Sensing the electrical properties of roots: A review

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Abstract

Thorough knowledge of root system functioning is essential to understand the feedback loops between plants, soil, and climate. In situ characterization of root systems is challenging due to the inaccessibility of roots and the complexity of root zone processes. Electrical methods have been proposed to overcome these difficulties. Electrical conduction and polarization occur in and around roots, but the mechanisms are not yet fully understood. We review the potential and limitations of low-frequency electrical techniques for root zone investigation, discuss the mechanisms behind electrical conduction and polarization in the soil–root continuum, and address knowledge gaps. A range of electrical methods for root investigation is available. Reported methods using current injection in the plant stem to assess the extension of the root system lack robustness. Multi-electrode measurements are increasingly used to quantify root zone processes through soil moisture changes. They often neglect the influence of root biomass on the electrical signal, probably because it is yet to be well understood. Recent research highlights the potential of frequency-dependent impedance measurements. These methods target both surface and volumetric properties by activating and quantifying polarization mechanisms occurring at the root segment and cell scale at specific frequencies. The spectroscopic approach opens up a range of applications. Nevertheless, understanding electrical signatures at the field scale requires significant understanding of small-scale polarization and conduction mechanisms. Improved mechanistic soil–root electrical models, validated with small-scale electrical measurements on root systems, are necessary to make further progress in ramping up the precision and accuracy of multi-electrode tomographic techniques for root zone investigation.

Abbreviations: ARSA, absorbing root surface area; DC, direct current; ECF, extracellular fluids; EDL, electrical double layer; EIM, earth impedance method; EIS, electrical impedance spectroscopy; EIT, electrical impedance tomography; ERM, electrical resistance measurements; ERT, electrical resistivity tomography; ICF, intracellular fluids; MALM, mise-a-la-masse; PCA, principal component analysis; SIP, spectral induced polarization.

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1 | INTRODUCTION

The rhizosphere provides the necessary nutrients, water, and anchorage for plants and is one of the most active parts of soil in terms of nutrient and carbon cycling and maintaining biodiversity (De-la-Peña & Loyola-Vargas, 2014; Nguyen, 2009). It is home to numerous interactions between soil, plant, and microorganisms. Understanding rhizosphere processes is therefore crucial for sustainable agriculture and food production. Traditional methods of soil and root investigation such as excavation, soil cores, ingrowth bags, or mini-rhizotrons (Maeght, Rewald, & Pierret, 2013) provide insights into the structure, composition, and variability of the subsurface. However, they have clear limitations in terms of temporal and spatial resolution, mainly caused by the spatial heterogeneity of the root zone and its dynamics. Root growth, plant water and nutrient uptake, and transient boundary water fluxes create highly heterogeneous and dynamic patterns in root and soil properties that can be difficult to capture, even with a high-density network of point sensors (Amundson, Richter, Humphreys, Jobbágy, & Gaillardet, 2007; Jayawickreme, Jobbágy, & Jackson, 2014). In addition, intrusive sampling (e.g., rhizotubes) can interfere with the natural state of the system and may affect the processes being monitored, thereby potentially altering the experimental results.

Geophysical methods offer the possibility to infer properties and structures of the rhizosphere and pedosphere, as well as flow and transport processes at various spatial scales ranging from the single root to the field scale. Geophysical properties can be related to soil state variables (e.g., soil moisture content, salt concentration), soil properties (e.g., clay content, cation exchange capacity), and root properties (e.g., root mass, root surface area) as summarized by Vanderborght et al. (2013). Geophysical methods have been widely used in the past two decades to monitor moisture patterns associated with water flow (Deiana et al., 2007; Huisman, Sneyvangers, Bouten, & Heuvelink, 2002; Looms, Jensen, Binley, & Nielsen, 2008; Lu & Sabatier, 2009; Oberdörster, Vanderborght, Kemna, & Vereecken, 2010; Weihermüller, Huisman, Lambot, Herbst, & Vereecken, 2007; Zhou, Shimada, & Sato, 2001) and root water uptake (Beff, Günther, Vandoorne, Couvreur, & Javaux, 2013; Cassiani, Boaga, Vanella, Perri, & Consoli, 2015; Dick, Tetzlaff, Bradford, & Soulsby, 2018; Garré, Javaux, Vanderborght, Pages, & Vereecken, 2011; Jayawickreme, Van Dam, & Hynman, 2008; Mares, Barnard, Mao, Revil, & Singha, 2016; Michot et al., 2003; Srayeddin & Doussan, 2009; Vanella et al., 2018). The application of geophysical techniques in an agricultural context to study how agricultural production is affected by environmental variables (e.g., water availability, salinity) and agricultural management (e.g., impact of fertilizer and irrigation application), as well as to study fundamental soil–root interactions, is now referred to as “agrogeophysics” (Allred, Daniels, & Ehsani, 2008; Vereecken, Binley, Cassiani, Revil, & Titov, 2007).

Various agrogeophysical applications have been reported in the literature, ranging from mapping of soil compaction or determination of plow-pan depth (Besson, Cousin, Samouëlian, Boizard, & Richard, 2004; Doolittle, Sudduth, Kitchen, & Indorante, 1994; Kitchen, Sudduth, & Drummmond, 1999; Lu, Hickey, & Sabatier, 2004; Jonard et al., 2013) to the assessment of irrigation efficiency (Tresoldi et al., 2019). Soil mapping for precision agriculture and crop modeling has been realized using on-the-go electrical resistivity tomography (ERT; Andreelli et al., 2013; Rossi et al., 2013), electromagnetic induction (Brogi et al., 2019; Corwin & Lesch, 2003; Jaynes, Novak, Moorman, & Cambardella, 1995; Kachanoski, Wesenbeeck, & Gregorich, 1988; Robert et al., 1995; Sudduth, Drummmond, & Kitchen, 2001; von Hebel et al., 2018), and ground-penetrating radar (Freeland, Yoder, & Ammons, 1998; Ristic, Petrovacki, & Vrtunski, 2014; Yoder, Freeland, Ammons, & Leonard, 2001) Salinity mapping for whole farm planning is also feasible with geophysical methods (Williams, Walker, & Anderson, 2006). One of the key challenges in agrogeophysics is the noninvasive investigation of the spatial distribution of the root system (Amato et al., 2009; Konstantinovic, Wöckel, Schulze Lammers, Sachs, & Martinov, 2007) and its impact on soil state variables. Due to the vital role that root biomass and architecture play in plant breeding, drought tolerance, and carbon sequestration, measurements of root extent and distribution are necessary for proper understanding and modeling of plant growth, root water uptake, and the carbon balance of agricultural systems.

The aim of this review is to provide an overview of the applications of electrical geophysical methods to root investigation, to critically analyze recent accomplishments, and to identify remaining knowledge gaps. We limit ourselves to low-frequency electrical methods, from direct current (DC) to a few tens of kilohertz, due to their potential to investigate individual fine crop roots (diameter <1 mm). Recent reviews focused on imaging of root zone processes (Corona-Lopez, Sommer, Rolfe, Podd, & Rieve, 2019; Zhao et al., 2019), whereas others provided a short summary of the

Core Ideas
- Electrical methods offer a noninvasive approach for root investigation.
- Impedance measurements at different frequencies are preferable due to more information content.
- Interpretation of impedance spectra at the microscopic level needs to be considered.
FIGURE 1 Schematic illustration of (a) the plant cell, showing some of the organelles (vacuole, nucleus, and nuclear membranes), the cell wall, and the three-layer (protein-lipid-protein) cell membrane. The frequency-dependent current paths through plant tissues, showing (b) low-frequency current pathway and (c) high-frequency current pathway. (d) Description of counterion polarization mechanism in root segments occurring at low frequencies (Kessouri et al., 2019; Weigand & Kemna, 2019). The strength of the electrical double layer (EDL) polarizability depends on the composition and concentration of ions in the extracellular fluid (ECF) and electric potential distribution at the cell membrane (Kinraide, 2001; Kinraide & Wang, 2010; Wang et al., 2011). ICF, intracellular fluids

application of electrical impedance spectroscopy (EIS) to study various parts of plants (e.g., fruits, vegetables, and leaf water; Jocsac, Vegvari, & Vozary, 2019). To the best of our knowledge, this review is the first attempt to provide a comprehensive overview of all the active electrical methods (based on electric excitation) for root studies that are used by different research communities at various spatial scales. In particular, we reviewed all active electrical methods used to study the capability of roots to conduct electric current and to store electrical energy (in terms of charge separation).

This review is organized as follows. After a brief introduction to electrical conduction and polarization in plants, we will first present the different categories of measurement techniques and highlight their potentials and limitations. After that, we will discuss the common aspects, identify knowledge gaps for this active field of research, and provide suggestions for future research.

