



- Conductivity Sensor

(Product No. 3135PK)

The Conductivity Adaptor (Product No. 3135)

Range 1: 0 to 100 μ S Range 2: 0 to 1000 μ S Range 3: 0 to 10 mS (10,000 μ S) Range 4: 0 to 100 mS (100,000 μ S)

The Conductivity Electrode (Product No. 3136)

Built -in automatic temperature compensation Cell constant tolerance: $\pm 20\%$

🐼 DATA HARVEST

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Introduction

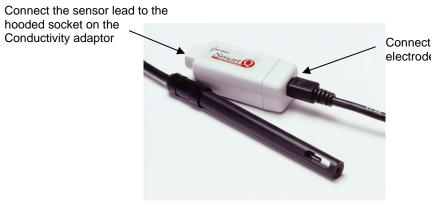
The *Smart* Q Conductivity adaptor enables a Conductivity electrode to be connected to an EasySense unit. The EasySense unit will detect when the Conductivity adaptor is connected and which of the ranges it is set to.

The *Smart Q* Conductivity sensor is used to measure the conductivity of a solution. The Conductivity electrode is a simple epoxy-bodied carbon (graphite plate) type selected for its durability in field use. It is ideal for measuring salinity and changes in conductivity of a water sample. It can be used by chemists for investigating the difference between ionic and molecular compounds in an aqueous solution. Although it will not identify the specific ions that are present, it can be used to determine the total concentration of ions in a sample.

The SI unit of conductance is Siemens per cm (S/cm). This was formerly known as a mhos (equal to S/cm) or CF units (mS/cm x 10). The siemen is a large unit, so the value for aqueous samples is commonly expressed in microsiemens (μ S).

Connecting

- Push one end of the sensor cable (supplied with the EasySense unit) into the hooded socket on the adaptor.
- Connect the Conductivity electrode (with the locating arrow facing upward) into the socket on the other end of the Conductivity adaptor.
- Connect the other end of the sensor cable to an input socket on the EasySense unit.
- The EasySense unit will detect that the Conductivity sensor is connected and display values using the currently selected range. If the range is not suitable for your investigation, set to the correct range.



Connect the Conductivity electrode to the other end



Real Time Logging Wizard

Sensor 2

Sensor 3



Ranges

All Smart Q sensors are supplied calibrated. The stored calibrations for this Sensor are the 100 uS. 1000 uS. 10 mS. and 100 mS ranges.

It is also possible for the user to adjust the calibration constants of the electrode for more accurate readings - these are stored as 'User' ranges, i.e.100 µS User, 1000 µS User, 10 mS User and 100 mS User. See 'User calibration' on page 4.

With some EasySense units it is possible to set the range from the unit, please refer to the unit's user manual.

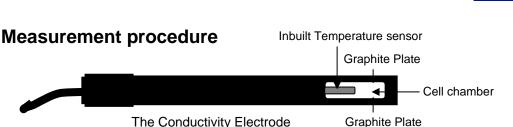
To alter the range in the EasySense software:

- 1. Select EasyLog from the Home screen.
- 2. Select the New recording wizard icon.
- 3 Click on the sensor's name.
- 4. A set sensor range window will open. Select the required range, then OK.
- Select Finish to exit the wizard. 5.

OR

- 1. From the Home screen select Sensor Config from the Settings menu.
- 2. Select the Conductivity sensor from the list and then the Change Range button.
- The current range will be highlighted. Select the required range and OK. 3.
- 4. Close Sensor Config.

The range setting will be retained until changed by the user.



The Conductivity Electrode

- If possible, soak the tip of the Conductivity electrode in deionised water for about 30 minutes to make sure the electrodes are clean. If not, rinse the tip thoroughly with deionised water before use.
- Wipe the outer part of the electrode body with a clean paper towel. Shake vigorously to remove any droplets from the cell chamber. If possible, wash the tip in a sample of the solution to be tested.
- Place the Conductivity electrode in the sample to be tested. The sample must be at least 3 cm deep to ensure the cell chamber is fully submerged.
- Stir the solution gently to get rid of any air bubbles that could be trapped in the cell chamber. • Wait for 10 seconds to allow the readings to stabilize.

Note: If you are taking readings in a solution that has a temperature below 10°C or above 35°C, allow more time for the readings to stabilize.

If the value is above, below or near to the maximum or minimum value in the current range, • alter the selected range of the Sensor.

