

Ion Selective Electrodes



Ammonium Ion-Selective Electrode	(Order Code NH4-BTA or NH4-DIN)
Calcium Ion-Selective Electrode	(Order Code CA-BTA or CA-DIN)
Chloride Ion-Selective Electrode	(Order Code CL-BTA or CL-DIN)
Nitrate Ion-Selective Electrode	(Order Code NO3-BTA or NO3-DIN)

The Vernier Ion-Selective Electrodes can be used to measure the concentration of a specific ion in aqueous samples. The species available include Nitrate (NO_3^-), Chloride (Cl^-), Calcium (Ca^{2+}), and Ammonium (NH_4^+). These electrodes can be used to perform a wide variety of tests or planned experiments, including the following:

- **Water Hardness (Calcium ISE):** Calcium is very useful as an indicator of hardness of water in freshwater samples. Even though water hardness is also due to other ions (Mg^{2+} and Fe^{3+}), because Ca^{2+} is normally the predominate hardness ion, hardness as Ca^{2+} is a reliable indicator of water hardness.
- **Nitrate Monitoring (Nitrate ISE):** Students can use this ISE to qualitatively determine the concentration of the nitrate ion, NO_3^- , in freshwater samples. This ion may appear in samples due to waste or fertilizer runoff. Increased levels of nitrate may also be present in waste from plants or in water samples with animal decay.
- **Salinity and Chloride Monitoring in Seawater (Chloride ISE):** Use this ISE to determine the concentration of chloride ion, Cl^- , or salinity levels in saltwater samples. The wide range of this electrode allows you to measure salinity (as sodium chloride), chloride concentration, seawater, or brackish water samples.
- **Monitoring Streams for Ammonium (Ammonium ISE):** Ammonium (NH_4^+) is often present as a result of water runoff from fertilized fields. This ISE gives you an easy way to check for this environmentally important ion.

Inventory of Items Included with the ISE

- One Ion-Selective Electrode (Nitrate, Calcium, Chloride, or Ammonium).
- One bottle of High Standard solution with MSDS sheet (1000 mg/L for Ca^{2+} and Cl^- , or 100 mg/L for NO_3^- and NH_4^+).
- One bottle of Low Standard solution with MSDS sheet (10 mg/L for Ca^{2+} and Cl^- , or 1 mg/L for NO_3^- and NH_4^+).
- One ISE Soaking Bottle (empty bottle with insertion cap).

NOTE: This product is to be used for educational purposes only. It is not appropriate for industrial, medical, research, or commercial applications.

Collecting Data with the ISE

This sensor can be used with the following interfaces to collect data:

- Vernier LabQuest® as a standalone device or with a computer
- Vernier LabQuest® Mini with a computer
- Vernier LabPro® with a computer, TI graphing calculator, or Palm® handheld

- Vernier Go!® Link
- Vernier EasyLink®
- Vernier SensorDAQ®
- CBL 2™

Data-Collection Software

This sensor can be used with an interface and the following data-collection software.

- **Logger Pro 3** This computer program is used with LabQuest, LabQuest Mini, LabPro, or Go!Link.
- **Logger Pro 2** This computer program is used with ULI or Serial Box Interface.
- **Logger Lite** This computer program is used with LabQuest, LabQuest Mini, LabPro, or Go!Link.
- **LabQuest App** This program is used when LabQuest is used as a standalone device.
- **EasyData App** This calculator application for the TI-83 Plus and TI-84 Plus can be used with CBL 2, LabPro, and Vernier EasyLink. We recommend version 2.0 or newer, which can be downloaded from the Vernier web site, www.vernier.com/easy/easydata.html, and then transferred to the calculator. See the Vernier web site, www.vernier.com/calc/software/index.html for more information on the App and Program Transfer Guidebook.
- **DataMate program** Use DataMate with LabPro or CBL 2 and TI-73, TI-83, TI-84, TI-86, TI-89, and Voyage 200 calculators. See the LabPro and CBL 2 Guidebooks for instructions on transferring DataMate to the calculator.
- **Data Pro** This program is used with LabPro and a Palm handheld.
- **LabVIEW** National Instruments LabVIEW™ software is a graphical programming language sold by National Instruments. It is used with SensorDAQ and can be used with a number of other Vernier interfaces. See www.vernier.com/labview for more information.