1.1 Electrical conduction and polarization in biological tissue

Biological tissues are made up of cells delimited by membranes. Cells contain tiny cellular structures that perform specific functions (organelles) and intracellular fluids (ICF), together called the cytoplasm (see Figure 1a). In plant cells, membranes are surrounded by cell walls mainly made of polysaccharides. The individual plant cells form a porous network called the apoplast, and the pore space between the cells is filled by extracellular fluids (ECF) (Raven, Johnson, Mason, Losos, & Singer, 2017). The membranes are made of
Table 1  Polarization mechanisms in biomaterials

<table>
<thead>
<tr>
<th>Polarization mechanism</th>
<th>Description</th>
<th>Frequency range</th>
</tr>
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<tbody>
<tr>
<td>Counterion or ionic polarization</td>
<td>Ionic diffusion in the electrical double layers adjacent to charged surfaces, such as cell membranes, creates polarization (Kao, 2004; Schwan &amp; Takashima, 1991).</td>
<td>α range &lt;20 kHz</td>
</tr>
<tr>
<td>Interfacial polarization (Maxwell–Wagner effect)</td>
<td>Restricted movement of ions in the intercellular space as a result of complex structural properties of biological tissue creates high-conductivity zones within the matrix of low conductivity material (e.g., cell interiors surrounded by cell membranes; Kao, 2004; Schwan &amp; Takashima, 1991)</td>
<td>β range (kHz–MHz)</td>
</tr>
<tr>
<td>Dipole or orientational polarization</td>
<td>Electrical dipoles can freely orient in response to the application of an electrical field and align themselves with the field (Kao, 2004)</td>
<td>γ range (MHz–GHz)</td>
</tr>
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phospholipids and proteins, which control water and nutrient entry into the cytoplasm due to their selective permeability. This helps to maintain the composition and concentration of the ICF, which is distinctly different from that of the ECF (Raven et al., 2017), as shown in Figure 1d. The radial transport of water and nutrients in roots takes place in the apoplast but also from cell to cell through plasmodesmata (i.e., tiny connections between cells, called the symplastic pathway; Couvreur et al., 2018). These two pathways also play an important role for electrical current flow.

Current conduction within a root depends on the resistance of the apoplast and the extracellular fluid, whereas all membranes and walls play an important role for the storage of electrical charges (polarization). Polarization occurs at the cell membrane interface because charged particles such as Na⁺, Ca²⁺, K⁺, and Cl⁻ ions and amino acids cannot diffuse across the cell membrane. Instead, they can only cross the membranes through ion pumps and ion channels (see the Glossary) whose opening and closing are regulated by the potential difference across the membrane (voltage). Electrical charges at the surface of the cell membrane lead to the formation of an electrical double layer (EDL), consisting of a layer of surface charges of the cell membrane and a layer of associated counterions (Figure 1d) (Kessouri et al., 2019; Kinraide, 2001; Kinraide & Wang, 2010, Weigand & Kemna, 2019). When applying an alternating current, the polarization strength depends on frequency, intensity, and duration of the injection, surface charge density of the cell membrane, transmembrane potential difference, ionic concentration, water content, tissue composition, and tissue health or structural heterogeneity (Ackmann, 1993; Bera, Nagaraju, & Lubineau, 2016; Repo, Cao, Silvennoinen, & Ozier-Lafontaine, 2012). Polarization is expected to occur at both the outer root surface (root–soil interface) and in the inner root system. Here, it is important to realize that the EDL of the outer root surface depends on the concentration of ions in the external fluid, whereas the EDL of the inner root system depends on the ionic composition of the cellular fluid (Weigand & Kemna, 2017, 2019). Polarization is assumed to occur in three frequency ranges, which are referred to as the α, β, and γ range (Prodan, Prodan, & Miller, 2008; Schwan, 1957, 1963, 1988). Table 1 summarizes the three frequency ranges and the specific polarization mechanisms associated to them (Repo et al., 2012; Schwan & Takashima, 1991).

At low frequencies, the voltage gated ion channels of the cell membranes are closed because a low-frequency electric field produces small changes in the membrane potential difference that are too small to significantly alter the properties of the ion channels (Mathie, Kennard, & Veale, 2003). In this case, the EDL prevents any passage of current through the cell membrane (i.e., the impedance of the cell membranes is so high that it does not allow current passage). Instead, the current passes through the apoplast (Bera, Bera, Kar, & Mondal, 2016; Repo et al., 2012), as shown in Figure 1b. In this low-frequency case, the total impedance will be mainly determined by the resistance of the extracellular fluid (Bera, Bera, et al., 2016; Repo et al., 2012); this is termed “counterion polarization” (Table 1). As the frequency increases, the applied electrical field at the outer surface of the membrane changes the transmembrane potential difference that regulates the gating of ion channels and ion fluxes across the cell membranes (Hille, 2001; Kinraide, 2001; Mathie et al., 2003). The opening of ion channels leads to decreased negativity of the membrane surface potential and an increased cation flux into the cell that results in a weaker EDL (Kessouri et al., 2019; Kinraide, 2001; Kinraide & Wang, 2010; Wang, Kinraide, Zhou, Kopittke, & Peijnenburg, 2011; Weigand & Kemna, 2019), and a lower polarizability of the EDL. At these high frequencies, current thus flows through the entire cell and crosses different interfaces (Figure 1c) such that the resulting total impedance will be a combination of the properties of the apoplast and the extracellular fluid, the cell membranes, the cytoplasm, and the ICF (Bera, Bera, et al., 2016; Repo et al., 2012). In this case, the polarization is considered to be interfacial (Table 1). In addition, the influx of cations into the cytoplasm due to concentration gradients that exist across the cell membrane when the ion channels open alters the composition and concentration of the ICF (Kessouri et al., 2019; Weigand & Kemna, 2019), which might explain the large polarization observed in roots at high frequencies (Ehosioke et al., 2018; Kessouri et al., 2019). However, more studies at the root scale,
across various species and growth conditions, are still needed to better understand high frequency polarization in roots. Conduction is assessed by measuring the resistance and then calculating resistivity or conductivity, whereas polarization is assessed by measuring the capacitance of the biological tissue. Both conduction and polarization can also be assessed simultaneously by measuring the electrical impedance of a biological tissue and then calculating effective material properties (complex resistivity or conductivity) by using a geometric factor $K$, which accounts for the geometric dimensions of the measurement (see the Glossary). In this case, the real part is associated with electrical conduction, whereas the imaginary part is associated with electrical polarization.

1.2 Overview of electrical methods for root system investigation

These methods target either electrical resistance, capacitance, or impedance and are classified into eight categories, in order of appearance in the following sections: electrical resistance measurements (ERM), ERT, mise-a-la-masse (MALM), electrical capacitance measurements (ECM), spectral induced polarization (SIP), earth impedance method (EIM), electrical impedance spectroscopy (EIS), and electrical impedance tomography (EIT). Some of these methods share the same physical principles, but they differ in their application scale (root scale, plant scale, and plot scale), the frequency of the electric field (e.g., single frequency or multiple frequencies), and the amount and position of electrodes, as illustrated in Figure 2.

Methods at the root and plant scales have mainly been used in fundamental studies or to obtain information on key parameters that can be used to interpret results from measurements at the field scale. They are also useful to determine if sufficient contrast in the electrical signature can be expected in field measurements. In this review, we will give an overview of these different measurement techniques coming from various scientific communities and will discuss their functioning and potential for root system investigation.

2 METHODS TARGETING ELECTRICAL RESISTIVITY

2.1 ERM

Some studies have used relatively simple ERM to investigate plant roots (Anderson & Higinbotham, 1976; Cao, Repo, Silvennoinen, Lehto, & Pelkonen, 2010). An ERM setup consists of a multimeter and two electrodes connected along a
FIGURE 3 Application of electrical resistance measurement (ERM) for root studies. (a) Resistivity variation along a segment of willow roots, segments with same letter code show no significant difference (Reproduced from Cao et al., 2010, with permission of Oxford University Press). (b) Resistivity variation across species (Ehosioke et al., 2017).

single root segment (root scale) or with one electrode in the plant stem and the second one in the growth medium (soil and hydroponics; plant scale). The electrodes are simultaneously used for current injection and potential measurements. The main output of such measurements is the electrical resistance or resistivity of the studied system provided that an appropriate geometrical factor can be identified. The experimental setup is simple and typically operates at the centimeter scale. The understanding of the electrical response of single root segments is important since it is a first step towards discriminating root electrical signals from those of other soil properties and state variables, especially in combination with modeling (Rao et al., 2019).

Anderson and Higinbotham (1976) pioneered the study of electrical resistance of root segments by measuring longitudinal electrical resistance along 2-cm segments of maize (Zea mays L.) roots cut at varying distances from the root apex. The older root segments further away from the apex had a greater electrical resistance than the younger segments, except for the section just before the apex, which showed the greatest resistance. They attributed this to the proportionately large number of cells in the meristem and in the early stages of the elongation zone. However, only the resistance was measured, and the corresponding root diameters were not reported. Therefore, it is not possible to discriminate possible geometrical factors (i.e., differences in root diameter) from inherent electrical root properties, which compromises the interpretation of these measurements.

Cao et al. (2010) used ERM in a hydroponic system to study root morphology and the variation of axial resistivity along a single root segment of willow (Salix schwerinii E. Wolf). They calculated axial resistivity at varying distances from the apex (Figure 3a). They found the highest resistivity just before the root tip and observed a gradual reduction in resistivity towards the root collar. The high axial resistivity next to the root tip was attributed to the presence of a hydraulically isolated segment with immature metaxylem and mature protoxylem (Frensch & Steudle, 1989). The root tip itself showed the lowest resistivity, which was attributed to underdeveloped cell walls with a very permeable membrane that allow fast movement of solute and water between the root tip and soil (Frensch & Steudle, 1989).

