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IMPORTANT NOTES FOR SULFIDE ISE SYSTEM

Calibration and Cleaning of Sulfide Selective Ion Measurement System For Waste Water Sulfide Analysis

All calibration solutions and Process Grab Sample should be calibrated and tested at identical temperatures to the process temperature. The actual temperature of the process solution (and thereby the calibrating solutions as well) is not as critical as the fact that they are calibrated at the same temperature. For greatest overall accuracy of the Sulfide (S2-) measurement, however, measurement should be performed as close to 25 degrees Celsius (room temperature process solutions) as possible.

Calibration Point 1 (always the lower concentration value) and Calibration Point 2 (always the higher concentration value) determines the response of the 54e Analyzer to a given Sulfide Ion Selective Sensor (AB 6120). The calibration solutions values for Calibration Point 1 is the low ppm value and Calibration Point 2 is high ppm value

The One Point Re-Calibration Standard (usually a process grab sample or one of the two Sulfide standard solution) is always performed by using the standardize option from the Calibrate Main Menu. A one point calibration should never be performed by using the 2-point calibration option from the Calibrate Main Menu.

At all times the calibration solutions should be kept clean and out of direct sunlight and/or other high-energy radiation sources. New sensors should be conditioned in Sulfide standard solution for 3 - 5 minutes before beginning calibration procedure as described in this quick calibration guide.

The acceptable pH range of the AB 6120, AB 6120A & AB 8120 Sulfide Ion Selective Sensor is 11 - 14. pH compensation is required for all pH values to obtain a meaningful (pH independent) total Sulfide analytical data.

To enter custom ion menu, hold F3 for 5-7 seconds, then enter secret code of 20000 into system.

The following steps are required for an installation of a new Sulfide sensor:

- 1) Enter the nominal ISO Voltage -145 mV and Slope (- 28.00 mV per decade) as given on the proceeding page.
- 2) Perform a two point calibration to empirically determine slope. See attached procedures for 2-point calibration.
- 3) Perform a one-point calibration (standardize) at the defined (expected) measurement value. See the proceeding section one point calibration for further details on use of this calibration function.
- 4) Place sensor into process and allow it to find electrochemical and thermal equilibrium.
- 5) To account for any differences between the ionic strength and interfering ions between calibration and measured solution, a grab sample should be taken and analyzed by an alternate analysis system, and the online Sulfide system adjusted to read the analyzed value. The sensor should be left in process and this grab sample calibration performed as may be required, unless the sensor seems to be losing sensitivity or giving erratic readings.

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INSTRUCTIONS FOR CONFIGURING 1056 ISE ANALYZER for Sulfide Measurements

For any 1056 ISE Analyzer provided by ASTI the ISE channel MUST be the first channel (Sensor 1). If you have a single channel ISE analyzer (1056-XX-22ISE-38-YY) this allows for only one option, but you should be aware for any future installation that are dual channel ISE/pH analyzers (1056-XX-22ISE-33-YY) that the ISE sensor should ALWAYS be hooked into channel 1. You should check that your ASTI ISE sensor is properly wired according to the official ASTI hook-up schematic for ASTI sensors with preamplifier or without preamplifiers to the 1056 analyzer: http://www.astisensor.com/Rosemount 1056 With Preamp Hookup.pdf http://www.astisensor.com/Rosemount 1056 No Preamp Hookup.pdf

The 1056 analyzers will support both ASTI ISE sensors with and without preamplifiers. The ISE channel MUST ALWAYS be set to Custom ISE (If your analyzer shows the measurement for Sensor 1 as anything OTHER THAN Custom ISE <u>IT IS IMPROPERLY CONFIGURED</u>). If your analyzer has a setting of anything other than Custom ISE, it has undoubtedly been improperly reconfigured after the ASTI factory configuration and testing. Here is how to restore it:

Main Menu -> Program -> Measurement -> Sensor 1 -> Custom ISE Setup ->{{ENTER SLOPE, FORM. WT., ISO PCON, & ISO VOLTAGE}}

