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An Ion selective electrode is a sensor which converts the activity of a specific ion (dissolved in a solution) into a voltage (potential), which can be measured by a mV or Ion meter. The voltage is theoretically dependent on the logarithm of the ion activity, as described by the Nernst Equation.

E = E⁰ + (2.303*RT/nF)log(A)

The sensing part of the electrode is usually made from an ion specific membrane, coupled together with a reference electrode (either separate or as a combination).

There are five main types of ion selective membranes which are detailed hereunder.

One

Glass membrane electrodes are formed from special ionically conducting glass. By altering the composition and constituents of the glass, changes can be made to selectivity, chemical resistance, response time, and impedance. The most common glass membrane ion selective electrode is the pH electrode. Another common one is the sodium selective electrode.

Two

Crystalline or solid state membrane electrodes are made from relatively insoluble ionically conducting inorganic salts. These are available in homogeneous and heterogeneous forms. They have good selectivity since only ions which can introduce themselves into the crystal lattice can interfere. Examples include the Fluoride electrode which uses a doped LaF3 crystal, and the Chloride electrode which uses silver chloride powder.

Three

Polymer Membrane Electrodes are based on special organic polymer membranes which contain various ion-exchange ionophores incorporated into an inert matrix. These are used in electrodes to measure ions such as Potassium, Calcium, and Nitrate.

Four

Gas sensing electrodes have gas permeable membranes and an internal solution. Gas molecules diffuse across the membrane and react with the solution. This pH change is then detected by the pH electrode. Common gas sensing electrodes include ammonia and carbon dioxide.

Five

Enzyme electrodes are based on the reaction of an enzyme reacting with a specific substrate, and the resultant product of this reaction (usually H+ or OH-) is detected by an electrode such as a pH electrode. These reactions occur within a special membrane. An example is the Glucose electrode.

The Reference Electrode

An important part of the measurement is the use of a stable reference electrode. Many ISE's incorporate their own reference electrode; these are usually either a single junction refillable type Ag/AgCl type, or the double junction type, which is used for ISE's such as chloride, bromide etc. These types of reference electrodes allow the user to select an appropriate electrolyte for the particular application. For instance Potassium Nitrate is commonly used as a filling solution for ISE's such as Chloride, Bromide, Iodide, Cyanide, Silver and Sulphide.Whenever choosing a filling solution, it is also important to select one that is close to being equitransferant to minimise junction potential errors. The E⁰ factor in the Nernst equation is the sum of all the liquid junction potentials present in the system, and any variation in this during analysis can introduce major sources of potentail drift and error in measurements.

Use of ISA/TISAB

For accurate and reproducible measurements, adjustment of the ionic strength is often required. The reason is to raise the the ionic strength of all standards and samples to a uniformly high level so that the activity coefficient is the same for the ions in both solutions. This enables the concentration of the samples to be more accurately determined. ISA's can alo be used to adjust the pH of samples. TISAB (Total Ionic Strength Adjuster Buffer), used for Fluoride Ion analysis is a common example of this. This solution adjusts the ionic strength, adjusts the pH and can also minimise interferences.

Sample Pre-Treatment

It is important to note that Ion selective electrodes measure the activity of the ionic analytes in solution. If the ion to be measured is complexed or bound, sample pre-treatment may be required before analysis. This can involve preliminary steps such as drying, washing and grinding, as well as extraction or dry ashing. This is particularly the case with samples such as plant material and soils. In the event of interferences and oxidation effects, adding an appropriate reagent to both samples and standards can be beneficial.

ISE Analytical Techniques



Simple procedure used for measuring a large number of samples. This can also easily be automated with sample changers for unattended analysis. Calibration is performed using a series of standards, preferably bracketing the expected sample range. ISA is added to both samples and standards in the same ratios, and the concentration of the samples is determined by comparison to the standards.

Standard Addition

The electrode is immersed in the sample solution and an aliquot of a standard solution containing the measured species is added to the sample. From the change in potential before and after the addition, the original sample concentration is determined.

Analate Addition

Often used to measure viscous samples, small or very concentrated samples. It can be used to overcome the effects of complex sample matrics, but is not suitable for dilute samples. Total concentration is measured even in the presence of complexing agents. The electrode is immersed in a standard solution containing the ion to be measured, and an aliquot of the sample is added to the standard. The original sample concentration is determined from the change in potential before and after the addition.

Analate Subtraction

Used for the measurement of ions for which no ISE exists. The electrode is immered in a reagent solution that contains an ion that the electrode senses, and which reacts with the sample. It is not suited for dilute samples. An example is using a Lead ISE to measure Sulphate.

Titration Indicators

Ion selective electrodes are very useful as end point detectors in titrations because they are unaffected by colour or turbidity. A common example is use of a chloride ISE (or a silver billet electrode) for determining the salt content of dairy products by silver nitrate titration. Titrations are much more precise (approx. 10x) than direct ISE calibration, but they are more time consuming and they have a narrower dynamic range. ISE's can also be used in titrations to determine ions for which there is no current ISE. For example, Lanthanum ions can be determined in this way using a Fluoride ISE as indicator.

Why Use Ion Selective Electrodes?

The main reasons Ion selective electrodes are popular:-

The initial set up is inexpensive (typically one only needs a pH/mV meter or Ion meter, the electrodes, a stirring stand, and some basic chemicals)

The measurements are unaffected by colour or turbidity in the sample

The sample pre-treatment is usually simple

The measurements can be done in "real time", and can be easily automated

Wide dynamic range-usually several decades

Where Are They Used?