Collecting Data with the ISE

Here is the general procedure to follow when using the ISE:

1. The Ion-Selective Electrode (ISE) must be soaked in the High Standard solution (included with the ISE) for approximately 30 minutes. **Important:** Make sure the ISE is not resting on the bottom of the container, and that the small white reference contacts are immersed. Make sure no air bubbles are trapped below the ISE.
2. If the ISE needs to be transported to the field during the soaking process, use the Short-Term ISE Soaking Bottle. Remove the cap from the bottle and fill it 3/4-full with High Standard. Slide the bottle's cap onto the ISE, insert it into the bottle, and tighten. **Important:** Do not leave the ISE soaking for more than 24 hours. Long-term storage should be in the Long-Term ISE Storage Bottle.
3. Connect the ISE to the interface.
4. Start the data-collection software¹.

¹ If you are using Logger Pro 2 with either a ULI or SBI, the sensor will not auto-ID.

5. The software will identify the ISE. See below for calibration instructions.

Calibrating the ISE with a Computer

1. Choose Calibrate from the Experiment menu and then click .
2. High Standard Calibration Point: The ISE should still be soaking in the High Standard. Enter the concentration value of the High Standard (e.g., 100 for 100 mg/L or 1000 for 1000 mg/L) in the edit box.
3. When the displayed voltage reading for Reading 1 stabilizes (~1 minute), click .
4. Low Standard Calibration Point: Remove the ISE from the High Standard, rinse well with distilled water from a wash bottle, and gently blot dry with a paper towel or lab wipe. Place the electrode into the Low Standard (included with your ISE). Important: Make sure the ISE is not resting on the bottom of the container, and that the small white reference contacts are immersed. Make sure no air bubbles are trapped below the ISE.
5. Enter the concentration value of the Low Standard (e.g., 1 for 1 mg/L or 10 for 10 mg/L). After 60 seconds, click and then .
6. After calibrating, rinse off the end of the ISE, and blot it dry with a paper towel or lab wipe.
7. Insert the tip of the ISE into the sample to be tested. Important: Make sure the ISE is not resting on the bottom of the container, and that the small white reference contacts are immersed. Make sure no air bubbles are trapped below the ISE.
Note: Do not completely submerge the sensor. The handle is not waterproof.
8. Hold the ISE still for 60 seconds and record the displayed reading.

Calibrating the ISE with TI Graphing Calculators or Palm Handhelds

Enter the calibration routine for your data-collection program. Consult the reference material for the software on your handheld to get specific information concerning the calibration steps.

- High Standard Calibration Point: The ISE should still be soaking in the High Standard. Enter the concentration value of the High Standard (e.g., "100" for 100 mg/L or "1000" for 1000 mg/L).
- Low Standard Calibration Point: Remove the ISE from the High Standard, rinse well with distilled water from a wash bottle, and gently blot dry with a paper towel or lab wipe. Place the electrode into the Low Standard (included with your ISE). Important: Make sure the ISE is not resting on the bottom of the container, and that the small white reference contacts are immersed. Make sure no air bubbles are trapped below the ISE. After 60 seconds, enter the concentration value of the Low Standard (e.g., "1" for 1 mg/L or "10" for 10 mg/L).

Collecting Data

- After calibrating, rinse off the end of the ISE, and blot it dry with a paper towel or lab wipe.

Open an experiment file for the Colorimeter in the Probes & Sensors folder.

- Insert the tip of the ISE into the sample to be tested. Important: Make sure the ISE is not resting on the bottom of the container, and that the small white reference contacts are immersed. Make sure no air bubbles are trapped below the ISE.
Note: Do not completely submerge the sensor. The handle is not waterproof.
- Hold the ISE still for 60 seconds and record the displayed reading.

Storing Your Ion-Selective Electrodes

Proper care and storage of your ISE are important considerations for optimal longevity, especially with the PVC membrane ISEs (Ammonium, Calcium, and Nitrate).

- Long-term storage of ISEs (longer than 24 hours) should be done in the Long-Term ISE Storage Bottle, which is included. The sponge in the bottom of the bottle should be kept moist with distilled water. When you finish using the ISE, simply rinse it off with distilled water and blot it dry using a paper towel or lab wipe. Loosen the lid of the bottle and insert the ISE. NOTE: The tip of the ISE should NOT be touching the sponge. Check to be sure the white ceramic junctions are inside, rather than outside the bottle or under the grommet. Tighten the lid. This storage system keeps the electrode in a humid environment, which prevents the reference junctions from completely drying out.
- Short-term, wet storage (less than 24 hours) can be done by using the Short-Term ISE Soaking Bottle included with the electrode. Fill the bottle 3/4 full with High Standard. Loosen the cap, insert the electrode into the bottle, and tighten.