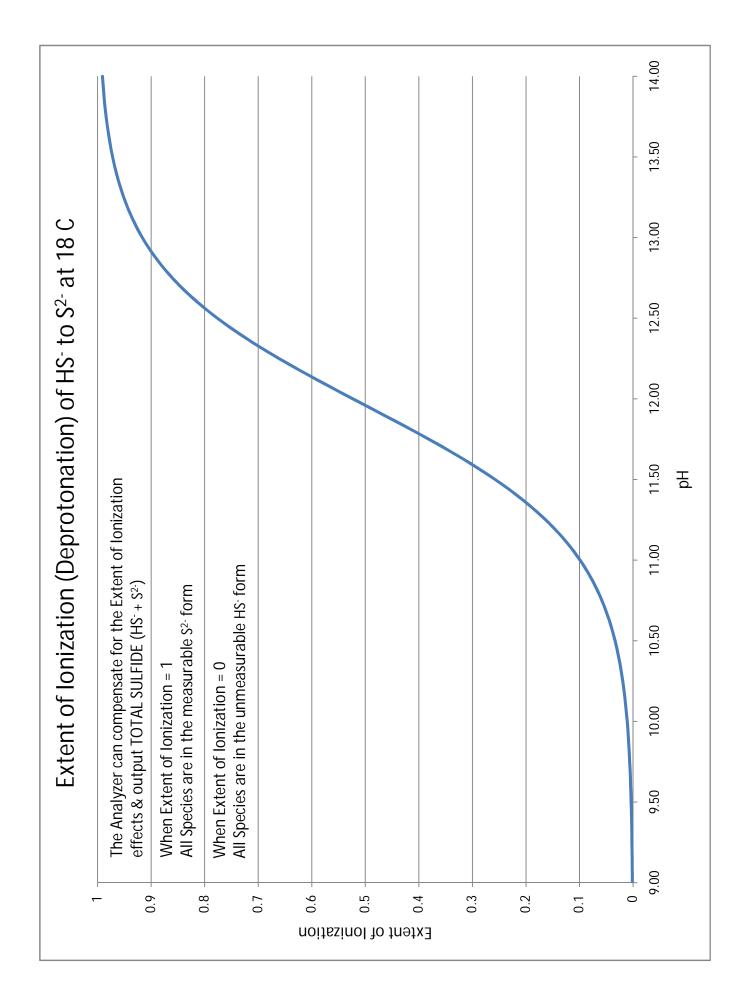
The values for these four Custom ISE variables are provided below. Normally the user would not need to enter these values as all ASTI provided 1056 ISE analyzer are preconfigured and tested with the ISE sensor provided at the factory. Only an alteration of the factory configuration would necessitate a restoration of ASTI factory configuration (NOT THE SAME AS RESTORE TO ROSEMOUNT FACTORY DEFAULT!!!)

Custom ISE Variable	Description of Variable	NOTES
32.07 grams per mol	IONIC WEIGHT	Defined by Selective Ion Measurement – DO NOT MODIFY
	Form Wt. in the 1056 analyzer terminology	
3.50515	ISOPOTENTIAL CONCENTRATION	Factory Defined – DO NOT MODIFY
	Iso pCon in the 1056 analyzer terminology	
-28.00 mV per decade	DEFAULT SLOPE	Will be changed when 2-point calibration is performed
	Slope in the 1056 terminology	
-145 mV	ISOPOTENTIAL VOLTAGE	Will be changed when either a 2-point slope or 1-point offset
	Iso Voltage in the 1056 analyzer terminology	standardize calibration is performed

Please note that after restoring the analyzer to the proper Custom ISE configuration (as described above), you will need to repeat your 2-point calibration using calibration solutions that are one decade (10X) apart in value. In addition, you will need to once again place the ISE sensor back into service and allow it to equilibrate. You will then also need to repeat your 1-point grab sample calibration (standardize in the 1056 terminology). This means taking a sample from the process and determining the ISE concentration (in ppm) for the grab sample and then using the 1-point standardize calibration feature of the 1056 analyzer to bring the online reading in accordance with the laboratory gram sample determination. In addition, once you reset your analyzer with the correct Custom ISE configuration, you can choose to run the temperature compensation in the automatic mode (this is the default) or in the manual temperature compensation mode. The control of the temperature compensation settings are identical for the Custom ISE as for pH namely in this particular case:

Main Menu -> Temperature -> Sensor1 -> {{ Set for AUTO or MANUAL }}

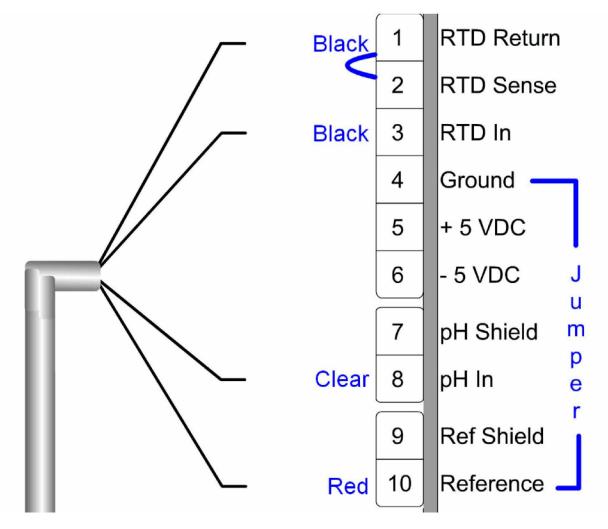
Please also note that the activity of free Sulfide ion in solutions is pH dependent over some pH ranges. The extent of ionization (HS- conversion to the measurable S2- ion form) is a pH and temperature dependent process. The following page describes such dependence for your consideration.





Connection Diagram of Iotron[™] pH / ORP / ISE Sensors **Without** Preamplifiers to Rosemount 1056 pH / ORP / ISE Analyzers

Connection from lotron[™] Sensor to Terminal Block in Rosemount Transmitter



Note 1: The temperature compensation element is 100 or 1000 Ohm Platinum (autoswitched).

Note 2: For ORP and Ion Selective Sensors, please put the active signal (clear) to terminal 8 (indicated as pH In).

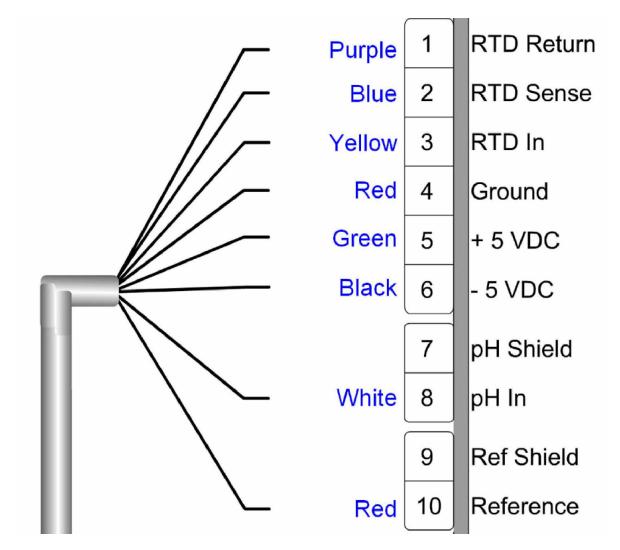
Note 3: Terminals 4 & 10 and terminals 1 & 2 must be tied together to satisfy the analyzer input requirements and disable the reference diagnostic features (pH glass diagnostics should still be available).

Note 4: For Dual Channel Analyzers, please ensure that the proper type of sensor is connected to the proper input board.



Connection Diagram of Iotron[™] pH / ORP / ISE Sensors **With** Preamplifiers to Rosemount 1056 pH / ORP / ISE Analyzers

Connection from lotron[™] Sensor to Terminal Block in Rosemount Transmitter



Note 1: The temperature compensation element is 100 or 1000 Ohm Platinum (autoswitched).

Note 2: The preamplifier does not support diagnostic features (if any).

Note 3: For ORP or Ion Selective Sensors, please put the active signal (white) to terminal 8 (indicated as pH In).

Note 4: For Dual Channel Analyzers, please ensure that the proper type of sensor is connected to the proper input board.