Ion selective electrodes or ISE's are commonly used in environmental, food and agriculture, power plants, research and in clinical applications.

Examples include: Fluoride in drinking water, Calcium in beer, Nitrates in plants, Chloride in canned food products, and Calcium in dental studies.

Limitations

- selectivity/interferences
- accuracy, typically 2-5% relative
- calibrating solutions and samples need to have the ionic strength adjusted
- lifetime and maintenance requirements of polymer membrane electrodes
- sample pre treatment is sometimes necessary for non-ideal samples

Care and Maintenance

Care must be taken to avoid damaging the membrane surface. Follow the instructions supplied with the electrodes to assure maximum life. After extensive use, the membranes may become coated with contaminants or become scratched. The solid state types such as Chloride and Fluoride can usually be easily repaired with fine emery and then washing with distilled water to remove debris. You should never touch the membranes of ISE's such as Calcium and Potassium, because it is very easy to damage the surface. These should instead be rinsed in standard solution and the deposits removed with a fine jet of water.

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IJ Series Intermediate Junction Electrodes

Setting the benchmark in super-long life, versatility and value for money since 1972, the flagship of the lonode Lab sensor range is the renowned IJ Series.

Built to exacting standards and available in pH, ISE, ORP and Reference, the IJ Series benefits are many ... A removable sleeve and renewable junction for longer life and easy cleaning, an isolated secondary reference to effectively defeat contamination, an annular ground-glass junction to provide

fast response times and resistance to clogging, selectable electrolyte and an optional toughened glass spear membrane for direct penetration. An exceptional choice in virtually any sample, the IJ Series is unequalled in food, wine, dairy, contaminated waste samples, industrial, mining, petro-chemical, electro plating and countless more. The IJ Series also features models with solvent resistant bodies and high temperature housings for the most demanding of environments.

Replaceable electrolyte

Removeable sleeve

Avaliable for pH, ISE & ORP

Polypropylene body

Way back In 1972,

History of the Intermediate Junction

Back in 1972, Ionode's R&D Chemists and Engineers set out to develop a sensor that would solve the the common problems of contamination and clogging etc. but still be tough, fast and super reliable. Working together, they decided it was possible and they set to work on what would become the IJ Series ... a sensor that would defeat contamination, work in any sample and have a lifetime far in excess of other single or double junction sensors. The benchmark in sensor versatility, reliability and performance was then born in the IJ Series. Featuring an renewable intermediate junction for easy cleaning, Ionode's proprietary glass formulae and a tough yet flexible body, the IJ Series quickly earned it's reputation for having the longest life in the industry and being the best value sensor available.

More Junctions, Less Contamination

IJ have unique junction mechanics which protect the primary reference from the contamination associated with traditional gel electrodes whilst providing the low maintenance and long life of a refillable sensor.

Removeable Sleeve

IJ electrodes have a removeable sleeve to make cleaning quick and easy.



IJ's Explained Why you need an intermediate junction probe

IJ's have unique junction mechanics which protect the primary reference from the contamination associated with traditional gel electrodes whilst providing the low maintenance and long life of a refillable sensor.





The IJ Range The full range of IJ electrodes

Model	Parameter	Measuring Range	Temp. Range	Sensor Type	Temperatur ← Swipe to see more information Compensat
IJ14	Ref		0 - 60	Reference	n/a
IJ44- Spear	рН	рН 0 - 14	0 - 60	A glass spear	Optional
IJ44- HT- Spear	рН	pH 0 - 14	0 - 100	A glass spear	Optional
IJ40	рН	рН 0 - 14	0 - 60	A glass bullet	Optional

IJ40- ALK	рН	рН 0 - 14	0 - 60	C Glass bullet	Optional
IJ40- HT- ALK	рН	рН 0 - 14	0 - 100	C Glass bullet	Optional
IJ64	ORP	±2000mV	0 - 60	Platinum Wire	n/a
IJ64D	ORP	±2000mV	0 - 60	Platinum Disc	n/a
IJ64- HT	ORP	±2000mV	0 - 100	Platinum Wire	n/a
IJAg	Ag⁺, Cl⁻	±2000mV	0 - 60	Silver Billet	n/a
IJAu	ORP	±2000mV	0 -60	Gold Wire	n/a

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Model	Parameter	Measuring Range	Temp. Range	Main Interferences	Slope
IJ-F	F ⁻	0.2 - 19,000 ppm	0 - 60	OH-	57mV/decade ± 5mV
IJ-Ag2S	S ²⁻	0.003 - 32,000 ppm	0 - 60	Hg ^{+/2+}	27mV/decade ± 3mV
	Ag+	0.01 - 108,000 ppm			57mV/decade ± 5mV
IJ-Cl	CI⁻	1.8 - 35,500 ppm	0 - 60	Br⁻, I⁻, S²⁻, CN⁻, S ₂ O ₃ ²⁻	57mV/decade ± 5mV
IJ-Br	Br⁻	0.2 - 79,900 ppm	0 - 60	I⁻, S²⁻, CN⁻	57mV/decade ± 5mV
IJ-I	l-	0.005 - 127,000 ppm	0 - 60	S ²⁻ , CN⁻	57mV/decade ± 5mV
IJ-CN	CN	0.2 - 260,000 ppm	0 - 60	S ²⁻ , I⁻	57mV/decade ± 5mV