This sensor is equipped with circuitry that supports auto-ID. When used with LabQuest, LabPro, Go! Link, SensorDAQ, EasyLink, or CBL 2, the data-collection software identifies the sensor and uses pre-defined parameters to configure an experiment appropriate to the recognized sensor.

Specifications

Range of ISE Electrodes

Ammonium (NH ₄ ⁺):	0.10 to 18,000 mg/L or ppm (5.5 × 10 ⁻⁶ M to 1.0 M)
Calcium (Ca ²⁺):	0.20 to 40,000 mg/L or ppm (5 × 10 ⁻⁶ M to 1.0 M)
Chloride (Cl ⁻):	1.8 to 35,500 mg/L or ppm (5 × 10 ⁻⁵ M to 1.0 M)
Nitrate (NO ₃ ⁻):	0.10 to 14,000 mg/L or ppm (7 × 10 ⁻⁶ M to 1.0 M)

Resolution

13-bit (SensorDAQ)	the resolution values for each range will be half as large as those shown for 12-bit interfaces.
12-bit (LabQuest, LabQuest Mini, LabPro, Go! Link, ULI, or SBI):	% of reading, low scale reading, high scale reading
Ammonium (NH ₄ ⁺):	0.7% of reading (0.10 ± 0.0007 or 18,000 ± 130 mg/L)
Calcium (Ca ²⁺):	1.4% of reading (0.20 ± 0.0028 or 40,000 ± 560 mg/L)
Chloride (Cl ⁻):	0.7% of reading (1.8 ± 0.013 or 35,500 ± 250 mg/L)
Nitrate (NO ₃ ⁻):	0.7% of reading (0.10 ± 0.0007 or 14,000 ± 98 mg/L)
10-bit (CBL 2):	the resolution values for each range will be four times as large as those shown for 12-bit interfaces.

pH Range:

Ammonium (NH ₄ ⁺):	4 to 7.5
Calcium (Ca ²⁺):	3 to 10
Chloride (Cl ⁻):	2 to 12
Nitrate (NO ₃ ⁻):	2.5 to 11

Interfering Ions:

Ammonium (NH ₄ ⁺):	K ⁺
Calcium (Ca ²⁺):	Pb ²⁺ , Hg ²⁺ , Sr ²⁺ , Cu ²⁺ , Ni ²⁺
Chloride (Cl ⁻):	CN ⁻ , Br ⁻ , I ⁻ , S ²⁻
Nitrate (NO ₃ ⁻):	ClO ₄ ⁻ , I ⁻ , ClO ₃ ⁻ , CN ⁻ , BF ₄ ⁻

Electrode Slope (log voltage vs. concentration):

Ammonium (NH ₄ ⁺):	+56 mV/decade
Chloride (Cl ⁻):	-56 mV/decade
Calcium (Ca ²⁺):	+28 mV/decade
Nitrate (NO ₃ ⁻):	-56 mV/decade

Approximate Calibration Voltages:

Ammonium (NH ₄ ⁺):	High (100 mg/L) 2.1 V	Low (1 mg/L) 1.3 V
Calcium (Ca ²⁺):	High (1000 mg/L) 1.9 V	Low (10 mg/L) 1.5 V
Chloride (Cl ⁻):	High (1000 mg/L) 2.0V	Low (10 mg/L) 2.8 V
Nitrate (NO ₃ ⁻):	High (100 mg/L) 1.6	Low (1 mg/L) 2.4 V

Electrode Resistance: 1 to 4 MΩ

Reproducibility: ± 5% of reading

Temperature range (can be placed in): 0 to 50°C (no temperature compensation)

Minimum immersion: 1 inch

Electrode Length: 155 mm

Body Diameter: 12 mm

Cap Diameter: 16 mm

Cable Length: 100 cm

How Your Ion-Selective Electrode Works

Vernier Ion-Selective Electrodes (ISEs) are membrane-based electrodes that measure a specific ion (e.g., Ca²⁺) in an aqueous solution. When the membrane of the electrode is in contact with a solution containing the specific ion, a voltage, dependent on the level of that ion in solution, develops at the membrane. All of our ISEs are combination style; that is, the voltage develops in relation to an internal Ag/AgCl reference electrode. The ISEs measure for the specific ion concentrations directly. Samples need to be aqueous to avoid contaminating or dissolving the membrane. Vernier Ion-Selective Electrodes (ISEs) have two types of membranes:

- Solid Polymer Membrane: This type of PVC membrane is used in our Nitrate, Calcium, and Ammonium ISEs. The membrane is a porous plastic disk,

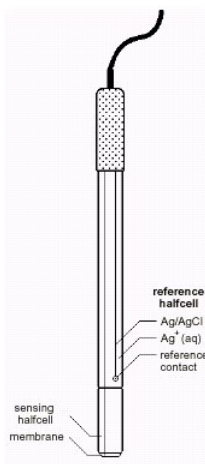


Figure 2

permeable to the ion exchanger, but impermeable to water. It allows the sensing cell to contact the sample solution and separates the internal filling solution from the sample. Our PVC membrane electrodes have replaceable membrane modules.

- Solid State Membrane: This thin crystal disk, used in our Chloride ISE, is an ionic conductor. It separates the internal reference solution from the sample solution. The Chloride ISE does not have replaceable membrane modules.

The voltage developed between the sensing and reference electrodes is a measure of the concentration of the reactive ion being measured. As the concentration of the ion reacting at the sensing electrode varies, so does the voltage measured between the two electrodes.

As described in the Nernst equation, ISE response is a linear equation:

$$E = E_o + m(\ln a)$$

where E is the measured voltage, E_o is the standard potential for the combination of the two half cells, m is the slope, ln is natural log, and a is the activity of the measured ion species.

Assuming the ionic strength is fairly constant, the Nernst equation may be rewritten to describe the electrode response to the concentration, C, of the measured ionic species:

$$E = E_o + m(\ln C)$$

A graph of the natural log of concentration (lnC) vs. potential (V) can be plotted:

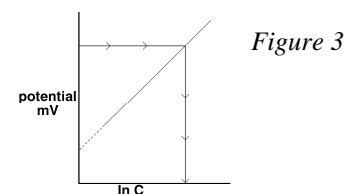


Figure 3

The method for determining the concentration of Ca²⁺, NH₄⁺, NO₃⁻, or Cl⁻ ions using one of our ISEs is based on the following:

- The interface reads a voltage that is related to ion concentration.
- Using the voltage reading and the principle described in Figure 3, the natural log of concentration of the specific ion can be determined.
- The ion concentration can be determined from the natural log of concentration.

Maintaining and Replacing the ISE Standard Calibration Solutions

Having accurate standard solutions is essential for performing good calibrations. The two standard solutions that were included with your ISE can last a long time if you take care not to contaminate them with a wet or dirty probe. At some point, you will need to replenish your supply of standard solutions.

Vernier sells replacement standards in 500 mL bottles. Order codes are:

Ammonium Low Standard: NH4-LST	Ammonium High Standard: NH4-HST
Calcium Low Standard: CA-LST	Calcium High Standard: CA-HST
Chloride Low Standard: CL-LST	Chloride High Standard: CL-HST
Nitrate Low Standard: NO3-LST	Nitrate High Standard: NO3-HST

To prepare your own standard solutions, use the information in Table 1.

Note: Use glassware designed for accurate volume measurements, such as volumetric flasks or graduated cylinders. All glassware must be very clean.

Table 1: Replacing ISE Standard Solutions

Standard Solution	Concentration (mg/L or ppm)	Preparation Method using High Quality Distilled Water
Ammonium (NH ₄ ⁺) ISE High Standard	100 mg/L NH ₄ ⁺ as N	0.382 g NH ₄ Cl / 1 L solution
Ammonium (NH ₄ ⁺) ISE Low Standard	1 mg/L NH ₄ ⁺ as N	Dilute the High Standard by a factor of 100 (from 100 mg/L to 1 mg/L).*
Calcium (Ca ²⁺) ISE High Standard	1000 mg/L as Ca	2.771 g CaCl ₂ / 1 L solution or 3.669 g CaCl ₂ •2H ₂ O / 1 L solution
Calcium (Ca ²⁺) ISE Low Standard	10 mg/L as Ca	Dilute the High Standard by a factor of 100 (from 1000 mg/L to 10 mg/L).*
Chloride (Cl ⁻) ISE High Standard	1000 mg/L as Cl	1.648 g NaCl / 1 L solution
Chloride (Cl ⁻) ISE Low Standard	10 mg/L as Cl	Dilute the High Standard by a factor of 100 (from 1000 mg/L to 10 mg/L).*
Nitrate (NO ₃ ⁻) ISE High Standard	100 mg/L NO ₃ as N	0.607 g Na NO ₃ / 1 L solution
Nitrate (NO ₃ ⁻) ISE Low Standard	1 mg/L NO ₃ as N	Dilute the High Standard by a factor of 100 (from 100 mg/L to 1 mg/L).*