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MENU for ROSEMOUNT 54e SULFIDE ANALYZER

Main Sub-Menu	Calibrate Main Menu	
Calibrate Sub-Menu	2-Point Calibration	
low ppm	Calibration Point 1	
high ppm	Calibration Point 2	
Enabled	Hold Option for Outputs 1 & 2: Read Rosemount 54e-pH/ORP Manual for Details	
Calibrate Sub-Menu	Standardize	
Variable	Standardize Standardize Concentration: Value = Grab Sample, Calibration Solution 1 or 2:	
, and the second s	See calibration instructions for details on this function; a.k.a. 1-point calibration	
Calibrate Sub-Menu	Adjust Temp	
User Defined	Manual Adjustment of temperature, not required for most installations	
Calibrate Sub-Menu	Temp Compensation	
Setting / Value	FUNCTION / PARAMETER	
Manual 25°C	Temperature Compensation (Factory Defined – DO NOT MODIFY)	
M ' C 1 14		
Main Sub-Menu	Diagnostic Variables	
Defined by 54e	Displays all currently defined variables in 54e Memory	
Main Sub-Menu	Program	
Program Sub-Menu	Alarm Setpoints	
Rosemount Default	Variable alarm setpoints for 3 @ relays	
0.000 ppm & 999.9 ppm	Any user defined value within measurement output range is acceptable	
Program Sub-Menu	Output Setpoints	
4mA= 0.000 ppm	4-20mA Output 1 – Process Sulfide ISE	
20mA = 10 ppm	These are only the factory defaults. These values can be altered.	
90 seconds	Time Average of Process Output (Dampening)	
$4mA = 5^{\circ}C$	4-20mA Output 2 – Temperature	
20mA = 50 °C	These are only the factory defaults. These values can be altered.	
0 seconds	Time Average of Temperature Output	
Program Sub-Menu	Diagnostics	
Disabled	Disabled diagnostics required to make Sulfide ISE measurement run properly	
	(Factory Defined – DO NOT MODIFY)	
Program Sub-Menu	Simulated Test	
Rosemount Default	Read Rosemount 54e-pH/ORP Manual for Details	
Program Sub-Menu	Configure	
ASTI Programmed	No Passcode has been assigned.	
Passcode for Configure	Various parameters in this function may affect output and display of Analyzer	
S2-	Selective Ion Measurement Parameter (Factory Defined – DO NOT MODIFY)	
ppm	Display units for S2- Measurement (Defined in Program-Configure Menu)	
°C	Display units for Temperature Measurement (Defined in Program-Configure Menu)	
-999 to + 999	ISE Input Range in mV's – Rosemount Defined	
-777 10 + 777	ISE input Kange in inv 5 – Kosemount Denned	
Custom Ion Curro	Press F3 for 5-7 seconds, enter Passcode of 20000	
Custom Ion Curve		
32.07 grams per mol 3.50515	IONIC WEIGHT (Defined by Selective Ion Measurement – DO NOT MODIFY)	
	pISOPOTENTIAL CONCENTRATION (Factory Defined – DO NOT MODIFY)	
-28.00 mV per decade	DEFAULT SLOPE (Will be changed when 2-point calibration is performed)	
-145 mV	ISOPOTENTIAL VOLTAGE (Will be changed when 2-point calibration is performed)	

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Sulfide (S2-) Probe Two Point Calibration

This calibration method should not need to be performed frequently. All new Sulfide sensors should be calibrated using a 2-point calibration first. Subsequent calibrations can be made using only the standardize option from the Calibrate Main Menu. This calibration determines the sensitivity or slope of each sensor, which is then stored in the 54e transmitter. A "standardize" (a.k.a. 1-point) calibration must be performed after every 2-point calibration. Details for the 1-point calibration are given in the proceeding page. MAKE SURE THAT THE "HOLD" FEATURE IS ON BEFORE STARTING ANY CALIBRATION. READ THE ROSEMOUNT 54E-PH/ORP MANUAL FOR FURTHER DETAILS ABOUT THE HOLD FEATURE.