Cao et al. (2010) also measured the resistance of the stem–root system at different immersion depths of the root system in a hydroponic solution. Afterwards, the roots were cut off and the resistance of the stem was measured. The excised roots were scanned to assess the surface area, and the number of lateral roots was obtained using image analysis (WinRhizo). They found that the measured resistance correlated significantly with root surface area ($r^2 = .75$) and the root length ($r^2 = .63$), which is expected and linked to the geometric factor, and to a lesser extent with the number of lateral roots ($r^2 = .56$). This suggests that ERM could be useful to study root morphology. The submerged part of the root system played a negligible role, which is probably due to current leakage at the stem–solution interface. They also found that the average resistivity of the roots (10.5 Ωm) was much smaller than that of the stem (108.3 Ωm), which was attributed to the anatomical differences between the root and stem. For example, the vascular tissue is thicker and more
Recently, Ehosioke et al. (2017) used ERM measurements to determine the electrical resistivity of root segments of three different plant species: maize, ryegrass (Lolium perenne L.), and Brachypodium. Their results showed that electrical resistivity of roots varied for different species (Figure 3b). It was found that the variation in electrical resistivity could be linked to the structural variation and anatomical differences between the species.

A major limitation of ERM is that it is hard to obtain knowledge on the structures that are contributing most to the electrical conduction. In addition, roots were excised before ERM measurements in some studies (Anderson & Higinbotham, 1976; Cao et al., 2010). In this case, the cut surface in contact with the nutrient solution may introduce a bias because of an increased absorbing surface area.

### 2.2 ERT

Electrical resistivity tomography, also called DC resistivity imaging (DCR) or electrical resistivity imaging (ERI), determines the distribution of electrical resistivity by performing a set of resistance measurements on the ground surface and/or in boreholes. Measurements are performed by injecting current via two electrodes, and the resulting voltage difference is measured at two other electrodes using various combinations of current and potential electrodes along a transect or grid. In order to determine the resistivity of the subsurface, a geophysical inversion of the measured resistances must be performed. The obtained resistivity distributions are typically presented as tomograms. Since the inverse problem is ill posed, the obtained resistivity model is nonunique and typically represents a smoothed representation of the actual resistivity distribution.

Electrical resistivity tomography is interesting for root system investigations because of its high two- or three-dimensional spatial coverage, acquisition speed, minimally invasive nature, and cost effectiveness. Some ERT studies focused on root biomass estimation, whereas others focused on physiological processes such as water and nutrient uptake by the root system, but these are mainly interpretation perspectives rather than a prior methodological difference, as both aspects can be captured by time-lapse ERT measurements.

Figure 4 compares results from several studies that have directly correlated bulk resistivity with root biomass with varying success (Amato et al., 2008, 2009; Rossi et al., 2011; Paglis, 2013). Bulk soil resistivity is known to be strongly influenced by soil moisture content, porosity, mineralogy, temperature, and salinity level and only to a lesser extent by roots. Thus, the contrast between roots and soil depends highly on soil type and saturation state (Rao et al., 2019) and is therefore varying with time. In addition, fine roots actively take up water and nutrients and release different exudates. These processes also affect resistivity at several temporal scales, ranging from daily (night vs. day, sunny vs. cloudy days) to seasonal (growth period vs. winter or drought season).
Based on these considerations, it is clear that the relationships shown in Figure 4 are site specific and without generality. Overall, ERT may lack direct sensitivity to root properties due to the variations in the contrast between bulk soil and root electrical resistivity and the relatively low volume fraction of roots. Using resistivity as a predictor for root biomass without taking these factors into account is therefore highly uncertain (Rao, Lesparre, Orozco, Wagner, & Javaux, 2020).

Electrical resistivity tomography has also been used to investigate physiological processes such as water and nutrient uptake, which is promising given the high sensitivity to water content in the root zone. A range of studies have reported areas of contrasting resistivity (in time and space) that were attributed to root water uptake processes beneath herbaceous plants (Beff et al., 2013; Garré, Günther, Diels, & Vanderborght, 2012; Michot et al., 2003; Moreno, Arnon-Zur, & Furman, 2015; Panissod, Michot, Benderitter, & Tabbagh, 2001), and some shrubs and trees (Hagrey, Meissner, Werban, Rabbel, & Ismaeil, 2004; Hagrey & Michaelson, 2002; Jayawickreme, Van Dam, & Hyndman, 2010; Katul, Todd, Pataki, Kabala, & Oren, 1997; Koumanov, Hopmans, & Schwanik, 2006; Robinson, Slater, & Schäfer, 2012). Electrical resistivity tomography has also been used in orchards to study soil moisture dynamics and irrigation patterns near cash crops such as fruit trees (Lazzari et al., 2008; Loperte et al., 2006). These studies found an increasing resistivity in the root zone due to drying through root water uptake processes, but some root zones show lower resistivity than the background as shown in Figure 5 (Lazzari et al., 2008; Petersen & Al Hagrey, 2009; Rodriguez-Robles, Arredondo, Huber-Sannwald, Ramos-Leal, & Yépez, 2017; Werban, Attia al Hagrey, & Rabbel, 2008). The low resistivity of fine roots and high resistivity of tree roots reported in the studies discussed above is an indication that roots are a contributing factor, although masked by other factors, as discussed above. The effect of root biomass can only be clearly distinguished with additional independent measurements.

Many examples exist where time-lapse ERT has been used to quantify water and solute fluxes in a range of agricultural systems, including maize (Beff et al., 2013; Michot et al., 2003), barley (Hordeum vulgare L.; Garré et al., 2011), mixed cropping (Garré et al., 2013), and orchards (Cassiani et al., 2015; Moreno et al., 2015; Vanella et al., 2018). Interestingly, none of these studies accounted for the potential effect of root biomass on the measured resistivity. This is most probably because the sensitivity of ERT to soil moisture is larger than to root biomass in most cases, especially when looking at time-lapse differences. Nevertheless, Rao et al. (2019) already warned that this might not always be the case. Using coupled electrical and root water uptake models, they showed that the bias in soil moisture estimates may be up to 50% when roots are not properly accounted for. Although ERT is less sensitive to roots than other system properties (soil texture, porosity, stone fraction, mineralogy) and state variables (soil temperature, fluid conductivity, soil moisture), the influence of roots can be consistent. Therefore, there is a need for auxiliary data to separate root effects from other properties and variables.

### 2.3 MALLM method

In the MALLM method (Parasnis, 1967), an electric current is injected into a conductive body and the resulting voltage is measured at the ground surface located at certain distances around the body. The resulting voltage contours are then used to delineate the extent of the conductive body. In root studies, the conductive body is the plant stem–root system. This setup is based on the assumption that the current injected into a tree stem will be released into the soil through the root system, but only at the root–soil interface where roots absorb water and minerals from the soil.
Mary et al. (2018) used the MALM method jointly with ERT to characterize vine plant root water uptake at the plant scale. They injected current directly into the tree stem in one experiment and directly into the soil close to the stem in another experiment. It was observed that current injection into the stem produced very different voltage distributions in the soil compared with an injection into the soil. They concluded that current injected into the stem is redistributed in the soil by the root network. In a follow-up study, Mary, Vanella, Consoli, and Cassiani (2019) used MALM and ERT measurements to assess the root extent of citrus trees in an orchard under two different irrigation regimes (full and partial irrigation). The MALM measurements were used to map the locations of the active roots, whereas ERT was used to obtain quantitative information about the uptake of water by the roots. It was reported that the distribution of current flow for a stem injection was much different from that of a current injection into the soil in the case of partial irrigation. In contrast, no differences were observed in the case of full irrigation. This difference in behavior was attributed to differences in the resistivity contrast in the two cases. For partial irrigation, roots are likely more conductive than the surrounding soil, and the current flows preferentially through the roots. In the case of full irrigation, the surrounding soil is more conductive than the root, and the current is expected to leave the root system near the stem–soil interface, as also suggested by Urban, Bequet, and Mainiero (2011) and Peruzzo et al. (2020). Thus, the success of the MALM method depends on the contrast between the electrical properties of the root and that of the soil and on the contact resistance at the interface between the two. More studies are required to better understand root electrical properties, which are essential to improve the interpretation of MALM measurements.

2.4 Measurements focusing on electrical capacitance

2.4.1 ECM on single plants in soil

Electrical capacitance measurements are commonly used to study root systems at the plant scale (Chloupek, 1972; Dalton, 1995). Root capacitance is measured with an LCR meter (i.e., equipment that measures inductance, capacitance, and resistance of an electronic component) and two electrodes. One electrode is inserted into the stem, while the other is inserted into the growing medium. The capacitance is the amount of electric charge stored by the root system for a given electric potential (in farads). It quantifies polarization processes (stored charges) taking place in the root tissue upon current flow (Dalton, 1995; Rajkai, Végh, & Nacsá, 2002; Repo, Zhang, Ryypö, & Rikala, 2000). Table 3 summarizes ECM applications reviewed below according to their target plant property: root biomass, morphology, or physiology.

We reviewed 138 studies that reported correlations between the measured capacitance ($C_{\text{root}}$) and root dry mass ($R_{\text{dm}}$), whereas another 14 studies reported correlations between $C_{\text{root}}$ and root fresh mass ($R_{\text{fm}}$). The average correlation between $C_{\text{root}}$ and $R_{\text{dm}}$ was .77, whereas the average correlation between $C_{\text{root}}$ and $R_{\text{fm}}$ was .74. This suggests that ECM may be suitable for root biomass determination (see Figure 6). However, the range of correlation coefficients reported for $R_{\text{dm}}$ (.30–.98) and $R_{\text{fm}}$ (.51–.94) is large. This is due to other factors besides root mass that affect root capacitance, such as plant species (Aulen & Shipley, 2012), genotypes (Mcbride, Candido, & Ferguson, 2008), growing medium (Cseresnýés et al., 2017), and soil water status (Cseresnýés et al., 2018).