*Perform two serial dilutions as described below.

- Combine 100 mL of the High Standard with 900 mL of distilled water. Mix well.
- Combine 100 mL of the solution made in Step (a) with 900 mL of distilled water. Mix well.

Replacement Membrane Modules (Ammonium, Nitrate, and Calcium only)

The Nitrate, Calcium, and Ammonium ISEs each have a PVC membrane with a limited life expectancy. They are warranted to be free from defects for a period of twelve (12) months from the date of purchase; it is possible, however, that you may get somewhat longer use than the warranty period. If you start to notice a reduced response (e.g., distinctly different voltages or voltage ranges during calibration), it is probably time to replace the membrane module. **Important:** Do not order membrane modules far in advance of the time you will be using them; the process of degradation takes place even when they are stored on the shelf. Note that the Chloride ISE has a solid-state membrane, with a longer membrane life, and does not need a replacement module.

Using ISA (Ionic Strength Adjuster) Solutions to Improve Accuracy

For optimal results at low concentrations of calcium, chloride, and nitrate ions, a standard method for making measurements with ion-selective electrodes is to add ionic strength adjuster (ISA) solutions to each of your standard solutions and samples.

Note: Do not use an ISA for low concentrations of ammonium.

Adding an ISA ensures that the total ion activity in each solution being measured is nearly equal, regardless of the specific ion concentration. This is especially important when measuring very low concentrations of specific ions. The ISA contains no ions common to the Ion-Selective Electrode itself. Note: The additions of ISA to samples or standards described below do not need to have a high level of accuracy—combining the ISA solution and sample solution counting drops using a disposable Beral pipet works fine. The following are instructions for using ISA solutions with Vernier Ion-Selective Electrodes.

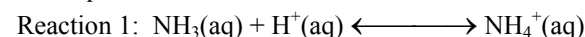
Calcium ISA: Add the 1.0 M KCl ISA solution (7.46 g KCl / 100 mL solution) to the Ca²⁺ standard or to the solution being measured, in a ratio of 1 part of ISA (by volume) to 50 parts of total solution (e.g., 1 mL of ISA to 50 mL of total solution, or 2 drops of ISA to 5 mL of total solution).

Chloride ISA: Add the 5.0 M NaNO₃ ISA solution (42.50 g NaNO₃ / 100 mL solution) to the Cl⁻ standard or to the solution being measured, in a ratio of 1 part of ISA (by volume) to 50 parts of the total solution (e.g., 1 mL of ISA to 50 mL of total solution, or 2 drops of ISA to 5 mL of total solution).

Nitrate ISA: Add the 2.0 M (NH₄)₂SO₄ ISA solution (26.42 g (NH₄)₂SO₄ / 100 mL solution) to the NO₃⁻ standard or to the solution being measured, in a ratio of 1 part of ISA (by volume) to 50 parts of total solution (e.g., 1 mL of ISA to 50 mL of total solution, or 2 drops of ISA to 5 mL of total solution).

Using the Ammonium Ion-Selective Electrode

The Ammonium ISE can be used to determine concentrations of NH₄⁺ ions in aqueous solutions, in units of mg/L, ppm, or mol/L. Concentrations of aqueous ammonium ions should not be mistaken for concentration of aqueous ammonia, or NH₃(aq). The concentrations of these two species, though different, are often involved in the same equilibrium reaction:



In a more acidic environment, higher concentrations of H⁺ ions will cause this reaction to shift toward the right, resulting in higher concentrations of NH₄⁺. In a more basic (alkaline) environment, the concentration of NH₄⁺ will be lower, causing the reaction to shift toward the reactants, producing higher concentrations of NH₃. At pH values greater than 10 (see Figure 4), most of the ammonium ions will be converted to ammonia. At pH values less than 7.5, most of the aqueous ammonia will be converted to ammonium ions.

