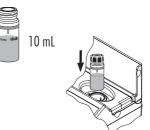
Code	Description	Quantity
HI93713-0	Phosphate Low Range Reagent	1 packet

REAGENT SETS

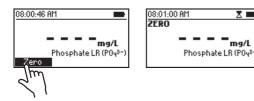
HI93713-01	Reagents for 100 tests
HI93713-03	Reagents for 300 tests
For other accessori	es see ACCESSORIES section.

MEASUREMENT PROCEDURE

- Select the Phosphate LR method using the procedure described in the METHOD SELECTION section.
- Rinseand replace the plastic stopper and the cap. Shake the cuvette several times with unreacted sample.
- Fill the cuvette with 10 mL of sample (up to the mark). Replace the plastic stopper and the cap.



- Insert the cuvette into the holder and close the lid.
- Press Zero. The display will show "-0.0-" when the meter is zeroed and ready for measurement.



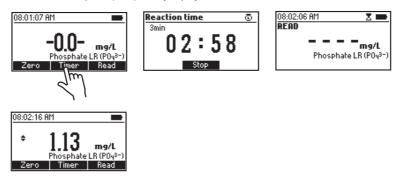
• Remove the cuvette and add the content of one packet of H193713-0 Phosphate Low Range Reagent. Replace the plastic stopper and the cap. Shake gently (for about 2 minutes) until the powder is completely dissolved.



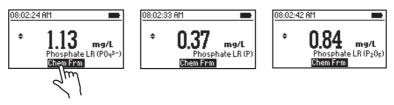
• Insert the cuvette into the holder and close the lid.



• Press **Timer** and the display will show the countdown prior to the measurement or wait 3 minutes and press **Read**. When the timer ends the meter will perform the reading. The instrument displays concentration in mg/L of phosphate (PO₄³⁻).



- Press the \blacktriangle or \blacktriangledown key to access the second level functions.
- Press Chem Frm to convert the result to mg/L of phosphorus (P) and phosphorus pentoxide (P_2O_5).



- Press the \blacktriangle or \blacktriangledown key to return to the measurement screen.

INTERFERENCES

- Iron, Silica above 50 mg/L
- Copper, Silicate above 10 mg/L
- Arsenate, Highly buffered samples, Hydrogen sulfide, Turbid samples

10.29. PHOSPHATE HIGH RANGE

SPECIFICATIONS

Range	0.0 to 30.0 mg/L (as PO_4^{3-})
Resolution	0.1 mg/L
Accuracy	\pm 1.0 mg/L \pm 4% of reading at 25 °C
Light Source	LED with narrow band interference filter $@$ 525 nm
Method	Adaptation of Standard Methods for the Examination of Water and
	Wastewater, 18 th Edition, Amino Acid Method

REQUIRED REAGENTS

Code	Description	Quantity
HI93717A-0	Phosphate High Range Reagent A	10 drops
HI93717B-0	Phosphate High Range Reagent B	1 packet

REAGENT SETS

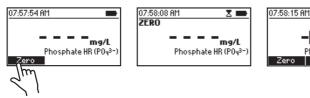
HI93717-01	Reagents for 100 tests
HI93717-03	Reagents for 300 tests
For other accessorie	es see ACCESSORIES section

MEASUREMENT PROCEDURE

- Select the Phosphate HR method using the procedure described in the METHOD SELECTION section.
- Fill the cuvette with 10 mL of unreacted sample (up to the mark). Replace the plastic stopper and the cap.



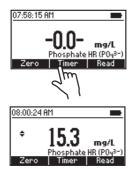
• Press Zero. The display will show "-0.0-" when the meter is zeroed and ready for measurement.



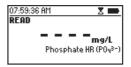




- Add 10 drops of HI93717A-0 Phosphate HR Reagent A.
- Add one packet of HI93717B-0 Phosphate HR Reagent B to the cuvette. Replace the plastic stopper and the cap. Shake gently until completely dissolved.
- Insert the cuvette into the holder and close the lid.
- Press Timer and the display will show the countdown prior to the measurement or wait 5 minutes and press Read. When the timer ends the meter will perform the reading. The instrument displays the results in mg/L of phosphate (PO₄³⁻).





