## Field Analysis

Many nutrients as well as salinity (EC) can easily be measured with test kits (colorimeter), handheld meters or laboratory analysis. Some sources for such testing equipment are listed here:

Spectrum Technologies - 815-436-4440 - www.specmeters.com Hach Company - 970-669-3050 - www.hach.com Myron L Company - 760-438-2021 - www.myronl.com

Measuring pH with suction extraction can be done successfully, however the pH of the solution changes because some  $CO^2$  is lost due to the vacuum process. The loss can be calculated and thus compensated for during interpretation. Additional information is available upon request.

In any event, analysis of the field sample cannot be done in a "vacuum". The sample analysis needs to be compared to some base. The best "standard" to use would be the tissue (petiole) analysis done by a lab and in conjunction with known agronomic facts concerning the particular crop and its nutrient uptake rate in relation to the amount of that nutrient in the soil water. This is not totally precise, but once you establish a correlation, or ratio, you have a guideline for future adjustments in the soil water/nutrient balance.

If nothing else, once you establish this relationship, and look at a given soil water sample "reading," you will be able to verify that you do not need to add more nitrogen, for example. Further, by analyzing soil solution samples below the root system, you can verify deep percolation loss of nitrogen due to over-irrigation.

# Summary

Soil solution sampling by a farmer is a convenient, inexpensive and valuable practice. It allows for a much better informed management decision, which directly relates to crop production. In most cases, it can result in significant cost savings on the fertilizer bill. And finally, it can help the farmer mitigate the deep percolation of nitrates into the ground water supply that results from over-fertilization and over-irrigation.

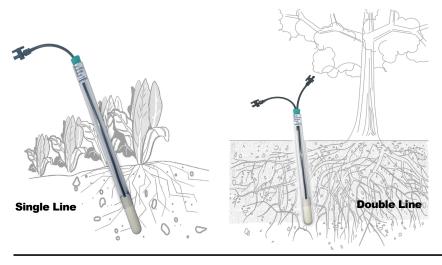
WARRANTY: The IRROMETER COMPANY warrants its products against defective workmanship or materials under normal use for one year from date of purchase. Defective parts will be replaced at no charge for either labor or parts if returned to the manufacturer during the warranty period. The seller's or manufacturer's only obligation shall be to replace the defective part and neither seller nor manufacturer shall be liable for any injury, loss or damage, direct or consequential, arising out of the use of or inability to use the product. This warranty does not protect against abuse, shipping damage, neglect, tampering or vandalism, freezing or other damage whether intentionally or inadvertently caused by the user.

# IRR METER®

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# **INSTALLATION AND OPERATING INSTRUCTIONS**

### **SSAT - Soil Solution Access Tube**



# Soil Water Sampling - Suction Lysimetry

This technology is a simple, inexpensive and non-destructive method of extracting a soil water sample for chemical analysis. The early work in this field was principally done in the area of salinity monitoring. Today, most of the application of this technology is in the area of nutrient management because of fertigation, the practice of introducing nutrients, principally nitrogen, in soluble form through a low volume (micro/drip/trickle) irrigation system. Many commercial agriculture producers involved in drip irrigated vegetable crops have adopted the use of this technology to monitor and control their nitrogen injection systems.

#### **Practical Field Use**

In the area of nutrient management, where drip systems are used by a farmer to "spoon feed" nitrogen to a growing crop, the technique of soil water sampling has proven to be a useful tool in manipulating the rate at which soluble nutrients are introduced through the irrigation water. Tissue analysis is still the baseline practice, but this can be time consuming and costly. And with the "spoon feeding" of nitrogen, sometimes done with every irrigation cycle, tissue analysis results are always lagging the actual event. Further, the technique of soil water sampling can help to monitor any excessive leaching of mobile nutrients beyond the root system, which would be caused by over-irrigation.

Certain cautions should be recognized when using this technique, particularly the fact that adequate sampling sites should be used to mitigate variations that may be encountered.

# When to Sample

The best advice is to extract your solution sample when the soil is at field capacity. This condition usually occurs anywhere from a few hours after, to the day following, an irrigation.

Soil moisture sensors (IRROMETERS OR WATERMARKS) placed in the active root system of the crop (and in the area wetted by the emitter) are the most convenient way to verify this condition of field capacity (15-25 centibars of soil water suction). At this time you are also sampling the soil solution, which contains the nitrogen introduced with the irrigation. And, with readily available soil water, the time required to draw the solution sample into the SSAT (Soil Solution Access Tube) is lessened. Attempting to draw a solution sample from soil that has dried beyond 35 centibars of suction can be frustrating, because in this condition the soil water environment contains a much greater volume of air.