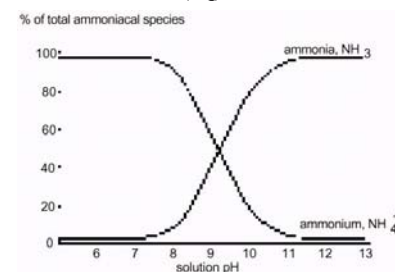


Figure 4

Sampling Freshwater Samples for Ammonium Concentration

While permissible levels of ammonium in drinking water should not exceed 0.5 mg/L, streams or ponds near heavily fertilized fields may have higher concentrations of this ion. Fertilizers containing ammonium sulfate, (NH₄)₂SO₄, or ammonium nitrate, NH₄NH₃, may result in runoff from fields containing higher levels of the ammonium ion, NH₄⁺. Monitoring ammonium levels on a stream that borders fertilized fields may show significant seasonal differences in NH₄⁺ concentrations. In this kind of study,

you may also take pH measurements in your water samples; as indicated in the previous paragraph, higher or lower pH values can greatly affect the ratio of NH_4^+ / NH_3 in a sample. Since the Ammonium ISE measures only NH_4^+ levels, you may want to adjust your samples to the same pH value each time you make measurements; this may not be necessary if you have relatively “hard” water. Hard water is naturally buffered against changes in pH.

Using the Ammonium ISE at Low Concentration Levels (0.1-0.5 mg/L NH_4^+ -N)

If the concentration of your sample is less than 0.5 mg/L NH_4^+ -N, you will obtain better results if you calibrate the electrode with 0.1 mg/L and 1.0 mg/L NH_4^+ -N. This is because the slope of the calibration changes at these low levels. You can make up the 0.1 mg/L Ammonium Standard by following the directions in Table 1 to make the 1.0 mg/L NH_4^+ -N Standard, then diluting one more time using 100 mL of the 1.0 mg/L with 900 mL of high quality distilled water.

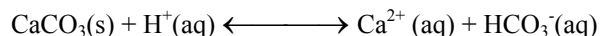
Expressing Ammonium Concentration

Concentrations of ammonium are often expressed in units of mg/L NH_4^+ as N. Here is a calculation for a 100 mg/L NH_4^+ as N standard solution that is prepared by adding solid NH_4Cl to distilled water:

$$\frac{100 \text{ mg } \text{NH}_4^+}{1 \text{ L}} \times \frac{1 \text{ g } \text{NH}_4^+}{1000 \text{ mg } \text{NH}_4^+} \times \frac{53.5 \text{ g } \text{NH}_4\text{Cl}}{14.0 \text{ g } \text{NH}_4^+ - \text{N}} = 0.382 \text{ g } \text{NH}_4\text{Cl}/\text{L solution}$$

Using the Calcium Ion-Selective Electrode

Your Calcium Ion-Selective Electrode can be used to determine the concentration of aqueous Ca^{2+} ions, in the range of 1.8 to 40,100 mg/L. It can be especially useful in determining “hardness of water.” Calcium ions are often found in freshwater samples as a result of water flowing over soil and mineral deposits containing limestone, chalk, magnesite, or dolomite. In one common reaction, limestone is dissolved according to the reaction



This reaction and others similar to it produce water with a relatively high concentration of Ca^{2+} ions, and lesser concentrations of Mg^{2+} and Fe^{3+} ions—known as hard water.

Using the Calcium ISE to Determine Water Hardness as Ca^{2+}

Many methods of determining water hardness use “total hardness,” or the sum of hardness due to Mg^{2+} and Ca^{2+} . Since the Ca^{2+} concentration of freshwater usually exceeds that of Mg^{2+} , determining the Ca^{2+} concentration alone is a good indicator of water hardness—we will refer to this measurement as “calcium hardness.” For best results, calibrate the Calcium ISE using the 10 mg/L and 1000 mg/L standards.