Based on 14 studies relating ECM to root morphology, there appears to be a correlation between $C_{\text{root}}$ and root length or root surface area, with correlation coefficients ranging from .56 to .94 (average $R = .79$). Ellis, Murray, and Kavalieris (2013) tested the use of a four-electrode setup for ECM, which eliminated the high contact resistance present in the classical two-electrode setup. They observed very low correlations of $C_{\text{root}}$ with root mass, area, and length ($R = .21–.31$). It was therefore concluded that electrical capacitance was a poor predictor of root dimensions in their analysis. Since $C_{\text{root}}$ depends on tissue density and thus the position of stem electrodes, the setup of Ellis, Murray, and Kavalieris (2013) involved two stem electrodes. In particular, the second stem electrode (a potential electrode) is closer to the soil surface and thus reduces the influence of tissue density. This might explain the poor correlation they observed.

Investigations of root development with time using ECM showed that there are important temporal variations in the measured capacitance (Cseresnýés et al., 2018; Cseresnýés, Rajkai, & Takács, 2016; Cseresnýés, Takács, Füzy, & Rajkai, 2014; Cseresnýés, Takács, Füzy, Végh, & Lehoczyk, 2016; Dalton, 1995). Cseresnýés et al. (2014) found that root water uptake and $C_{\text{root}}$ increased from the emergence of a seedling to the beginning of flowering and then decreased continuously during the fruit setting for cucumber (Cucumis sativus L.) and bean (Phaseolus vulgaris L.). Similarly, ECM presented in Cseresnýés et al. (2018) identified peak activity with a capacitance value of 96.4 nF at a period that coincided with the completion of flowering 92 d after sowing. Afterwards, the capacitance decreased to 6.3 nF at Day 161, which coincided with the phenological stage of senescence. Senescence is associated with suberization and cell death, which is assumed to result in a lower capacitance. Cseresnýés et al. (2018) argued that ECM is more reliable for the estimation of root activity than for the estimation of root mass, length, and surface area. We tend to agree with this argument because root activity is a contributing factor in root biomass and morphology assessment using ECM, since most of these
measurements were performed on active plants. The large disparity among studies could be a reflection of the physiological state of the plant at the time of measurements.

Electrical capacitance measurements are usually performed using an LCR meter. Since capacitance is only clearly defined for an ideal capacitor, interpretation of the measured values relies on the selection of an appropriate equivalent circuit model. Dalton (1995) proposed a linear model assuming that roots behave as cylindrical capacitors and that their capacitances can be added together as if wired in parallel, suggesting that capacitance depends on the submerged root mass. Dietrich, Bengough, Jones, and White (2012) have shown that this model is too simplified and proposed an alternative equivalent circuit model (Figure 7) in which (a) the capacitance of the growth medium is much larger than the capacitance of the plant tissue, (b) the capacitances of tissues along an unbranched root can be considered as connected in series, (c) the capacitances of multiple unbranched roots comprising the whole root system act in parallel but reduce to the equivalent of a single capacitor, and (d) the capacitances of individual roots are directly proportional to their cross-sectional area or circumference. The model of Dietrich et al. (2012) indicates that the total capacitance is dominated by the aboveground part of the plant between the stem electrode and the growth medium. This may explain the wide range of results that have been obtained with ECM, since the distance between stem electrode and growth medium was not standardized in these studies.

An important point to realize is that soil water content affects capacitance measurements (a principle commonly used in the design of soil moisture sensors). However, ECM cannot distinguish the plant from the soil response. Many studies have circumvented this problem by making ECM on plants grown in hydroponic conditions. In situ monitoring of root activity may be feasible for ECM, but only under controlled conditions and after careful calibration. Such calibration is required, since capacitance is known to be influenced by species (Aulen & Shipley, 2012), genotypes (Mcbride et al., 2008), tissue density (Ellis, Murray, & Kavalieris, 2013; Ellis, Murray, et al., 2013), electrode type (Kormanek, Głąb, & Klimek-Kopyra, 2016), and growing media (Cseresnyés et al., 2017).

2.5 Electrical impedance as a comprehensive target property

2.5.1 EIM for single plants in soil

The EIM has been used to study physiological characteristics of root systems in situ, such as the absorbing root surface area (ARSA, Table 2). The setup relies on four electrodes, two current electrodes $C_1$ and $C_2$ for current injection, and two potential electrodes $P_1$ and $P_2$ to measure the resulting voltage. The current electrode $C_1$ is hammered into the stem, whereas $C_2$ is placed into the soil at a distance of about 10–20 m from the stem (depending on the tree size). The potential electrode $P_1$ is fixed into the root collar, whereas $P_2$ is inserted into the soil and moved repeatedly along the line connecting $C_1$ and $C_2$ (see Figure 8a). The measurements are used to calculate the grounding impedance $I_g$ and to estimate ARSA (Aubrecht, Stanek, & Koller, 2006; Čermák, Ulrich, Stanek, Koller, &...
Aubrecht, 2006) using

\[
\text{ARSA} = \rho \frac{L_{\text{eff}}}{I_g}
\]  

(1)

Where \(I_g\) is the grounding impedance, \(L_{\text{eff}}\) is the distance to the stem, and \(\rho\) is the wood resistivity. The wood resistivity is measured with two or four electrodes (depending on the tree size) while assuming that the tree is a cylinder (Aubrecht et al., 2006).

Aubrecht et al. (2006) described the theoretical basis of the EIM. The applicability of this method is based on the following crucial assumptions: (a) the xylem resistance is negligible and is not considered in the calculation of ARSA, (b) there is no radial conduction in woody or suberized root, which implies that electrical current is expected to flow through the entire root system and exit via the fine absorbing roots, (c) every absorptive part of a root branch is expected to contribute equally to the overall conductance, (d) root branches are expected to be electrically discrete resistors and tree root segments to be measured separately, (e) the ARSA is predominantly determined by the contact impedance between the root and the soil, and finally, (f) the soil impedance is a negligible part of the grounding impedance of a tree. Urban et al. (2011) assessed the applicability of the EIM to investigate tree root systems by testing these assumptions in several experiments, but they could not confirm that the assumptions listed above apply. Instead, their results clearly showed that a tree–root–soil continuum is a serial circuit where the xylem and soil impedance dominate the contact impedance. In particular, they showed that xylem electrical resistance is an important component of \(I_g\) and that \(I_g\) increases significantly with diameter at breast height (DBH) (Figure 8b). Therefore, the observed relationships between ARSA and DBH were site and soil specific. They concluded that the EIM cannot reliably assess the ARSA of distal fine roots because most of the charge carriers exit the root system in the proximal parts of the root–soil interface. Čermák et al. (2013) compared...
results from the EIM with those obtained from scanning and microscopy of Norway spruce \textit{[Picea abies (L.) Karst.]} at two different sites. The root absorptive index value obtained from EIM (0.7 m$^2$ m$^{-2}$) was the lowest for both sites compared with the 2.2 and 1.2 m$^2$ m$^{-2}$ obtained for the scan and microscopy, respectively. This suggests that ARSA obtained by the EIM is not reliable and thus agrees with the conclusion of Urban et al. (2011). In the field, various determinants of soil and xylem resistivity exist: for example, soil texture and porosity (Chloupek, 1972), root density (Amato et al., 2008), moisture of the soil and xylem ( Hagrey, 2007; Zenone et al., 2008), and also cell anatomy (van Beem, Smith, & Zobel, 1998; Zhang, Willison, & Willison, 1991). These factors vary in space and time, causing $I_g$ and $L_{	ext{eff}}$ to vary independently of ARSA, which makes it difficult to determine ARSA from the EIM alone.

### 2.5.2 The value of measurements at multiple frequencies

Measurements at multiple frequencies offer more potential for root investigation because the use of different frequencies allows to separate processes that have different time or spatial scales. Methods using multiple frequencies are classified here as EIS, SIP, and EIT.

### 2.5.3 EIS and SIP

Electrical impedance spectroscopy determines the impedance of the soil–plant continuum across a range of frequencies (hertz to megahertz). This method has been used in many studies in plant science and medicine (Bera, Bera, et al., 2016; Bera, Nagaraju, & Lubineau, 2016; Coster, Chilcott, & Coster, 1996; Hayden, Moyse, Calder, Crawford, & Fensom, 1969; Inaba, Manabe, Tsuji, & Lwamuto, 1995; Klauke et al., 2005; Lin, Chen, & Chen, 2012; Macdonald, 1992; Shrivastava, Barde, Mishra, & Phadke, 2014a, 2014b). For root studies, EIS has been used to assess the morphological and physiological properties of roots such as root growth (Ozier-Lafontaine & Bajazet, 2005; Repo, Laukanen, & Silvennoinen, 2005), estimation of root system size (Cao, Repo, Silvennoinen, Lehto, & Pelkonen, 2011), and mycorrhizal colonization of roots (Cseresnyés, Takács, Végh, Antón, & Rajkai, 2013; Repo, Korhonen, Laukanen, Lehto, & Silvennoinen, 2014; Repo, Korhonen, Lehto, & Silvennoinen, 2016), as summarized in Table 4. An EIS setup consists of an impedance analyzer and two or three electrodes; one is typically inserted into the root collar, while the other(s) is fixed into the growth substrate. The electrode can be a stainless steel needle (Repo et al., 2014), a tension clamp (Cseresnyés, Takács, et al., 2013; Cseresnyés, Rajkai, & Vozáry, 2013),