# **Depths of Sampling**

In drip irrigation, the crop root system tends to be massed more towards the soil surface. This is the area where the crop extracts water and nutrients and thus the top 18" – 24" (45-60 cm) is the most critical for tree and vine crops. With shallower rooted crops such as vegetables, the top 12" (30 cm) is most significant. Mobile nutrients, such as nitrogen, are quickly transported beyond the root system if irrigation is excessive. And this fact emphasizes the need to monitor irrigation scheduling very carefully to prevent excessive leaching. Sampling of the soil water below the root system can be useful to spot heavy concentrations of nitrogen being lost to deep percolation. But the basic rule, for nutrition management, is to sample the active root system.

#### Installation

To install the SSAT (Soil Solution Access Tube) in the soil or container, make a 7/8" (22 mm) diameter access hole to the desired depth with a soil coring tool, IRROMETER Installation tool, a piece of 1/2" galvanized pipe, or 7/8" (22 mm) diameter solid rod. This provides for a snug fit (close contact with the soil). In hard or rocky soil, when coring a hole is difficult, a soil auger can be used to make the hole. The removed soil cuttings should be mixed with water to make a heavy slurry to "grout" the SSAT in place to establish good contact. Compact the soil at the surface around the top of the tube to avoid water channeling down the tube to the ceramic tip. Also, avoid installing in depressions, or low areas, which could retain standing water around the tube. Longer tubes can be used to sample from shorter depths to provide more vacuum capacity. They can be installed at an angle to avoid having too much of the tube exposed above ground to avoid possible accidental damage.

# Sample Collection

The SSAT should have the vacuum applied when the soil is at or near field capacity (10-25 centibars of soil water suction). Vacuum is applied with the hand vacuum pump (#1002-SSAT) twisted in to the valve (see figure 1) on the end of the tubing with the valve in the open position, handle parallel to the valve body/tubing. Pump until the vacuum is between 70 and 80 centibars on the pump gauge. Then close the valve, handle perpendicular to the body/tubing. (See figure 2) Then the pump can be twisted off to remove.







Figure 1

Figure 2

Figure 3

## Sample Collection (continued)

The vacuum will draw soil water into the SSAT through the ceramic tip. How long it takes to collect a sufficient sample size depends on how wet the soil is near the tip and how much vacuum is applied to the tube. On a very short SSAT (6" to 12", 15 to 30 cm) the plastic syringe (#DS-50CC) can be used to place a partial vacuum, which can be sufficient in very moist soils.

After sufficient time has passed to collect a significant volume of soil water (4 to 8 hours), the vacuum is released by opening the valve. Connect the plastic extraction syringe (DS-50CC) to the tubing by twisting it onto the valve and draw up on the syringe plunger to extract the soil solution for analysis. (**see figure 3**) If difficult to draw up, twist the rubber stopper loose to make the sample easier to extract.

### **DoubleLineSSAT**

The Double Line SSAT has two small tubing lines. One is the suction line, which goes all the way down into the tip and is for collecting the soil water sample. The other is short and is for pressurizing the tube to push the sample out. For depths 36" (90cm) and longer, the use of pressurized extraction is needed as the extraction syringe will not create enough lift to pull the sample out of the tube. Vacuum is applied with the hand vacuum pump (#1002-SSAT) twisted in to the valve on the end of the suction line tubing with the valve in the open position. Pump until the vacuum is between 70 and 80 centibars on the pump gauge. Then close the valve. Now the pump can be twisted off to remove.

For extraction with the Double Line SSAT, open both valves. The exposed end of the longer suction line should be placed in an open collection vessel, such as a sterile bottle or jar. The short or pressure line can be connected to an air pressure source. The extraction syringe (DS-50CC) is convenient to use for this. Pull the plunger all the way back, then attach it to the short or pressure line valve and slowly depress the plunger to push the collected sample out. To push more air in, detach the syringe before pulling the plunger back to repeat the procedure until the sample is completely extracted. Other air pressure sources, such as a bicycle tire pump or canister of pressurized air can also be used. As pressure is applied, the air forces the collected soil water to be extracted through the suction line and into the collection vessel. Only a small amount of pressure is needed, otherwise the rubber stopper can be pushed out of the tube. Be certain to completely evacuate the tube to prevent contaminating the next sample.