Set Sensor Range	
Sensor 1	
Select Range:	
Conductivity	
1000uS 10mS 100mS 100uS User 1000uS User 10mS User 100mS User	
OK	

Conductivity

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• Clean thoroughly when testing is complete to avoid any contamination when the electrode is next used.

User calibration

All *Smart* Q Sensors are supplied calibrated. The stored calibrations for this Sensor are utilized when any of the default ranges are selected i.e.**100** μ **S**, **1000** μ **S**, **10 mS**, **100 mS**. These ranges are suitable for most investigations.

Conductivity is measured by putting a voltage across two electrodes (graphite plates), which have a fixed area and are a fixed distance apart. The electric current that flows between the two graphite plates varies as the conductivity of the solution changes. Each Conductivity electrode has its own physical characteristics (a small difference in the exact area and distance apart from one graphite plate to another). This is defined as the cell constant (K). See **Theory** on page 6. If you are performing an experiment that requires a very accurate calibration, it is possible for the user to adjust the calibration to the K-value of the electrode (the **User** calibration).

The accuracy of the User calibration will depend upon the condition of the electrode and the accuracy of the calibration solution. It is best to perform and save the calibration with the electrode while it is new and uncontaminated.

Only one conductivity calibration solution is required to calculate the K value for an electrode. Select a calibration solution that is closest to the estimated value of your sample solutions. An accurate calibration solution is essential. It is best to purchase prepared solutions from a supplier such as Scientific and Chemical Supplies Ltd, (web site: www.scichem.co.uk). Although recipes are available for preparing calibration solutions, it is extremely difficult to make an accurate solution.

- Ensure the Conductivity electrode is uncontaminated. Soak the tip of the electrode in deionised water for about 30 minutes. Wipe the outer body of the electrode with a clean dry paper towel. Shake vigorously to remove any droplets from the cell chamber and allow to air dry.
- Connect the Conductivity sensor to the EasySense unit.
- Start the EasySense program and select **Sensor Config** from the **Settings** menu.
- Select the Conductivity sensor from the list (it will be listed using its current range) and click on Change Range. Select a User range suitable for the value of the calibration solution e.g. for an 84 µS solution select the 100 µS User range, OK.
- Select Calibrate Sensor.
- Wash the tip of the Conductivity electrode in a sample of the Conductivity calibration solution. Place the electrode in the calibration solution, which must be at least 3 cm deep. Stir the solution gently and wait for 10 seconds to allow the readings to stabilize.
- Type the value of the calibration solution (in μS or mS) into the second calibration point box and click on Next.
 Note: 1.000 μS = 1 mS.
- The K value for the electrode will be calculated by the program e.g. K Value = 0.99876. This value can either be applied to the other User ranges (click on Yes) or just to the selected range (click on No).
- A message to say 'The User range has been reprogrammed' will be displayed. Disconnect the Conductivity sensor from the EasySense unit and then reconnect for the calibration to be applied.

The User calibration values will be stored in the Sensor and will be retained until reconfigured by the user.

First	0.0	mS	1424
Second	▶ 100.0	mS	
Instructions			
			N 1 - 1
Put the ser	nsor in to the soli	ution and pres	sNext
			Next
Current Read	lina usina		
Default Calib			0.0
Current User	0.0		
			1312



Practical information

Some typical conductivity ranges of hydrous solutions are:

Sample	Conductivity (µS/cm)	Sample	Conductivity (µS/cm)
Ultra pure water	0.055	Brackish water	1000 - 8000
Distilled water	0.5 - 5	KCI 0.01M	1,410
Rain water	20 -100	MgSO4	5,810
Drinking water	50 - 200	KCI 0.1M	12,900
Tap water	100 - 1500	Ocean water	53,000
River water	250 - 800	H2SO4	82,600