Using the standard solutions described here, your results will be in units of mg/L of Ca^{2+} . Units of calcium hardness are usually expressed as “calcium hardness as CaCO_3 ”. To convert from units of mg/L of Ca^{2+} (150 mg/L is used in this example) to units of calcium hardness as CaCO_3 , in mg/L, you would use this expression:

$$\frac{150 \text{ mg } \text{Ca}^{2+}}{1 \text{ L}} \times \frac{100 \text{ g } \text{CaCO}_3}{40 \text{ g } \text{Ca}^{2+}} = 374 \text{ mg/L (hardness as } \text{CaCO}_3\text{)}$$

It is important to remember that total hardness, taking into account both the Ca and Mg levels, will be about 1.5 times higher than your calcium hardness value. Water hardness varies considerably in different parts of the United States, from levels of less than 60 mg/L (total hardness as CaCO_3) in Washington, Oregon, Louisiana, Mississippi, Tennessee, and New England, to levels exceeding 250 mg/L or higher in Midwestern states (Ohio, Indiana, Illinois, Iowa, Nebraska, South Dakota, and Oklahoma). Water with a hardness as CaCO_3 level greater than 120 mg/L is considered to be “hard,” while levels exceeding 180 mg/L are referred to as “very hard.” equivalence point occurs first, with the Mg^{2+} equivalence point following; therefore Ca^{2+} can be determined as well as total hardness (as Ca^{2+} and Mg^{2+}). A plot of $\ln [\text{Ca}^{2+}]$ (natural log of calcium ion concentration) vs. volume is used to determine the equivalence point. The second derivative can be used to calculate the point of maximum inflection at the equivalence point of the titration.

Using the Chloride Ion-Selective Electrode

Chloride ions are found in freshwater samples as a result of water flowing over salt-containing minerals. These salts might include either sodium chloride (NaCl) or potassium chloride (KCl). The EPA maximum contamination level for chloride concentration in drinking water is 250 mg/L. The chloride ion concentration in seawater is approximately 19,400 mg/L—well below the upper limit of the Chloride ISE of 35,500 mg/L.

When the response of the Chloride ISE begins to slow, the membrane may need polishing. Cut a small piece (about 1 inch square) from a polishing strip. Wet the end of the electrode and the dull side of the polishing strip thoroughly with distilled water. Using only moderate pressure, polish the end of the electrode by gently rubbing in a circular motion. This will remove the inactive layer of the membrane which impedes measurement. Rinse thoroughly with distilled water and recalibrate in the usual manner.

Sampling Freshwater Samples for Chloride Concentration

For best results, calibrate the Chloride ISE using the 10 mg/L and 1000 mg/L standards.

Measuring Chloride Concentration of Saltwater or Brackish Water

When measuring chloride concentration in seawater or brackish water, calibrate the Chloride ISE using the 1000 mg/L standard included with your Chloride ISE for one calibration point (or 1.806 parts per thousand, or ppt). For the second calibration point, prepare a standard that is 20,000 mg/L Cl^- by adding 32.96 g of solid NaCl to enough distilled water to prepare 1 L of solution:

$$\frac{20000 \text{ mg } \text{Cl}^-}{1 \text{ L}} \times \frac{1 \text{ g } \text{Cl}^-}{1000 \text{ mg } \text{Cl}^-} \times \frac{58.5 \text{ g } \text{NaCl}}{355 \text{ g } \text{Cl}^-} = 32.96 \text{ g } \text{NaCl} / \text{L solution}$$

If you are calibrating in ppt, call this solution 36.13 ppt.

Determining Salinity of Saltwater or Brackish Water

Salinity is the total of all salts dissolved in water, expressed either as mg/L (equal to parts per million, ppm) or in parts per thousand (ppt). Seawater contains a fairly

constant quantity of chloride ions. From your measurement of chloride ion concentration (in the previous section), salinity can be calculated using the following formula:

$$\text{Salinity (mg/L or ppm)} = 1.8066 \times [\text{Cl}^- \text{ concentration, mg/L}]$$

Using this formula, the salinity of saltwater is calculated to be:

$$\text{Salinity (mg/L or ppm)} = 1.8066 \times (19400 \text{ mg/L}) = 35,000 \text{ mg/L}$$

The level of salinity of seawater in parts per thousand, or ppt, would be:

$$\text{Salinity (ppt)} = 35000 / 1000 = 35 \text{ ppt}$$

Using the Nitrate Ion-Selective Electrode

Nitrate ions, NO_3^- , may be found in freshwater samples from a variety of sources. Sewage is often the primary source. Sometimes nitrates are present due to runoff from fertilized fields. Nitrates can also result from the runoff from cattle feedlots and barnyards. In all of these cases, as plant and animal organisms die, bacterial action breaks down the protein into ammonia, NH_3 . Some ammonia is converted into ammonium ions, NH_4^+ . Other bacterial action converts some of the ammonia and ammonium ions into nitrite ions, NO_2^- , and then into nitrate ions, NO_3^- .