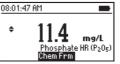


∛ ×10

- Press the \blacktriangle or \blacktriangledown key to access the second level functions.
- Press Chem Frm to convert the result to mg/L of phosphorus (P) and phosphorus pentoxide (P_2O_5).



08:01:2	3 AM	
÷	5.0	mg/L
	Chem Frm	hate HR (P)



• Press the \blacktriangle or \blacktriangledown key to return to the measurement screen.

INTERFERENCES

Interference may be caused by:

- Sulfide
- Chloride above 150000 mg/L
- Magnesium above 40000 mg/L CaCO₃
- Calcium above 10000 mg/L CaCO₃
- Iron (Ferrous) above 100 mg/L

10.30. SILICA LOW RANGE

SPECIFICATIONS

Range	0.00 to 2.00 mg/L (as SiO ₂)
Resolution	0.01 mg/L
Accuracy	\pm 0.03 mg/L \pm 3% of reading at 25 °C
Light Source	LED with narrow band interference filter @ 610 nm
Method	Adaptation of the ASTM Manual of Water and Environmental Technology,
	D859. Heteropoly Molybdenum Blue Method

REQUIRED REAGENTS

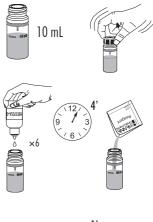
Code	Description	Quantity
HI93705A-0	Silica Low Range Reagent A	6 drops
HI93705B-0	Silica Low Range Reagent B	1 packet
HI93705C-0	Silica Low Range Reagent C	1 packet

REAGENT SETS

HI93705-01	Reagents for 100 tests
HI93705-03	Reagents for 300 tests
For other accessories	see ACCESSORIES section.

MEASUREMENT PROCEDURE

- Select the Silica LR method using the procedure described in the METHOD SELECTION section.
- Fill the cuvette with 10 mL of unreacted sample (up to the mark).
- Add 6 drops of H193705A-0 Silica LR Reagent A. Replace the plastic stopper and the cap. Swirl the solution.
- Press Timer and the display will show the countdown prior to adding H193705B-0 Silica LR Reagent B or wait 4 minutes.
- Add one packet of HI93705B-0 Silica LR Reagent B and shake until it is completely dissolved.
- Press Continue and the display will show the countdown or wait 1 minute.



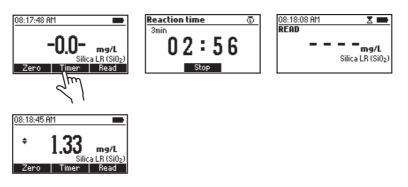


- Insert the cuvette into the holder and close the lid.
- Press Zero. The display will show "-0.0-" when the meter is zeroed and ready for measurement.



- Remove the cuvette.
- Add one packet of H193705C-0 Silica LR Reagent C and shake until it is completely dissolved.

- Insert the cuvette into the holder and close the lid.
- Press Timer and the display will show the countdown prior to the measurement or wait 3 minutes and press Read. When the timer ends the meter will perform the reading. The instrument displays result in mg/L of silica (SiO₂).



• Press the \blacktriangle or \blacktriangledown key to access the second level functions.

• Press Chem Frm to convert the result to mg/L of silicon (Si).



• Press the \blacktriangle or \blacktriangledown key to return to the measurement screen.

INTERFERENCES

Interference may be caused by:

- Phosphate above 75 mg/L, causes an 11% reduction in reading
- Phosphate above 60 mg/L, causes a 2% reduction in reading
- Sulfide and high concentration of iron
- Eliminate color and turbidity interferences by zeroing the meter with the original water sample

10.31. SILICA HIGH RANGE

SPECIFICATIONS

Range	0 to 200 mg/L (as SiO ₂)
Resolution	1 mg/L
Accuracy	\pm 1 mg/L \pm 5% of reading at 25 °C
Light Source	LED with narrow band interference filter @ 466 nm
Method	Adaptation of the US EPA Method 370.1 for Drinking, Surface and Saline
	Waters, Domestic and Industrial Wastes & Standard Method 4500-SiO ₂

REQUIRED REAGENTS

Code	Description	Quantity
HI96770A-0	Silica High Range Reagent A	1 packet
HI96770B-0	Silica High Range Reagent B	1 packet
HI96770C-0	Silica High Range Reagent C	1 packet

REAGENT SETS

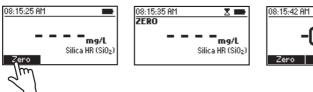
HI96770-01	Reagents for 100 tests
HI96770-03	Reagents for 300 tests
For other accessories	see ACCESSORIES section.