- The Conductivity electrode needs to be kept clean and free of deposits and other types of build-up. This epoxy cell type of electrode has 'easy to clean' graphite plates which are corrosion safe and less easily contaminated.
- If used in solutions with a high ion concentration, it is possible for the graphite plates to become contaminated. Soak the electrode cell portion in water with a mild detergent for 15 minutes. Then soak in a dilute acid solution such as 0.1 mol dm⁻³ hydrochloric acid or 0.5 mol dm⁻³ acetic acid for another 15 minutes. Rinse well with distilled water.
- The most common reason for inaccurate measurements is cross contamination of samples. Take care not to transfer droplets of one sample to another. Wash the electrode with distilled water between test solutions and shake vigorously to remove droplets. Ideally airdry and then wash the cell in a sample of the solution to be measured.
- Be sure that samples are capped to prevent evaporation. It is best to fill sample bottles to the brim to prevent a gas such as carbon dioxide dissolving in the water sample.
- Do not use in a situation that could result in damage to the graphite plates in the cell chamber. Do not attempt to blot or wipe the inside of the cell.
- The automatic temperature compensation for this electrode operates over the range 10°C to 35°C, but it can be placed in solutions within a temperature range of 0 to 80°C.
- There may be a decrease in accuracy when used in solutions that have a conductance of above 20 mS (20,000 μ S). Measurements above 50 mS will be approximate. If samples need to be diluted, use fresh deionised water (to reduce inaccuracy caused by variation in the conductivity of the deionised water).
- The Conductivity electrode not only measures conductivity between the graphite plates but also, to a lesser extent, in a field to the sides the electrode. In a narrow vessel, the walls may interfere with this field. If the electrode is held too close to the top of the liquid level or other objects (e.g. the bottom of a beaker) an incorrect reading may result.
- Interference may occur between electrochemical sensors (**pH**, **Oxygen**, and **Conductivity**) if they are placed in the same solution at the same time and connected to the same EasySense unit. This is because these sensors make an electrical connection to the solution; therefore an electrical path exists between the sensors through the solution. Maximise the distance between the Sensors to minimise the effect, the distance required will depend on the conductivity of the solution.
- If the Sensors are being used in a solution that has a fairly high conductance e.g. seawater, either connect the Sensors to different EasySense units or take readings from the Sensors individually. (Place one sensor in the solution, take a reading, and remove from the solution. Place the other sensor in the solution, take a reading and remove).
- Total solids (TDS) can be estimated in ppm (part per million) i.e. mg/L by multiplying the reading in microSiemens by 0.67, i.e. 200 μ S x 0.67 = 134 ppm.



Do not place the electrode in

- viscous organic liquids such as heavy oils, glycerine or ethylene glycol
- in acetone
- non-polar solvents such as pentane or hexane.

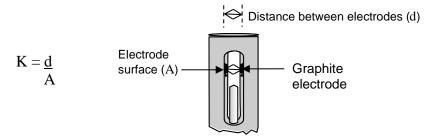
Theory

Electrical conductivity is an inherent property of most materials. Electrons carry electric current in metal; in water electrical current is carried by charged ions. The conductivity of a solution is determined by the ability of a solution to conduct an electric current between two electrodes. This will depend on the number of charge carriers, how fast they move, and how much charge each one carries.

For most water solutions, the higher the concentration of dissolved salts, and therefore more ions, the higher the conductivity. Low conductivity will indicate an absence of ions and therefore purity of water.

Conductivity is measured by putting a voltage across two electrodes, which have a fixed area and are a fixed distance apart. The electric current that flows between the two electrodes varies as the conductivity of the solution changes.

Each Conductivity electrode has its own physical characteristics. This is defined as the cell constant (K) and is dependent on the distance (d) between the two electrodes, and the area of the electrode surface (A).



The Conductivity electrode supplied has a nominal cell constant (K) of 1.0 cm⁻¹.

Alternating Current is used to prevent complete ion migration to the two electrodes. With each A.C. cycle the polarity of the electrodes are reversed which in turn reverses the direction of ion flow. This will prevent electrolysis and polarisation from occurring.

Although the presence of ions due to dissolved salts will increase the conductivity of water, it is not specific and can only be used to determine solution concentration if a single known salt is present.

Some species ionise more completely in water than others do, and their solutions are more conductive as a result. Each acid, base or salt has its own characteristic curve for concentration vs. conductivity. The relationship persists until very large ion concentrations are reached.

Temperature compensation

Temperature has a large effect on conductivity. The Conductivity electrode incorporates an inbuilt temperature sensor that is used to compensate for changes in the conductivity of solutions with temperature.

The temperature compensation is to counteract real changes in the conductivity of solution with temperature, not variations in the electrode with temperature.

For example: a 0.01mol dm⁻³ KCl solution has a conductivity of 1413 μ S at 25°C. Its conductivity changes from 1020 μ S at 10°C to 1552 μ S at 30°C. The *Smart* Q Conductivity Sensor will read a value of approximately 1413 μ S at any temperature in the 10 to 35°C range.

This allows the conductivity of sample solutions to be compared without bringing them all too exactly the same temperature.





Cleaning, storage and maintenance of the electrode

Depending on the sample application, the electrode may require cleaning periodically to ensure accurate measurements.

