O₂ dynamics in the rhizosphere of young rice plants (Oryza sativa L.) as studied by planar optodes

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Abstract
Background and aims Radial O₂ loss (ROL) strongly affect the O₂ availability in the rhizosphere of rice. The ROL create an oxic zone around the roots, protecting the plant from toxic reduced chemical species and regulates the redox chemistry in the soil. This study investigates the spatio-temporal variability in O₂ dynamics in the rice rhizosphere.

Method Applying high-resolution planar optode imaging, we investigated the O₂ dynamics of plants grown in water saturated soil, as a function of ambient O₂ level, irradiance and plant development, for submerged and emerged plants.

Results O₂ leakage was heterogeneously distributed with zones of intense leakage around roots tips and young developing roots. While the majority of roots exhibited high ROL others remained surrounded by anoxic soil. ROL was affected by ambient O₂ levels around the plant, as well as irradiance, indicating a direct influence of photosynthetic activity on ROL. At onset of darkness, oxia in the rhizosphere was drastically reduced, but subsequently oxia gradually increased, presumably as root and/or soil respiration declined.

Conclusion The study demonstrates a high spatio-temporal heterogeneity in rhizosphere O₂ dynamics and difference in ROL between different parts of the rhizosphere. The work documents that spatio-temporal measurements are important to fully understand and account for the highly variable O₂ dynamics and associated biogeochemical processes and pathways in the rice rhizosphere.

Keywords Planar optode · Rice (Oryza sativa L.) · Radial oxygen loss · Rhizosphere · Plaques imaging · Oxygen

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Introduction

Due to the importance of rice (*Oryza sativa*) as a staple food source in global human nutrition, it is likely the most studied plant growing in flooded and waterlogged soils. However, studies on O$_2$ dynamics in the rice rhizosphere remain a challenge and the biogeochemical functioning of the rice rhizosphere is still rudimentary. Flooded conditions are a widespread phenomenon among rice and it is estimated that about 75% of all rice are grown under O$_2$ depleted conditions in flooded and waterlogged soils, during part or all of the cropping period (Roger et al. 1993). Rice plants growing in O$_2$ depleted soils are potentially exposed to high levels of harmful reduced/toxic chemical species, like H$_2$S and Fe$^{2+}$ (Armstrong and Armstrong 2005; Mongon et al. 2014). To overcome these challenges a range of wetland plants, including some rice genotypes, have evolved two adaptations (I) development of a barrier against solutes and gasses e.g., by increased suberization in exodermis and/or lignification in sclerenchyma cells (Greenway et al. 2006; Kotula et al. 2009) and/or (II) leakage of O$_2$ to establish a protective oxygenated zone around the roots, a strategy facilitated by well-developed gas conducting aerenchyma inside the roots (Armstrong 1980; Kirk 1993). The two strategies vary extensively between different species of wetland plants and even among genotypes of the same species of wetland land plants and even among genotypes of the same plant.

The widespread occurrence of ROL in rice plants and the ability to form a barrier to ROL when grown in stagnant solutions is extensively reported e.g., (Colmer et al. 1998; Insalud et al. 2006; Kotula et al. 2009). However, the ability to form a barrier to ROL varies among different genotypes as demonstrated by Colmer et al. (1998) and Colmer (2003), who investigated different rice genotypes of lowland rice from drained and waterlogged soils. The studies demonstrated that genotypes to various extend showed an increase in the root porosity and a development of a tight barrier to ROL, when grown in stagnant O$_2$ depleted conditions, as compared to oxygenated conditions (Insalud et al. 2006; Kotula et al. 2009; Shiono et al. 2011). The plasticity in root physiology presumably contributes to the ability of rice to grow in diverse environments at different levels of waterlogging (Colmer 2003).