MEASUREMENT PROCEDURE

- Select the Silica HR method using the procedure described in the METHOD SELECTION section.
- Fill the cuvette with 10 mL of unreacted sample (up to the mark). Replace the plastic stopper and the cap.



- Insert the cuvette into the holder and close the lid.
- Press Zero. The display will show "-0.0-" when the meter is zeroed and ready for measurement.





- Remove the cuvette.
- Add one packet of H196770A-0 Silica HR Reagent A. Replace the plastic stopper and the cap. Shake vigorously until completely dissolved.
- Add one packet of H196770B-0 Silica HR Reagent B. Replace the plastic stopper and the cap. Shake vigorously until completely dissolved.

08:15:42 AM

Zero

• Press Timer and the display will show the countdown prior adding H196770C-0 Silica HR Reagent C or wait 10 minutes.

mg/L Silica HR (SiO₂) **Reaction time**

10min

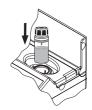
- Add one packet of H196770C-0 Silica HR Reagent C. Replace the plastic stopper and the cap. Shake vigorously until completely dissolved.
- Insert the cuvette into the holder and close the lid.



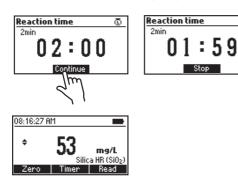


۵.

58

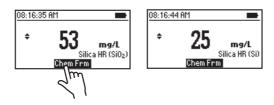


 Press Continue and the display will show the countdown prior to the measurement or wait 2 minutes and press Read. When the timer ends the meter will perform the reading. The instrument displays the results in mg/L silica (SiO₂).



© 08:16:20 AM ∑ ■ READ Silica HR (SiO₂)

- Press the \blacktriangle or \blacktriangledown key to access the second level functions.
- Press Chem Frm to convert the result to mg/L of silicon (Si).



- Press the \blacktriangle or \blacktriangledown key to return to the measurement screen.

INTERFERENCES

Interference may be caused by:

- Phosphate above 75 mg/L, causes an 11% reduction in reading
- Phosphate above 60 mg/L, causes a 2% reduction in reading
- Sulfide and high concentration of iron
- Eliminate color and turbidity interferences by zeroing the meter with the original water sample

10.32. ZINC

SPECIFICATIONS

Range	0.00 to 3.00 mg/L (as Zn)
Resolution	0.01 mg/L
Accuracy	\pm 0.03 mg/L \pm 3% of reading at 25 °C
Light Source	LED with narrow band interference filter @ 575 nm
Method	Adaptation of Standard Methods for the Examination of Water and Wastewater,
	18 th Edition, Zincon Method

REQUIRED REAGENT

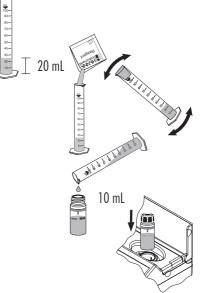
Code	Description	Quantity
HI93731A-0	Zinc Reagent A	1 packet
HI93731B-0	Zinc Reagent B	0.5 mL

REAGENT SETS

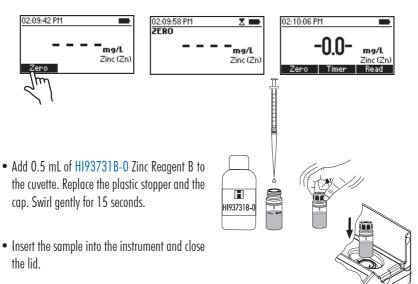
HI93731-01	Reagents for 100 tests
HI93731-03	Reagents for 300 tests
For other accessori	es see ACCESSORIES section.