Set-up requirements:

- Two 250 mL GLASS OR PLASTIC BEAKERS.
- Low Sulfide Standard Solution (low ppm)
- High Sulfide Standard Solution (high ppm)

Follow the on-screen directions in 2-point calibration submenu in the 54e Analyzer. Calibration Points 1 & 2 have been preprogrammed into your instrument. The low standard solution (low ppm) will always be Calibration Point 1 & the high standard solution (high ppm) will always be Calibration Point 2. Do not change the preprogrammed values for Calibration Point 1 & 2.

Important Notes about Calibration:

- Fill a 250 mL GLASS beaker with enough standardization solution such that the entire tip of the Sulfide sensor will be submersed
- Please read the attached sheet on the cleaning procedure for this ion selective sensor. The cleaning procedure should usually be performed when transferring a sensor in or out of solution. Sensors can also be cleaned before being placed into different concentration standardization solutions and/or grab sample solutions. Cleaning is only required if fouling appears on the ISE membrane or on the reference junction. If no contamination is apparent, then the simple rinsing procedure given below can be used.
- Thoroughly rinse the sensors with DI water and gently blot dry with a clean paper towel. Be careful not to scratch or damage the sensitive solid state Sulfide ion selective membrane
- Allow a minimum of 3 5 minutes for the sensor to stabilize once it has been removed from the process and placed into the low standard solution (low ppm).
- Allow a minimum of 3 5 minutes for the sensor to stabilize between the low and high Calibration Solutions (low ppm to high ppm).

Key Sequence for 2-Point Calibration

- 1. Press any Function Key
- 2. Highlight Calibrate and Enter Function
- 3. Select Hold Mode to On and Continue
- 4. Select 2-Point Calibration and Enter Function
- 5. Sulfide sensor should be in low ppm standard solution already cleaned and conditioned. Press Continue. Instrument will display that Calibration Point 1 is stabilizing. The instrument will take about 20 seconds to stabilize and determine mV value for the first calibration point.
- 6. After Cal Point 1 has stabilized select edit and input low ppm into instrument. This value must be entered even if it already is correct on the display. After low ppm has been entered into Analyzer, press save to continue to second calibration point.
- 7. Sulfide sensor should be in high ppm standard solution already cleaned and conditioned. Press Continue. Instrument will display that Calibration Point 2 is stabilizing. The instrument will take about 20 seconds to stabilize and determine mV value for the second calibration point.
- 8. After Cal Point 2 has stabilized select edit and input high ppm into instrument. **This value must be entered even if it already is correct on the display.** After high ppm has been entered into Analyzer, press save.

THE ROSEMOUNT ANALYZER AT THIS POINT SHOULD DISPLAY "2-POINT CALIBRATION DONE". ONLY IF THIS MESSAGE APPEARS HAS THE TWO-POINT CALIBRATION BEEN SUCCESFULLY PERFORMED, OTHERWISE REPEATED STEPS AS OUTLINED ABOVE UNTIL THE ANALYZER ACCEPTS THE TWO-POINT CALIBRATION.

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Single Point Calibration -

This is the correct method to Adjust for Sensor Drift

Sulfide Sensor One Point Calibration

- Only use the "Standardize" function from the Calibrate Menu. Any other method will destroy the previous 2-point calibration
- Can be performed DAILY if required.

When the Sulfide system has been calibrated by the 2-point method it can then be used in the "single point" calibration mode (on a daily basis if required). MAKE SURE THAT THE "HOLD" FEATURE IS ON BEFORE STARTING ANY CALIBRATION. READ THE ROSEMOUNT 54E-PH/ORP MANUAL FOR FURTHER DETAILS ABOUT THE HOLD FEATURE.

Set-up requirements:

- Two 250 mL GLASS OR PLASTIC BEAKERS
- Low Sulfide Standard Solution (low ppm) or
- High Sulfide Standard Solution (high ppm) or
- Process Grab Sample Solution

Follow the on-screen directions in the "Standardize" submenu of the Calibration Menu in the 54e Analyzer. No value has been preprogrammed into your instrument for the standardization concentration. Both the low and high standardization solutions can be used to perform the "1-point" calibration performed by the standardize function. In addition, grab samples can be taken from the process and analyzed by an alternate method for ion concentration. The separately determined concentration of the process sample can then be entered into the standardize menu. In this way, the Sulfide sensor can be standardized without ever having to remove the sensor from the process line.