<table>
<thead>
<tr>
<th>Method*</th>
<th>Application category</th>
<th>Specific root property</th>
</tr>
</thead>
<tbody>
<tr>
<td>ERM</td>
<td>Morphology</td>
<td>Surface area, length, and number of lateral roots</td>
</tr>
<tr>
<td>ECM</td>
<td>Anatomy</td>
<td>Structural variation along a single segment</td>
</tr>
<tr>
<td></td>
<td>Biomass estimation</td>
<td>Dry mass and fresh mass</td>
</tr>
<tr>
<td></td>
<td>Morphology</td>
<td>Length and surface area</td>
</tr>
<tr>
<td>EIM</td>
<td>Physiology</td>
<td>Root activity (water and nutrient uptake)</td>
</tr>
<tr>
<td>MALM</td>
<td>Physiology</td>
<td>Location of active roots</td>
</tr>
<tr>
<td>EIS/SIP</td>
<td>Anatomy</td>
<td>Structural change in roots with age</td>
</tr>
<tr>
<td></td>
<td>Biomass estimation</td>
<td>Resistivity and polarization contrast between root and soil</td>
</tr>
<tr>
<td>ERT</td>
<td>Biomass estimation</td>
<td>Root mass density</td>
</tr>
<tr>
<td>EIT</td>
<td>Anatomy</td>
<td>Structural change in roots</td>
</tr>
<tr>
<td></td>
<td>Physiology</td>
<td>Root decay due to nutrient deprivation</td>
</tr>
<tr>
<td></td>
<td>Biomass estimation</td>
<td>Resistivity and polarization contrast between root and soil</td>
</tr>
<tr>
<td></td>
<td>Morphology</td>
<td>Root spatial extent</td>
</tr>
</tbody>
</table>

*ERM, electrical resistance measurements; ECM, electrical capacitance measurements; EIM, earth impedance method; MALM, mise-a-la-masse; EIS, electrical impedance spectroscopy; SIP, spectral induced polarization; ERT, electrical resistivity tomography; EIT, electrical impedance tomography.
<table>
<thead>
<tr>
<th>Reference</th>
<th>Plant species</th>
<th>Root properties</th>
<th>Correlation</th>
<th>Growth medium</th>
<th>Observations</th>
</tr>
</thead>
<tbody>
<tr>
<td>Chloupek, 1972</td>
<td><em>Zea mays</em></td>
<td>Dry mass</td>
<td>.728</td>
<td>Sand</td>
<td></td>
</tr>
<tr>
<td></td>
<td><em>Helianthus annuus</em></td>
<td>Fresh mass</td>
<td>.692</td>
<td>Clay soil</td>
<td></td>
</tr>
<tr>
<td>Chloupek, 1977</td>
<td><em>Helianthus annuus</em></td>
<td>Fresh mass</td>
<td>.54, .566</td>
<td>Sand</td>
<td></td>
</tr>
<tr>
<td></td>
<td><em>Daucus carota</em></td>
<td>Fresh mass</td>
<td>.514</td>
<td>Loam soil</td>
<td></td>
</tr>
<tr>
<td>Kendall et al., 1982</td>
<td><em>Trifolium pretense</em></td>
<td>Dry mass</td>
<td>.672</td>
<td>Hydroponics</td>
<td></td>
</tr>
<tr>
<td></td>
<td><em>Medicago sativa</em></td>
<td>Dry mass</td>
<td>.50</td>
<td>Silt loam</td>
<td></td>
</tr>
<tr>
<td>Dalton, 1995</td>
<td><em>Solanum lycopersicum</em></td>
<td>Dry mass</td>
<td>.77</td>
<td>Hydroponics</td>
<td>Linear relation</td>
</tr>
<tr>
<td>van Beem et al., 1998</td>
<td><em>Zea mays</em></td>
<td>Fresh mass</td>
<td>.73</td>
<td>Vermiculite</td>
<td></td>
</tr>
<tr>
<td></td>
<td><em>Zea mays</em></td>
<td>Fresh mass</td>
<td>.53</td>
<td>Loam (field)</td>
<td></td>
</tr>
<tr>
<td>Preston, McBride, Bryan, &amp; Candido, 2004</td>
<td><em>Populus deltoids</em></td>
<td>Fresh mass</td>
<td>.866</td>
<td>Potting soil</td>
<td></td>
</tr>
<tr>
<td>Rajkai, Vég, &amp; Nacs, 2005</td>
<td><em>Helianthus annuus</em></td>
<td>Fresh mass</td>
<td>.832, .921</td>
<td>Sandy soil</td>
<td>Needle &amp; clamp electrodes</td>
</tr>
<tr>
<td>Ozier-Lafontaine &amp; Bajazet, 2005</td>
<td><em>Solanum lycopersicum</em></td>
<td>Dry mass</td>
<td>.987, .829</td>
<td>Hydroponics &amp; clay loam</td>
<td>Linear and exponential relation</td>
</tr>
<tr>
<td></td>
<td><em>Amaranthus tricolor</em></td>
<td>Fresh mass</td>
<td>.937</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mcbride et al., 2008</td>
<td><em>Zea mays</em> (inbreeds and hybrids)</td>
<td>Dry mass</td>
<td>.779, .647, .761, .823, .364, .846</td>
<td>Hydroponics + Turface</td>
<td>Different genotypes showed unique capacitance relation with dry mass</td>
</tr>
<tr>
<td>Bengough et al., 2009</td>
<td><em>Triticum aestivum</em></td>
<td>Dry mass</td>
<td>.753</td>
<td>Gravel–sand mix</td>
<td></td>
</tr>
<tr>
<td>Tsukahara, Yamane, Yamaki, &amp; Honjo, 2009</td>
<td><em>Prunus persica</em></td>
<td>Fresh mass</td>
<td>.897</td>
<td>Soil (field)</td>
<td></td>
</tr>
<tr>
<td></td>
<td><em>Prunus persica</em></td>
<td>Dry mass</td>
<td>.806</td>
<td>Soil (potsoil)</td>
<td></td>
</tr>
<tr>
<td>Chloupek, Dostál, Středa, Psota, &amp; Dvořáčková, 2010</td>
<td><em>Daucus carota</em></td>
<td>Fresh mass</td>
<td>.525</td>
<td>Soil (field)</td>
<td></td>
</tr>
<tr>
<td>Pitre et al., 2010</td>
<td><em>Salix viminalis</em></td>
<td>Dry mass</td>
<td>.81</td>
<td>Soil (potsoil)</td>
<td></td>
</tr>
<tr>
<td></td>
<td><em>Salix schwerinii</em></td>
<td>Dry mass</td>
<td>.49</td>
<td>Sandy soil (field)</td>
<td></td>
</tr>
<tr>
<td>Aulen &amp; Shipley, 2012</td>
<td>Herbaceous species (10 different plants)</td>
<td>Dry mass</td>
<td>.30</td>
<td>Compost</td>
<td>Requires species-specific calibration, regression strength not strong</td>
</tr>
<tr>
<td>Dietrich et al., 2012</td>
<td><em>Hordeum vulgare</em></td>
<td>Fresh mass</td>
<td>.869</td>
<td>Hydroponics</td>
<td>No linear relation when roots were raised from solution Proposed a new model in place of Dalton model</td>
</tr>
<tr>
<td></td>
<td></td>
<td>C/s area</td>
<td>.806, .771</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Dietrich, Bengough, Jones, &amp; White, 2013</td>
<td><em>Triticum aestivum</em></td>
<td>Dry mass</td>
<td>.753</td>
<td>Sand</td>
<td>Confirmed the model proposed by Dietrich et al., 2012 EC measured in wet substrate ≠ direct assessment of root mass</td>
</tr>
<tr>
<td>Ellis, Murray, &amp; Kavalieris 2013</td>
<td><em>Vicia faba</em></td>
<td>Root length</td>
<td>.56</td>
<td>Sand</td>
<td>EC is a poor predictor of root size and depends on tissue density</td>
</tr>
<tr>
<td>Ellis, Murray, et al., 2013</td>
<td>Various</td>
<td>Root length</td>
<td>.94</td>
<td>Moist clay-loam (plantation)</td>
<td>Position of stem electrode affected EC values</td>
</tr>
</tbody>
</table>