MEASUREMENT PROCEDURE

- Select the Zinc method using the procedure described in the METHOD SELECTION section.
- Fill the graduated glass vial up to the 20 mL mark with the sample.
- Add one packet of HI93731A-0 Zinc Reagent A, close the cylinder. Invert several times to mix until completely dissolved.
- Fill a cuvette with 10 mL of the reacted sample (up to the mark). Replace the plastic stopper and the cap.
- Insert the cuvette into the holder and close the lid.



• Press Zero. The display will show "-0.0-" when the meter is zeroed and ready for measurement.



• Press **Timer** and the display will show the countdown prior to the measurement or wait 3 minutes and 30 seconds and press **Read**. When the timer ends the meter will perform the reading. The instrument displays the results in **mg/L** of **zinc (Zn)**.



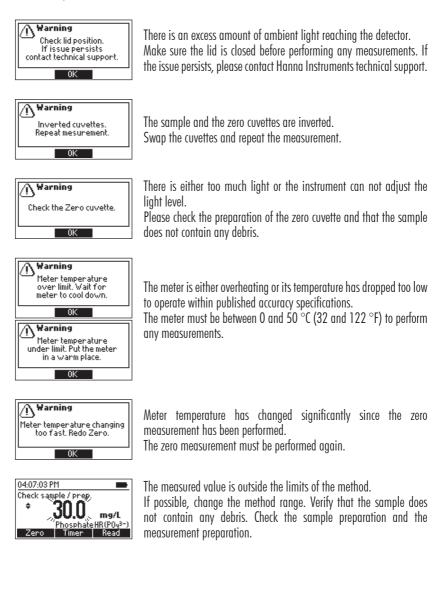
INTERFERENCES

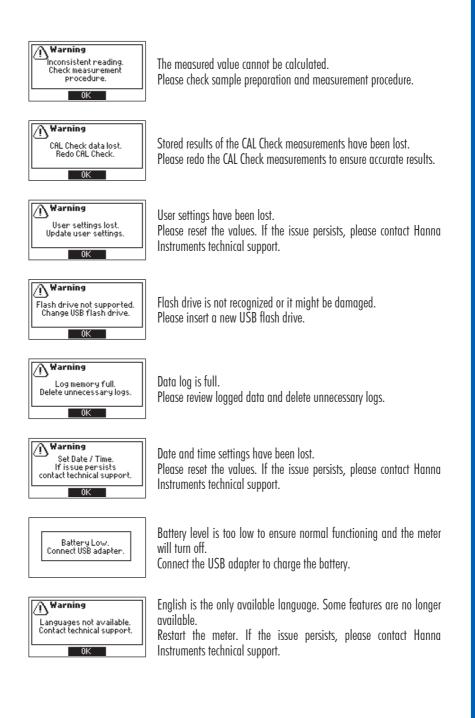
Interference may be caused by:

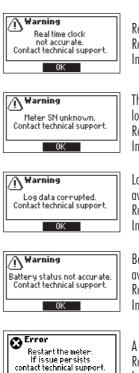
- Iron above 7 mg/L
- Aluminum above 6 mg/L
- Copper, Manganese, Nickel above 5 mg/L
- Cadmium above 0.5 mg/L

11. WARNINGS & ERRORS

The instrument shows clear warning messages when erroneous conditions appear and when measured values are outside the expected range. The information below provides an explanation of the errors and warnings, and recommended action to be taken.







Real time clock is not accurate. Some features are no longer available. Restart the meter. If the issue persists, please contact Hanna Instruments technical support.

The device serial number can not be identified. Some features are no longer available.

Restart the meter. If the issue persists, please contact Hanna Instruments technical support.

Logged data is no longer accessible. Some features are no longer available.

Restart the meter. If the issue persists, please contact Hanna Instruments technical support.

Battery charge level is not accurate. Some features are no longer available.

Restart the meter. If the issue persists, please contact Hanna Instruments technical support.

A critical error has occurred.

Restart the meter. If the issue persists, please contact Hanna Instruments technical support.