Important Notes about 1-point "Standardize" Calibration:

- Fill a 250 mL GLASS beaker with enough standardization solution such that the entire tip of the Sulfide sensor will be submersed
- Read the attached sheet on the cleaning procedure for this ion selective sensor. The cleaning procedure should usually be performed when transferring a sensor in or out of solution. Sensors can also be cleaned before being placed into different concentration standardization solutions and/or grab sample solutions. Cleaning is only required if fouling appears on the ISE membrane or on the reference junction. If no contamination is apparent, then the simple rinsing procedure given below can be used.
- Thoroughly rinse the sensors with DI water and gently blot dry with a clean paper towel. Be careful not to scratch or damage the sensitive solid state Sulfide ion selective membrane
- Allow a minimum of 3 5 minutes for the sensor to stabilize once it has been removed from the tank and placed into either the low or high standard solution.

Key Sequence for 1-Point Calibration

- 1. Press any Function Key
- 2. Highlight Calibrate and Enter Function
- 3. Select Hold Mode to On and Continue
- 4. Select Standardize and Enter Function
- 5. Sulfide sensor should be in low ppm standard solution already cleaned and conditioned. Allow reading to stabilize. Press Edit. Enter low ppm into Analyzer and press save. The one point calibration is now complete. The reading on the display should be same as the entered standardize value. Alternatively, the high ppm calibration solution can be used for 1-point calibration if this is expected measurement value. If a grab sample is used, allow the sensor to find a stable reading in process. Enter the analyzed value of the process solution for the Sulfide concentration. The sensor should be left in service while this standardize calibration is performed.

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Cleaning and Maintenance of ASTI "S2-" Probe

Before a major 2-point calibration the sensor may need to be cleaned each time. The frequency of cleaning will depend on the quality of the process water and the build up of process reagents on the probe tip.

Note:

Any noticeable deposits on the tip of the sensor will result in a less accurate calibration and measurement.

CLEANING:

- 1. Thoroughly rinse the sensor tip with DI water. Gently blot the sensor tip dry.
- 2. Scrape the entire Kynar reference area clean with a sharp blade or Stanley knife. This reference is solid Kynar and cannot be damaged. Do <u>not</u> scratch the membrane.
- 3. Once this has been achieved the entire tip can be soaked in either the low or high standardization solution. After allowing sufficient time for conditioning; proceed to perform a 2-point or 1-point calibration.
- 4. The standardization solutions can always be used as conditioning solution for extended storage. Do not allow sensor to be exposed to air for prolonged periods of time. Always store sensor in standardization solution when not in service in process. For long-term storage, a standard solution should be placed into sensor protective cap. The cap should be sealed onto sensor body with Teflon tape.
- 5. The Sulfide ion selective electrode tip may become insensitive to measuring Sulfide ions due to an exposure to an interfering ion, mechanical damage or exposure to high levels of Sulfide. In this case, please follow the direction stated in the following pages which are extracted from the technical document Cleaning_Polishing_Solid-State_ISE_Procedure.pdf as posted on our technical support webpages.

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Miscellaneous

The decimal place can be moved in any screen of the 54e Analyzer by placing the cursor over the decimal place and using the up and down arrows to move the decimal point to any position.

Do not to allow air bubbles to get trapped near the Sulfide ion selective solid state membrane. This is will cause erroneous readings and drift.

The Sulfide sensor is comprised of a high-impedance solid state membrane system. Care should be taken not to move or touch the cable once a value is being stabilized. Touching the sensor cable can cause a noisy signal that may result in erroneous values and calibrations.

Please see the hook-up schematic found in the AB 6120, AB 6120A & AB 8120 Sulfide sensor shipping box, together with the specification sheet. Be sure to check whether that the jumper in the instrument is set for an internal preamplifier (preamplifier in analyzer setting).