Units of Nitrate Concentration

Nitrate ion concentration is usually expressed in units of mg/L of NO_3^- as N, also known as “nitrate-nitrogen.” This means that the concentration of nitrate is expressed as if the nitrate were only in the form of nitrogen itself. The standards that are included with your Nitrate ISE have concentrations of 1 and 100 mg/L of NO_3^- as N. Here is the calculation for making a 100 mg/L NO_3^- as N standard starting with solid NaNO_3 (as shown in Table 1). Notice that the atomic weight of N, 14.0, is used instead of the atomic weight of NO_3^- , 62.0.

$$\frac{100 \text{ mg N}}{1 \text{ L}} \times \frac{1 \text{ g N}}{1000 \text{ mg N}} \times \frac{85.0 \text{ g NaNO}_3}{14.0 \text{ g N}} = 0.607 \text{ g NaNO}_3 / \text{L solution}$$

Unpolluted waters usually have nitrate-nitrogen (NO_3^- as N) levels below 1 mg/L. Nitrate-nitrogen levels above 10 mg/L are considered unsafe for drinking water.

Test results are sometimes published in units of mg/L NO_3^- instead of NO_3^- as N. To convert 100 mg/L NO_3^- as N to mg/L NO_3^- , you would perform this conversion:

$$\frac{100 \text{ mg N}}{1 \text{ L}} \times \frac{62.0 \text{ g NO}_3}{14.0 \text{ g N}} = 443 \text{ mg/L NO}_3$$

Sampling Freshwater Samples for Nitrate Concentration

For best results, calibrate the Nitrate ISE, using the 1 mg/L and 100 mg/L standards.

How can I have my ISE read mV output instead of mg/L?

The amplification equation is: $V = 0.00727 * \text{mV} + 1.223$

Therefore, the reverse amplification equation, solving for mV, would be:

$$\text{mV} = 137.55 * V - 0.1682$$

Using ISEs with Other Vernier Sensors

It is very important for you to know ISEs will interact with some other Vernier sensors and probes, *if* they are placed in the same solution (in the same aquarium or beaker, for example), *and* they are connected to the same interface box (e.g., the same LabPro interface).

The following probes *cannot* be connected to the same interface as an ISE and placed in the same solution:

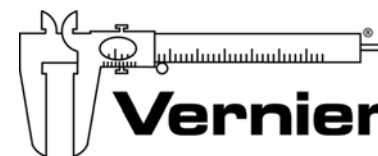
- Conductivity Probe
- pH Sensor
- Direct-Connect Temperature Probe
- DO Probe

If you wish to take simultaneous readings using any of the probe combinations listed above, here are some alternative methods:

- You can connect the probes to two different interfaces. If the two probes in question are connected to separate interfaces, the two probes will read correctly in the same solution.
- The Vernier Stainless Steel Temperature Probe (also shipped with CBL 2) can be used in the same container with ISEs.
- If you are sampling a lake or stream and want to use two of the probes with a single interface, you can connect the two probes in question to the same interface and load their respective calibrations. Place one probe in the water first and take its reading. Then remove it and place the second probe in the solution to take its reading.

Warranty

Vernier warrants this product to be free from defects in materials and workmanship for a period of five years from the date of shipment to the customer. This warranty does not cover damage to the product caused by abuse or improper use.



Measure. Analyze. Learn.™
Vernier Software & Technology

13979 S.W. Millikan Way • Beaverton, OR 97005-2886
Toll Free (888) 837-6437 • (503) 277-2299 • FAX (503) 277-2440
info@vernier.com • www.vernier.com

Rev 10/19/10

Logger Pro, Logger Lite, Vernier LabQuest, Vernier LabQuest Mini, Vernier LabPro, Go! Link, Vernier EasyLink and other marks shown are our trademarks or registered trademarks in the United States.

CBL 2 and CBL, TI-GRAPH LINK, and TI Connect are trademarks of Texas Instruments.

All other marks not owned by us that appear herein are the property of their respective owners, who may or may not be affiliated with, connected to, or sponsored by us.



Printed on recycled paper