12. STANDARD METHODS

Description	Range	Method
Aluminum	0.00 to 1.00 mg/L (as Al ³⁺)	Aluminon
Ammonia LR	0.00 to 3.00 mg/L (as NH ₃ -N)	Nessler
Ammonia MR	0.00 to 10.00 mg/L (as NH ₃ -N)	Nessler
Ammonia HR	0.0 to 100.0 mg/L (as NH ₃ -N)	Nessler
Bromine	0.00 to 8.00 mg/L (as Br ₂)	DPD
Chlorine Dioxide	0.00 to 2.00 mg/L (as ClO ₂)	Chlorophenol Red
Chlorine Dioxide, Rapid Method	0.00 to 2.00 mg/L (as ClO ₂)	DPD
Chlorine, Free	0.00 to 5.00 mg/L (as Cl ₂)	DPD
Chlorine, Total	0.00 to 5.00 mg/L (as Cl ₂)	DPD
Chromium(VI) LR	0 to 300 μ g/L (as Cr (VI))	Diphenylcarbohydrazide
Chromium(VI) HR	0 to 1000 µg/L (as Cr(VI))	Diphenylcarbohydrazide
Copper LR	0.000 to 1.500 mg/L (as Cu ²⁺)	Bicinchoninate
Copper HR	0.00 to 5.00 mg/L (as Cu ²⁺)	Bicinchoninate
Hydrazine	0 to 400 µg/L (as N2H4)	p-Dimethylaminobenzaldehyde
Iron LR	0.000 to 1.600 mg/L (as Fe)	TPTZ
Iron HR	0.00 to 5.00 mg/L (as Fe)	Phenanthroline
Iron(II)	0.00 to 6.00 mg/L (as Fe ²⁺)	EPA 315B
Molybdenum	0.0 to 40.0 mg/L (as Mo ⁶⁺)	Mercaptoacetic Acid
Nitrate	0.0 to 30.0 mg/L (as NO ₃ ⁻ - N)	Cadmium reduction
Nitrite LR	0 to 600 μ g/L (as NO $_2^-$ -N)	Diazotization
Nitrite HR	0 to 150 mg/L (as NO_2^{-1})	Ferrous Sulfate
Oxygen, Dissolved	0.0 to 10.0 mg/L (as 0_2)	Winkler
Oxygen Scavengers (Carbohydrazide)	0.00-1.50mg/L (as Carbohydrazide)	Iron Reduction
Oxygen Scavengers (DEHA)	0 to 1000 μ g/L (as DEHA)	Iron Reduction
Oxygen Scavengers (Hydroquinone)	0.00-2.50mg/L (as Hydroquinone)	Iron Reduction
Oxygen Scavengers (Iso-Ascorbic Acid)	0.00-4.50mg/L (as Iso-Ascorbic Acid)	Iron Reduction
рН	6.5 to 8.5 pH	Phenol Red

Description	Range	Method
Phosphate LR	0.00 to 2.50 mg/L (as PO ₄ ³⁻)	Ascorbic Acid
Phosphate HR	0.0 to 30.0 mg/L (as PO ₄ ³⁻)	Amino Acid
Silica LR	0.00 to 2.00 mg/L (as SiO ₂)	Heteropoly Blue
Silica HR	0 to 200 mg/L (as SiO_2)	EPA
Zinc	0.00 to 3.00 mg/L (as Zn)	Zincon

13. ACCESSORIES

13.1. REAGENT SETS

Code	Description
HI93700-01	100 ammonia LR tests
HI93700-03	300 ammonia LR tests
HI93701-01	100 chlorine free tests (powder)
HI93701-03	300 chlorine free tests (powder)
HI93701-F	300 chlorine free tests (liquid)
HI93701-T	300 chlorine total tests (liquid)
HI93702-01	100 copper HR tests
HI93702-03	300 copper HR tests
HI93704-01	100 hydrazine tests
HI93704-03	300 hydrazine tests
HI93705-01	100 silica LR tests
HI93705-03	300 silica LR tests
HI93707-01	100 nitrite LR tests
HI93707-03	300 nitrite LR tests
HI93708-01	100 nitrite HR tests
HI93708-03	300 nitrite HR tests
HI93710-01	100 pH tests
HI93710-03	300 pH tests
HI93712-01	100 aluminum tests
HI93712-03	300 aluminum tests
HI93713-01	100 phosphate LR tests
HI93713-03	300 phosphate LR tests
HI93715-01	100 ammonia MR tests
HI93715-03	300 ammonia MR tests
-	