(Continues)
<table>
<thead>
<tr>
<th>Reference</th>
<th>Plant species</th>
<th>Root properties</th>
<th>Correlation</th>
<th>Growth medium</th>
<th>Observations</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cseresnyés, Takács, et al., 2013</td>
<td>Zea mays</td>
<td>Dry mass Length Surface area</td>
<td>.9230 .9183 .9367</td>
<td>Pumice</td>
<td>Higher values and stronger correlation found in mycorrhiza-colonized roots</td>
</tr>
<tr>
<td>Postic &amp; Doussan, 2016</td>
<td>Triticum turgidum</td>
<td>Dry mass</td>
<td>.787</td>
<td>Loam</td>
<td>Recommended more than two-electrode setup and clamp electrode for stem</td>
</tr>
<tr>
<td>Cseresnyés, Rajkai, &amp; Takács, 2016</td>
<td>Glycine max</td>
<td>Dry mass</td>
<td>.844, .895, .936</td>
<td>Pumice</td>
<td>EC was affected by drought, EC increased with age (DAP), peaks during flowering and decreased after</td>
</tr>
<tr>
<td>Carlson &amp; Smart, 2016</td>
<td>Salix salicaceae</td>
<td>Dry mass</td>
<td>.88</td>
<td>Peat moss</td>
<td>Root dry mass relationship with EC is strongly linked with stem dry weight</td>
</tr>
<tr>
<td>Kormanek et al., 2016</td>
<td>Fagus sylvatica</td>
<td>Dry mass</td>
<td>.502 (cylinder), .747 (rect. el.)</td>
<td>Loam (forest nursery)</td>
<td>Rectangular electrode correlated better with all parameters than cylindrical electrode Proposed a two-dielectric model in place of Dalton model</td>
</tr>
<tr>
<td>Cseresnyés, Takács, Fuzy, Végh, &amp; Lehoczky, 2016</td>
<td>Zea mays A. theophrasti</td>
<td>Dry mass</td>
<td>.901 .954</td>
<td>Arenosol</td>
<td>EC increased with age, with peaks around 27–31 DAP, and then decreased.</td>
</tr>
<tr>
<td>Cseresnyés et al., 2017</td>
<td>Cucumis sativus</td>
<td>Dry mass</td>
<td>.688</td>
<td>Arenosol</td>
<td>Capacitive effect of the growth media increased with the complexity of the substrate Correction for dissipation factor (capacitive effect) improved correlation</td>
</tr>
<tr>
<td></td>
<td>Triticum aestivum</td>
<td>Dry mass</td>
<td>.879</td>
<td>Pumice</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Surface area</td>
<td>.683</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Length</td>
<td>.726</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Total root area</td>
<td>.641 (cylinder), .818 (rect. el.)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Total root length</td>
<td>.688 (cylinder), .802 (rect. el.)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cseresnyés et al., 2018</td>
<td>Zea mays</td>
<td>Dry mass</td>
<td>.882–.911</td>
<td>Chernozem soil</td>
<td>Root activity increased until flowering and decreased during maturity</td>
</tr>
<tr>
<td></td>
<td>Glycine max</td>
<td>Dry mass</td>
<td>.831–.874</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Note. C/s, cross-section; EC, electrical conductivity; DAP, days after planting; rect. el., rectangular electrode.

or a nonpolarizable Ag/AgCl electrode (Ozier-lafontaine & Bajazet, 2005).

Macdonald (1987) described two distinct ways to analyze the impedance spectra of a particular system. At a macroscopic level, the impedance spectra of the system can be analyzed using equivalent circuit models. Alternatively, interpretation can be done at a microscopic level by analyzing the impedance spectra in terms of the physicochemical properties of the system. Whereas the analysis of root impedance spectra at the microscopic level is still developing, equivalent circuit models are commonly fitted to impedance spectra measured on roots.

There are two types of equivalent electric circuit models that have been commonly used to describe plant tissue: (a) distributed models that consist of distributed circuit elements (DCE) where a mathematical expression is used to describe the complex impedance (Repo, Zhang, Ryyppö, Vapaavuori, & Sutinen, 1994; Repo et al., 2005), and (b) lumped models that consist of a few ideal resistors and capacitors (Cole, 1940; Hayden et al., 1969; Inaba et al., 1995;
**Table 4** Summary of electrical impedance spectroscopy (EIS) case application

<table>
<thead>
<tr>
<th>References</th>
<th>Focus</th>
<th>Plant species</th>
<th>Setup</th>
<th>Growth media</th>
<th>Observations</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Repo et al., 2005</td>
<td>Measurement of tree root growth</td>
<td>Salix myrsinifolia</td>
<td>40 Hz–340 kHz, two electrodes, silver</td>
<td>Hydroponics</td>
<td>Sum of the resistors in the distributed model decreased with increasing root mass but was constant when the increase stopped</td>
<td>Could not differentiate the influences of the root mass, root surface area, and stem</td>
</tr>
<tr>
<td>Ozier-Lafontaine &amp; Bajazet, 2005</td>
<td>Analysis of root growth</td>
<td>Lycopersicum esculentum</td>
<td>10 Hz–1 MHz, two electrodes, Ag/AgCl</td>
<td>Potsoil</td>
<td>Change in stem electrode position produced large variation in the IS; IS varied with plant age (DAP) in terms of $Z'$, $Z''$, and phase; their amplitudes decreased with root development (age)</td>
<td>Could not link the variations in IS with changes at soil–root interface or root internal medium during growth</td>
</tr>
<tr>
<td>Cao et al., 2011</td>
<td>Estimation of root system size</td>
<td>Salix schwerinii</td>
<td>60 Hz–60 kHz 2 electrodes, silver needle</td>
<td>Hydroponics</td>
<td>The root–solution interfacial EC correlated linearly with the contact area of roots in solution for both “root and stem” and “root” ($R^2 = .61$)</td>
<td>Effect of stem was found to be stronger than the effect of roots; the dominating effect of stem needs to be verified by more studies using seed grown plants</td>
</tr>
<tr>
<td>Cseresnyés, Takács, et al., 2013</td>
<td>Mycorrhizal colonization of roots</td>
<td>Zea mays (maize)</td>
<td>100 Hz–10 kHz, two electrodes; tension</td>
<td>Pumice</td>
<td>Mycorrhizal plants showed slightly lower impedance than the nonmycorrhizal plants; EIS showed a negative log relation with root dry mass, length, and surface area ($R^2 = .8223–.9740$); mycorrhizas increased root absorption surface area</td>
<td>Certain field conditions favor the use of EIS, whereas others favor the use of ECM; the impact of the growth medium on the result is not fully understood</td>
</tr>
<tr>
<td>Cseresnyés, Rajkai, &amp; Vozárny, 2013</td>
<td>Impact of phase angle on EIS of roots</td>
<td>Zea mays (maize)</td>
<td>100 H–10 kHz, two electrodes, tension</td>
<td>Arenosol</td>
<td>Negative log relation was found between impedance and root extent parameters, dry mass ($R^2 = .561$ and .946) at 100 and 1,090 Hz, respectively; surface area ($R^2 = .491$ and .916) at 100 and 1,300 Hz respectively</td>
<td></td>
</tr>
</tbody>
</table>

(Continues)
**TABLE 4 (Continued)**

<table>
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*Note.* IS, impedance spectra; DAP, days after planting; $Z'$, real part of impedance; $Z''$, imaginary part of impedance; EC, electrical capacitance; EI, electrical impedance; EIS, electrical impedance spectroscopy.

Zhang et al., 1991). Distributed models are used in cases where lumped models are not appropriate (e.g., highly differentiated tissues), since it may be necessary to match the measured spectra (Burr et al., 2001; Repo et al., 2005). However, it is challenging to formulate a circuit model that properly represents the root system and at the same time allows a physical interpretation of the model parameters (Repo et al., 2012). Typically, more than one equivalent circuit fits well to the impedance data, but most likely only one of them will provide a reasonable representation of the physical properties of the system (Ozier-Lafontaine & Bajazet, 2005; Repo et al., 2012; Srinivas, Sarah, & Suryanarayana, 2003). Current research therefore favors different interpretation methods such as principal component analysis (PCA) (Repo et al., 2014), Debye decomposition of the spectra (Weigand & Kemna, 2016), and fitting of impedance spectra using Cole–Cole models (Martin, Nordsiek, & Weller, 2015; Mary et al., 2017). Future studies should incorporate interpretations at the microscopic level based on the physicochemical properties of roots to improve the understanding of the polarization mechanisms at the root segment scale.

Ozier-Lafontaine and Bajazet (2005) measured the impedance spectra of a tomato (*Solanum lycopersicum* L.) root system at different growth stages. They found that the magnitude of the real and imaginary parts of the impedance decreased as the root age increased (Figure 9). They showed that root growth results in variations in root electrical properties, but the reasons for such variations (e.g., variations of the soil–root interface or the root internal properties) remained unclear.

Repo et al. (2014) classified the impedance spectra (5 Hz–100 kHz) of the root system of Scots pine (*Pinus sylvestris* L.) with PCA to investigate whether root colonization with mycorrhizal fungi (*Hebeloma* sp. and *Suillus luteus*) could be detected in situ. On average, the correlation between the impedance spectra and mycorrhizal fungi showed 29 and 38.5% change for *Hebeloma* sp. and *Suillus luteus*, respectively. This makes sense as mycorrhizal colonization and mutualism between tree and mycorrhiza is known to improve nutrient uptake of trees (Simard, Jones, & Durall, 2003).

Spectral induced polarization is a variant of EIS that is mainly used in the geophysical community. It uses a broad range of frequencies from millihertz to tens of kilohertz and four electrodes to reduce the influence of contact resistances and to improve the accuracy of the complex impedance measurements (Zimmerman et al., 2008). Spectral induced polarization has been successfully applied in various biogeoophysical investigations (Abdel Aal, Atekwana, Slater, & Atekwana, 2006; Atekwana & Slater, 2009; Atekwana, Werkema, & Atekwana, 2006; Kessouri et al., 2019). Mary et al. (2017) studied isolated coarse roots (in the laboratory) and found that coarse roots showed higher resistivity and polarized differently compared with soil, suggesting that SIP is a promising tool for root studies. Ehosioke et al. (2018) designed a new setup that allows the application of SIP to assess the electrical response of a single root segment without the influence of the stem and growing media. They observed a very large polarization response (0.4–0.8 rad) from crop root segments with peaks occurring around ~10 kHz. This is linked to interfacial polarization as discussed above, but more studies are needed to better understand this. Ehosioke et al. (2018) also observed a low phase error of 0.06 rad due to the contact impedance, which is very small compared with the polarization response from roots. This new setup was used to compare the electrical properties of primary root, seminal root, brace root, and stem of a maize plant grown in sand tubes at 27 d after sowing. The
results show that the stem is more resistive than the root and displays a stronger polarization than the root (Figure 10). This implies that the plant stem and roots do not contribute equally to the electrical circuit of the stem–root–soil continuum. The stronger influence of the stem is probably due to the anatomical differences that exist between the root and the stem (Longui et al., 2012).