- Water-soluble contamination can be removed by soaking in distilled water.
- Petroleum-based contamination can be removed by soaking in warm water and a mild detergent for 15 minutes.
- Ethanol may be used to clean the electrode as long as the wash time is limited to a maximum of 5 minutes.
- Lime or hydroxide coating can be removed by soaking in a dilute acid solution such as 0.1 mol dm⁻³ hydrochloric acid or 0.5 mol dm⁻³ acetic acid for 15 minutes.

After cleaning, rinse well with distilled water, shake vigorously and leave to air-dry.

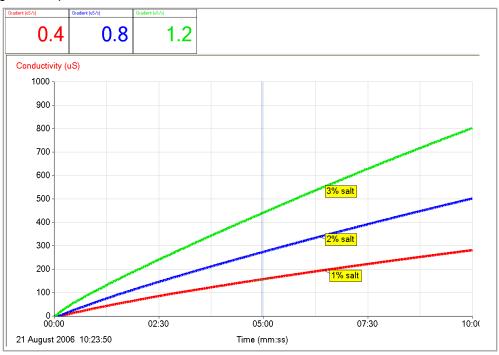
Store the electrode dry.

Investigations

- Electrolytes and non-electrolytes
- Conductivity of saltwater
- Equivalence point
- Conducting solutions
- Water quality using the $0 1000 \,\mu\text{S}$ range to test samples of water.
- Milk quality
- Speed of diffusion
- Strength of weak acids & bases
- Finding the concentration of unknown samples
- Diffusion through membranes

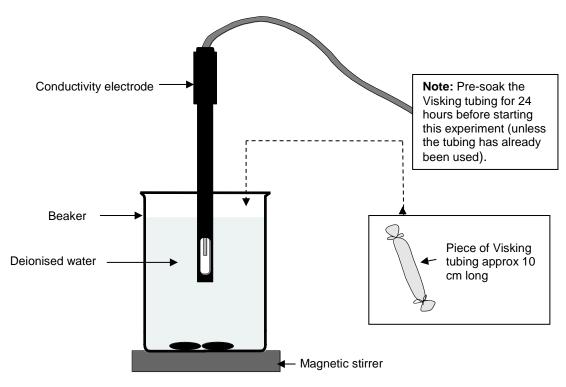
Does the concentration of a solution affect movement across a membrane?

For this graph we used analytical water with a conductivity of less than 10 μ S with 1%, 2% and 3% test solutions using the 0 – 1,000 μ S range. Gradient was used to calculate the rate of change at mid-point.









Test solutions:

No 1 = 1% w/v Salt solution

No 2 = 2% w/v Salt solution

No 3 = 3% w/v Salt solution

No 4 = 4% w/v Salt solution

- 1. Attach the Conductivity sensor to the EasySense unit.
- 2. Open the EasySense program and select **Graph** from the Home screen. From the Logging Wizard select a **10 minute** recording time with an interval of **1 second**.
- 3. Place the Conductivity electrode in the deionised water. Turn the magnetic stirrer on to give a slow agitation.
- 4. Select **Test Mode** from the **Tools** menu to determine the initial conductivity of the deionised water. Use this information to select a suitable range (see page 3).

Note: The $0 - 1,000 \,\mu\text{S}$ range would be suitable for water that has a conductivity of less than 50 μ S. If the value is between 50 - 400 μ S consider still using this range but reduce the percentage of salt in the test solutions e.g. 0.5%, 1%, 1.5%, 2%. If the value is higher than 400 μ S, use the 0 to 10 mS range and increase the percentage of salt in the test solutions.

- 5. Fill the Visking tubing with the 1% salt solution. Seal the open end. Rinse the Visking tube under running water. Place the tubing in the beaker of water and click on the **Start** icon to begin logging.
- 6. When the logging has finished discard the contaminated water and rinse the electrode with fresh deionised water.
- 7. Select Overlay.
- 8. Refill the beaker with the same amount of fresh deionised water and repeat the experiment using the other test solutions.



Limited warranty

For information about the terms of the product warranty, see the Data Harvest website at: https://data-harvest.co.uk/warranty.

Note: Data Harvest products are designed for **educational** use and are not intended for use in industrial, medical or commercial applications.



WEEE (Waste Electrical and Electronic Equipment) Legislation

Data Harvest Group Ltd is fully compliant with WEEE legislation and is pleased to provide a disposal service for any of our products when their life expires. Simply return them to us clearly identified as 'life expired' and we will dispose of them for you.