Code	Description
HI93716-01	100 bromine tests
HI93716-03	300 bromine tests
HI93717-01	100 phosphate HR tests
HI93717-03	300 phosphate HR tests
HI93721-01	100 iron HR tests
HI93721-03	300 iron HR tests
HI93723-01	100 chromium(VI) HR tests
HI93723-03	300 chromium(VI) HR tests
HI93728-01	100 nitrate tests
HI93728-03	300 nitrate tests
HI93730-01	100 molybdenum tests
HI93730-03	300 molybdenum tests
HI93732-01	100 dissolved oxygen tests
HI93732-03	300 dissolved oxygen tests
HI93731-01	100 zinc tests
HI93731-03	300 zinc tests
HI93733-01	100 ammonia HR tests
HI93733-03	300 ammonia HR tests
HI93738-01	100 chlorine dioxide tests
HI93738-03	300 chlorine dioxide tests
HI93746-01	50 iron LR tests
HI93746-03	150 iron LR tests
HI93749-01	100 chromium(VI) LR tests
HI93749-03	300 chromium(VI) LR tests
HI95747-01	100 copper LR tests
HI95747-03	300 copper LR tests
HI96770-01	100 silica HR tests

ACCESSORIES

Code	Description
HI96770-03	300 silica HR tests
HI96773-01	50 oxygen scavengers tests
HI96773-03	150 oxygen scavengers tests
HI96776-01	100 iron(II) tests
HI96776-03	300 iron(II) tests
HI96779-01	100 chlorine dioxide (rapid) tests
HI96779-03	300 chlorine dioxide (rapid) tests

13.2. pH ELECTRODES

Code	Description
HI10530	Triple ceramic, double junction, low temperature glass, refillable pH electrode with conical tip and temperature sensor
HI10430	Triple ceramic, double junction, high temperature glass, refillable pH electrode with temperature sensor
HI11310	Glass body, double junction, refillable pH/temperature electrode
HI11311	Glass body, double junction, refillable pH/temperature electrode with enhanced diagnostics
HI12300	Plastic body, double junction, gel filled, non refillable pH/temperature electrode
HI12301	Plastic body, double junction, gel filled, non refillable pH/temperature electrode with enhanced diagnostics
HI10480	Glass body, double junction with temperature sensor for wine analysis
FC2320	Double junction, open reference, non refillable, electrolyte viscolene, PVDF body with conical tip, pH/temperature electrode
FC2100	Double junction, open reference, non refillable, electrolyte viscolene, glass body with conical tip, pH/temperature electrode
FC2020	Double junction, open reference, non refillable, electrolyte viscolene, PVDF body with conical tip, pH/temperature electrode

Note: The enhanced diagnostics information are not displayed by meter.

13.3. pH SOLUTIONS

BUFFER SOLUTIONS

Code	Description
HI70004P	pH 4.01 buffer sachet, 20 mL (25 pcs.)
HI70007P	pH 7.01 buffer sachet, 20 mL (25 pcs.)
HI70010P	pH 10.01 buffer sachet, 20 mL (25 pcs.)
HI7001L	pH 1.68 buffer solution, 500 mL
HI7004L	pH 4.01 buffer solution, 500 mL
HI7006L	pH 6.86 buffer solution, 500 mL
HI7007L	pH 7.01 buffer solution, 500 mL
HI7009L	pH 9.18 buffer solution, 500 mL
HI7010L	pH 10.01 buffer solution, 500 mL
HI8004L	pH 4.01 buffer solution in FDA approved bottle, 500 mL
HI8006L	pH 6.86 buffer solution in FDA approved bottle, 500 mL
HI8007L	pH 7.01 buffer solution in FDA approved bottle, 500 mL
HI8009L	pH 9.18 buffer solution in FDA approved bottle, 500 mL
HI8010L	pH 10.01 buffer solution in FDA approved bottle, 500 mL

ELECTRODE STORAGE SOLUTIONS

Code	Description
HI70300L	Storage solution, 500 mL
HI80300L	Storage solution in FDA approved bottle, 500 mL