Recently, Tsukanov and Schwartz (2020) used SIP to investigate the relationship between the physical properties of wheat (*Triticum aestivum* L.) root grown in hydroponics and its electrical signature. They found a linear correlation between the electrical polarization in the 0.1 Hz to 1 kHz frequency range and root mass ($r^2 = .97$) and surface area ($r^2 = .82$). They also used SIP to monitor the electrical
response of wheat roots after poisoning the nutrient solution with cyanide. This led to a 30% decrease in polarization within 2 h. Their results further suggest that SIP is suitable to study morphology, physiology, and biomass of crop roots.

A key open challenge for EIS and SIP is the interpretation in the case of multicomponent systems. In geological systems, where the SIP method has been used widely and for decades, the physicochemical mechanisms of polarization for a single grain or pore throat are well understood, but the extrapolation of these mechanisms to a real porous media with heterogeneous characteristics (e.g., grain size, pore size, permeability, tortuosity, chemical heterogeneity, fluid phase distribution, etc.) is still a challenging task that requires theoretical developments (Kemna et al., 2012). Consequently, interpretation of complex impedance measurements in geosystems may be limited by the need to independently assess or assume the effects of chemical heterogeneity (Vaudelet, Revil, Schmutz, Franceschi, & Béгassat, 2011; Weller, Breede, Slater, & Nordsieck, 2011), temperature (Binley, Kruschwitz, Lesmes, & Kettridge, 2010; Martinez, Batzle, & Revil, 2012), saturation (Breede et al., 2012; Jougnot, Ghorbani, Revil, Leroy, & Cosenza, 2010), multiple fluid phases (Revil, Schmutz, & Batzle, 2011; Schmutz et al., 2010), grain size distribution (Revil & Florsch, 2010), pH (Skold, Revil, & Vaudelet, 2011), soil organic matter (Schwartz & Furman, 2015), or biological processes (Abdel Aal, Atekwana, & Atekwana, 2010; Atekwana & Slater, 2009; Ntarlagiannis, Williams, Slater, & Hubbard, 2005). In the particular case of root systems, interpretation of the complex impedance is difficult due to the complex architecture of roots and the inner root cell distribution and properties. Some setups involve the root–soil continuum, which adds the complexity of the geosystems to that of the root system with more variables to account for in the interpretation. Understanding of the root–soil–fluid interface from an electrochemical point of view and the inner root ionic distribution is necessary to properly interpret the overall signature when an external field is applied. More EIS and SIP studies at the root scale will help to improve the understanding of the main underlying mechanisms of polarization in roots, which is a

**Figure 9** Electrical impedance spectra of tomato root systems: (a) real part ($Z'$) of the impedance, (b) imaginary part ($Z''$) of the impedance, at different days after planting (DAP). The magnitude of the real and imaginary parts of the impedance decreased as the root age increased, which was linked to the increase in root system size with age (Figure 5 from reprinted Ozier-Lafontaine & Bajazet, 2005, reprinted with permission from Springer Nature)
FIGURE 10  (a) Electrical resistivity and (b) phase spectra of primary root, seminal root, brace root, and stem of a maize plant at 27 d after sowing. The stem was found to be more resistive and polarizable compared with the root sections. This indicates that the contribution of the stem will dominate that of the roots when impedance measurements are made on a stem–root continuum.

requirement for the development of a phytophysical model (Kessouri et al., 2019) that links root electrical signatures to root properties and root activity.

2.5.4  |  EIT

Electrical impedance tomography is similar to ERT in terms of setup and measurement, but it uses a frequency-dependent current injection to determine the real and imaginary part (or amplitude and phase) of the complex impedance. Electrical impedance tomography is the imaging counterpart of SIP and has been used to obtain information about root system anatomy, physiology, and morphology, as well as for biomass estimation (Table 2). The advantage of this type of measurement over other geophysical methods is that it has the diagnostic advantages of spectroscopy and the spatial resolution benefits of a tomographic approach as described by Kemna et al. (2012).

Weigand and Kemna (2017) investigated the suitability of multifrequency EIT to characterize and monitor plant root systems in a laboratory rhizotron experiment. They monitored the spatiotemporal distribution of the complex electrical resistivity of the root system of an oilseed plant (Brassica napus L. subsp. napus) in a water-filled rhizotron under nutrient deprivation using EIT. Total polarization parameters obtained from the multifrequency EIT results showed a low-frequency polarization response of the root system (Figure 11), which enabled a successful delineation of the spatial extent of the root system. The magnitude of the overall polarization response decreased along with the physiological decay of the root system due to nutrient deprivation. The length scales (i.e., frequencies) at which polarization processes occurred also changed due to the prolonged nutrient deficiency.

Mary et al. (2016) investigated the suitability of EIT for mapping tree roots in dikes. They concluded that EIT is useful to interpret anomalies produced by woody roots. In a follow-up study, Mary et al. (2017) used EIT to map roots embedded in an actual soil and observed that the maximum phase contrast between root and soil occurred at a frequency of 1 Hz. Anisotropy in the polarization was also observed such that the polarization signal was strongest when the current lines were aligned with the root. Electrical impedance tomography seems promising for root system investigations. However, some challenges need to be addressed in its application in order to explore its full potential. Most laboratory studies are conducted in hydroponics or in homogeneous substrate, allowing good contrast to be obtained. In agricultural fields, however, the topsoil is heterogeneous due to grain size variations (resulting from weathering), presence of microorganisms (e.g., earthworms), and human activities such as plowing. Obtaining a good contrast for root system characterization in a heterogeneous agricultural field may be
difficult (Weigand, 2017). Other challenges are the acquisition of high quality data in the field due to noise and high electrode impedances (van Treeck, Zimmermann, Weigand, Huisman, & Kemna, 2017), forward modeling errors resulting from imprecise electrode locations, inaccurate discretization, and so on (Zhao et al., 2019). Upscaling laboratory observations to the field scale will require suitable measurement devices, experimental setups, and procedures adapted to field conditions.

2.5.5 Joint discussion of stem injection methods

Measurement setups involving stem injection (Figure 2) of current as a means of characterizing root systems have commonly been used in a range of studies reviewed above. Although they adopted different methods such as ERM (Cao et al., 2010), ECM (Dietrich et al., 2012; Kendall, Pederson, & Hill, 1982), MALM (Mary et al., 2019; Peruzzo et al., 2020), EIM (Čermák et al., 2013; Urban et al., 2011), and EIS (Repo et al., 2005; Repo et al., 2012), it is interesting to note that all studies share a common conclusion. Proximal leakage of current at the stem–soil and solution interface appears as a common challenge in all measurement setups involving the stem–root–soil continuum. For example, Cao et al. (2010) observed that submerged roots did not contribute significantly to measured resistance, which is probably due to current leakage at the stem–solution interface. Urban et al. (2011) further supports this argument, as they concluded that distal roots conduct only a fraction of the current, and stem injection methods therefore cannot reliably estimate root system extent. Dietrich

**FIGURE 11** The spectral and spatial variation of polarization response of the root system of an oilseed plant: (a) the position of the bulk root system, and (b) the change in spectral shapes over time, showing decreasing polarization. The polarization occurred at a length scale indicated with dotted lines. (c, d) Spatial variation of the polarization response, showing (c) Time Step 1 and (d) Time Step 21. The change in spectral shape with time suggests that relaxation time distributions can provide additional information on the electrical response of root systems. The symbol $\sigma''$ is the imaginary conductivity; $z$ and $x$ are the vertical and horizontal dimensions of the rhizotron (modified from Weigand & Kemna, 2017, available under Creative Common Attribution 3.0)
et al. (2012) reported that excision of submerged roots did not affect capacitance and concluded that root capacitance is dominated by the tissue between the stem electrode and the solution interface, and closely related to the cross-sectional area of the root at the solution interface, thus agreeing with the findings of previous studies (Cao et al., 2010; Kendall et al., 1982; Repo et al., 2005; Repo et al., 2012; Urban et al., 2011). Mary et al. (2019) also observed proximal current leakage when the soil was wet and provided a more conductive pathway than the roots. Clearly, in situ applications of these methods should carefully consider other confounding variables, such as soil moisture (Csresnyés et al., 2018; Mary et al., 2019), soil texture and porosity (Chloupek, 1972), tissue density (Ellis, Murray, & Kavalieris, 2013; Ellis, Murray, et al., 2013), electrode type (Kormanek et al., 2016), and growth media (Csresnyés et al., 2017). In any case, careful calibrations will be required.

### Summary and Conclusions

#### 3.1 Resistance, capacitance, or impedance?

Plant scientists, earth scientists, and geophysicists all see potential in electrical methods to characterize the root system of plants given the observed correlations between electrical measurements or derived electrical properties with root biomass and/or root activity. Initial studies focused on resistance and capacitance measurements at a specific frequency, whereas recent studies have increasingly focused on impedance measurements at several frequencies; this approach allows both conduction and polarization effects to be studied simultaneously, giving information on roots at different time and spatial scales. Electrical conduction in bulk soil with roots is still mostly associated with soil moisture dynamics and is best used to study root physiological processes (e.g., soil water content changes associated with root water uptake). Electrical polarization of roots is related to the anatomy of the root tissue and cell structure (e.g., cell membrane properties), fluid composition, and plant metabolic state.

#### 3.2 Technical details of the measurement

Next to the targeted property, the technical details of the measurements (amount of electrodes, current injection strategy, electrode material, scale) determine the data quality and information content. In general, it was found that

- Four electrode measurements reduce problems with contact resistance and are therefore more reliable than measurements with two electrodes only.
- Injection schemes with an electrode in the stem (such as EIM, MALM, and EIS) are integrative measurements of the stem–root–soil continuum, and the results depend strongly on where the electrode is placed in the stem and the properties of the growing medium close to the stem, since the current may leave the system at the proximal parts of the root–soil interface.

#### 3.3 Interpretation models

Equivalent electrical circuit models are essential to interpret contributions of different components in the soil–plant continuum governing electrical conduction and polarization during plant- or field-scale measurements. Small-scale measurements such as ERM and SIP on root segments (with or without the plant still connected) are important to validate equivalent electrical circuit models and to determine the contribution of individual components on the behavior of the full root architecture in the soil. Equivalent circuit models, however, are not very accurate because of the problem of nonuniqueness as more than one circuit model can provide a good fit for the impedance data. Therefore, there is an increasing use of alternative interpretation methods such as classification-based approaches (e.g., PCA), fitting of empirical spectral models (e.g., Cole–Cole model), or Debye decomposition of the spectra. Recent studies have shown that low-frequency polarization of root systems probably originates from polarization at various double layers, be it at cell membranes, or larger ion-selective structures such as the Casparian strip or the root surface. This is important not only to study roots, but also to study soil moisture and solute transport dynamics in relation to plant growth, since the contribution of the presence of roots to the measured electrical signal is now typically neglected. Interpretations at the microscopic level based on the physicochemical properties of the root should be incorporated to improve the understanding of the mechanisms driving polarization at root segment scale.

#### 3.4 Future research perspectives

More SIP studies should be used to understand the variability of root electrical properties in time and space, across species and genotypes. To achieve this, controlled experiments will be required to obtain fundamental information, and these experiments must be well designed and constrained to study the effects of each key parameter separately. Although different methods have investigated the influence of anatomy and physiological processes on the root electrical response, this is not yet fully understood. More studies are needed to link root anatomy and physiology with electrical properties.
(conduction and polarization) of roots at different scales. Data obtained at root segment scale are necessary to validate phenomena underlying conduction and polarization occurring at the root segment and root architecture scale, and thus interpretation at the microscopic level will help to improve the understanding of the underlying mechanisms and establish models summarizing these phenomena.

Due to the complexity of the root system architecture embedded in the soil matrix, upscaling electrical properties measured at the root segment scale to plant scale or to field scale is a challenging task (Mary et al., 2016; Weigand & Kemna, 2017). Process-based multiscale numerical modeling of a soil–root continuum can thus pave a way for our understanding of upscaling mechanisms as we move from segment to plant and to field scale. A recent detailed numerical study at the plant scale shows that roots indeed affect ERT measurements and the effective electrical conductivity (Rao et al., 2019). However, explicitly representing root architecture (as in their study) in the computational domain demands a very high spatial resolution, thus increasing computational time and memory consumption. Further research on numerical approaches for such mechanistic models and the effect of the multiple scale levels involved in the underlying processes on the macroscopic properties is required. The parameterization of root segment electrical circuit models and how to link it to Cole–Cole or Debye decomposition models that will help interpret field measurements is also important.

4 | GLOSSARY

Alternating electric field: A voltage source in a medium usually produce an electric field that causes electric current to flow, if the medium is a conductor. If the electric charge (current) only flows in one direction, it is called direct current (DC). If the current changes direction periodically, it is known as alternating current (AC).

Capacitance (C): The electrical property of a capacitor that measures its ability to store an electrical charge on its two plates. The SI unit is farads (F).

Capacitors: These are devices that can store an electrical charge (current) when connected to a voltage source.

Cole–Cole model (CCM): A phenomenological model used to analyze SIP data. It is mostly used to describe the strength and frequency dependency of individual polarization peaks on SIP data. It is expressed as \( \rho^* = \rho_\infty + (\rho_0 - \rho_\infty) / [1 + (\omega \tau)^s] \) where \( \rho_0 \) and \( \rho_\infty \) are low-frequency and high-frequency electrical resistivity values, respectively. \( C = 1 - \infty \), representing the CCM exponent that describes the relaxation time distribution. This can also be expressed in terms of complex conductivity (\( \sigma^* \)), but complex electrical resistivity is used often in practical terms because it is calculated directly from measured electrical impedance.

Current injection: The process of transmitting current through an object of interest to assess its electrical properties.

Debye decomposition: The process of determining a relaxation time distribution in place of a fixed number of relaxation time.

Polarization: In an electric field, electric charges do not flow through dielectric materials as they do in electrical conductors; instead, they shift from their equilibrium position, and the process is termed dielectric polarization. A dielectric material is an electrical insulator that can be polarized when an electric field is applied to it. Dielectric properties refer to the storage and dissipation of electric and magnetic energy in materials.

Dielectric relaxation: Momentary delay (lag) in the dielectric constant of a material. The dielectric constant is the measure of a material’s ability to store electrical energy in an electric field.

Electrical impedance (Z), resistivity (\( \rho \)), and Conductivity (\( \sigma \)) (real, imaginary, magnitude, and phase): Electrical impedance is the measure of the opposition that a circuit offers to a current when a voltage is applied. Impedance represents the concept of resistance in AC circuits and has both magnitude and phase. In DC circuits, resistance could be expressed as impedance at zero phase angle. Impedance is a complex number, the SI unit is ohms (\( \Omega \)), just like resistance. Impedance (Z) is defined in Cartesian form as \( Z' = Z'' \) where \( w \) is the angular frequency, \( j \) is the imaginary number, \( Z \) is the real part (representing conduction), and the \( Z'' \) is the imaginary part (representing polarization).

Electrical resistivity (\( \rho \)) is an inherent property of a material that quantifies its opposition to the flow of electrical current. Its SI unit is ohm meters (\( \Omega \cdot m \)). Electrical conductivity (\( \sigma \)) is a measure of materials ability to conduct electric current. The SI unit is siemens per meter (S m\(^{-1} \)). It is the reciprocal of electrical resistivity. Complex resistivity (\( \rho^* \)) and conductivity (\( \sigma^* \)), which are effective material properties, can be calculated from the impedance using a geometric factor \( K \), which accounts for the geometric dimension of the measurement: \( \rho^*_0 = K Z^*_0 = \rho'_0 + j \rho''_0 = |\rho|e^{j\phi} = |\rho|e^{j\phi} = |\rho|e^{j\phi} \), where \( |\rho| \) is the resistivity magnitude, and \( \phi \) is the phase shift. \( \sigma^*_0 = 1/\rho'0 = 1/K Z'0 \).

Electrode: An electrical conductor used to establish a contact with a nonmetallic part of a circuit such as a semiconductor, an electrolyte, etc.

Electrolyte: Substances that dissolve in polar solvents such as water, to produce an electrically conducting solution.

Entropy: A measure of the degree of randomness or disorderliness in a system. A “negative entropy” (negentropy) refers to increasing orderliness in a system.

Equivalent circuits: Theoretical circuits that retain all the electrical properties of a given circuit.
Grounding impedance: The resistance offered by the ground electrode to the flow of AC current into the ground. This is the ratio of the potential of the ground electrode to the current injected by it (i.e., \( I_g = \text{potential to ground/current} = V/I \)).

Impedance spectrum and spectroscopy: Impedance spectrum is a plot of impedance \( \frac{Z}{\omega} \) against frequency, which reveals the change of impedance of a sample with frequency during excitation by alternating current. Spectroscopy is the process of using impedance spectrum to analyze the dielectric property of a material as a function of frequency.

Inversion and tomography: Tomography is the process of recording a projection series from different angles and then using computational reconstruction to produce the image of the object in three dimensions (tomograms).

Ion pumps and ion channels: These are assemblies of integral membrane proteins that regulate the transport of ions into and out of cells or organelles, resulting in the generation of electrical signals. “Ion pumps” actively transport ions against a concentration gradient, whereas “ion channels” allow a passive flow of ions down a concentration gradient.

Nonpolarizable electrodes: An electrode whose potential is not affected by the current passing through it. They are used to eliminate the electrode polarization effect that occurs when metals (e.g., stainless steel electrodes) are used.

Organelles: Tiny cellular structures that perform specific functions within a cell, examples include vacuole, nucleus, etc.

R-C circuit: An electric circuit composed of resistors and capacitors driven by a voltage or current source.

Relaxation time: The time required for a perturbed system to return back to equilibrium.

Resistor: A passive electrical component used to create resistance to the flow of current.

Senescence: The process of aging characterized by gradual deterioration of functional characteristics.

Suberization: This is the impregnation of cell walls with suberins resulting in the formation of cork tissues. These suberins are complex macromolecules (biopolymers) that form protective barriers to prevent water and nutrients from entering into the stele through the apoplast. In roots, suberin is formed in the cell walls of the endodermal cells, collectively known as the “Casparian strip.”

Vacuole: These are membrane-bound sacs within the cytoplasm of a cell. They are very important in providing support and also help to perform other functions such as storage, waste disposal, and protection.

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