ELECTRODE CLEANING SOLUTIONS

Code	Description
HI70000P	Electrode rinse sachet, 20 mL (25 pcs.)
HI7061L	General cleaning solution, 500 mL
HI7073L	Protein cleaning solution, 500 mL
HI7074L	Inorganic cleaning solution, 500 mL
HI7077L	Oil & fat cleaning solution, 500 mL
HI8061L	General cleaning solution in FDA approved bottle, 500 mL
HI8073L	Protein cleaning solution in FDA approved bottle, 500 mL
HI8077L	Oil & fat cleaning solution in FDA approved bottle, 500 mL

ELECTRODE REFILL ELECTROLYTE SOLUTIONS

Code	Description
HI7082	3.5M KCl electrolyte, 4x30 mL, for double junction electrodes
HI8082	3.5M KCl electrolyte in FDA approved bottle, 4x30 mL, for double junction

ACCESSORIES

13.4. OTHER ACCESSORIES

Code	Description
HI72083300	carrying case
HI731318	cloth for wiping cuvettes (4 pcs.)
HI731331	glass cuvette (4 pcs.)
HI731335N	cap for cuvette (4 pcs.)
HI731340	200 μ L automatic pipette
HI731341	1000 μ L automatic pipette
HI731342	2000 μ L automatic pipette
HI740034P	cap for 100 mL beaker (10 pcs.)
HI740036P	100 mL plastic beaker (10 pcs.)
HI740038	60 mL glass bottle and stopper
HI740142P	1 mL graduated syringe (10 pcs)
HI740143	1 mL graduated syringe (6 pcs.)
HI740144	pipette tip (6 pcs.)
HI740157P	plastic refilling pipette (20 pcs.)
HI740220	25 mL graduated glass vial (2 pcs.)
HI740223	170 mL plastic beaker
HI740224	170 mL plastic beaker (12 pcs.)
HI740225	60 mL graduated syringe
HI740226	5 mL graduated syringe
HI740227	filter assembly
HI740228	filter disc (25 pcs.)
HI740229	100 mL graduated cylinder
DEMI-02	demineralizer

Description
USB power adapter, European plug
USB power adapter, USA plug
electrode holder
CAL Check cuvette kit for HI83305
Sample preparation kit consisting of activated carbon for 50 tests, demineralizer bottle for 10 L of water, 100 mL graduated beaker with cap, 170 mL graduated beaker with cap, 3 mL pipette, 60 mL syringe, 5 mL syringe, graduated cylinder, spoon, funnel, filtter paper (25 pcs.)
USB to micro USB cable connector
cuvette cleaning solution (230 mL)
activated carbon (50 pcs.)

CERTIFICATION

All Hanna Instruments conform to the CE European Directives.



Disposal of Electrical & Electronic Equipment. The product should not be treated as household waste. Instead hand it over to the appropriate collection point for the recycling of electrical and electronic equipment which will conserve natural resources.

Disposal of waste batteries. This product contains batteries, do not dispose of them with other household waste. Hand them over to the appropriate collection point for recycling.

Ensuring proper product and battery disposal prevents potential negative consequences for the environment and human health. For more information, contact your city, your local household waste disposal service, the place of purchase or go to www.hannainst.com.



RECOMMENDATIONS FOR USERS

Before using this product, make sure it is entirely suitable for your specific application and for the environment in which it is used. Any variation introduced by the user to the supplied equipment may degrade the photometer's performance. For yours and the meter's safety do not use or store the photometer in hazardous environments.

WARRANTY

The HI83305 is warranted for two years against defects in workmanship and materials when used for its intended purpose and maintained according to instructions. This warranty is limited to repair or replacement free of charge. Damage due to accidents, misuse, tampering or lack of prescribed maintenance is not covered.

If service is required, contact your local Hanna Instruments Office. If under warranty, report the model number, date of purchase, serial number (engraved on the bottom of the meter) and the nature of the problem. If the repair is not covered by the warranty, you will be notified of the charges incurred. If the instrument is to be returned to Hanna Instruments, first obtain a Returned Goods Authorization (RGA) number from the Technical Service department and then send it with shipping costs prepaid. When shipping any instrument, make sure it is properly packed for complete protection.

Hanna Instruments reserves the right to modify the design, construction or appearance of its products without advance notice.

World Headquarters

Hanna Instruments Inc. Highland Industrial Park 584 Park East Drive Woonsocket, RI 02895 USA www.hannainst.com



MAN83305