The O$_2$ dynamic associated to roots without a tight barrier to ROL can have important implications for the redox conditions and the biogeochemical processes and pathways that take place in the rhizosphere, such as trace metal availability (Williams et al. 2014) and plaque formations (Mendelssohn et al. 1995). The formation of metal plaque in the rhizosphere of wetland plants is often associated to ROL as plaque mainly reflects the distribution of Fe-oxides (Wu et al. 2012). Plaque formation on roots have been demonstrated to be important for root nutrient and trace metal dynamics, generally limiting phosphorus uptake and acting as sorption sites for trace metal such Arsenic (Jiang et al. 2009; Deng et al. 2010; Seyfferth et al. 2010). Plaque formation on rice roots have generally been studies using extraction based methods (Deng et al. 2010; Cheng et al. 2014) and only few studies have investigated the spatial variation of plaque distributions (Seyfferth et al. 2010), however, the link between spatial variations in ROL and plaque is not well documented.

Traditionally ROL from rice roots have been studied using microelectrodes (Frenzel et al. 1992; Revsbech et al. 1999; Colmer and Pedersen 2008), bulk measurements (Klundze et al. 1993; Kirk and Du 1997) and cylindrical O$_2$ cathodes (Armstrong 1994; Insalud et al. 2006). The studies demonstrate that ROL from rice roots can be affected by e.g., irradiance (Waters et al. 1989; Revsbech et al. 1999; Colmer and Pedersen 2008) and flooding (Waters et al. 1989; Pedersen et al. 2009). However, these studies often rely on single point measurements of individual roots or plants grown in artificial medium such as hydroponics or agar medium. Single point measurements have provided valuable insight in root O$_2$ dynamics of rice plants, especially in cases where they have been performed on soil grown plants (Frenzel et al. 1992; Revsbech et al. 1999). However they only provide a limited understanding on the overall O$_2$ dynamic in the rhizosphere and still little is known about the spatio-temporal variations in ROL of intact rhizosphere grown under natural condition.

Recently, two-dimensional luminescent imaging (planar optodes) have been applied for rhizosphere and soil studies investigating 2D O$_2$, pH and pCO$_2$ dynamics (Frederiksen and Glud 2006; Blossfeld et al. 2013; Rudolph et al. 2013; Williams et al. 2014). Planar optodes are optical semitransparent sensors that allow real-time and two-dimensional imaging of e.g., O$_2$ by means of a luminescent and analyte specific indicator. By imaging the light emitted from the indicator, the analyte concentration can be quantified and related to physical structures in the soil. Planar optodes offer high
spatial (>20 μm) and temporal (1–30 s) resolution comparable to those of microelectrodes, but can resolve dynamics of larger areas of the soil rhizosphere (>20 cm²) as compared to a single-point measurement (Frederiksen and Glud 2006; Rudolph et al. 2013; Jovanovic et al. 2015). The use of planar optodes has provided new conceptual and quantitative understanding of solute dynamics in such environments and on the importance of microniche structure for the biogeochemical functioning in such environments (Jensen et al. 2005; Schreiber et al. 2012; Faget et al. 2013). Here we apply two-dimensional, high-resolution planar optode imaging to investigate the spatio-temporal variability in O₂ dynamics between individual roots and different parts of the root system of rice exposed to different environmental conditions. We furthermore demonstrate a simple system for plaque imaging. Plants were grown in flooded soil under natural conditions. The O₂ dynamics of the intact rhizosphere is investigated as a function of ambient O₂ levels, irradiance and plant development for emerged and submerged plants. The presented data are used to provide an improved conceptual and quantitative understanding of spatio-temporal O₂ dynamics in the rhizosphere of young rice plants.

Materials and methods

Plant, soil and rhizotrons

Acrylic rhizotrons (H×W×D: 40×10×1.5 cm) (Fig. 1a, b) were packed with soil sieved to<2 mm. The soil was loamy sand (sand: 510 g kg⁻¹ sand, 420 g kg⁻¹ silt, 70 g kg⁻¹ clay; pH(CaCl₂) 5.4; 23 g kg⁻¹ organic matter). The soil was added in layers and was carefully compacted to ensure an overall homogeneous soil structure. The porosity of the soil was calculated based on the mass of soil filled in the rhizotrons, the filled rhizotron volume and estimated soil particle density of 2.65 g cm⁻³. The depth of the soil was ~25 cm. A detachable front window of the rhizotron was opened and the soil surface was covered by a sheet of transparent Nuclepore membrane (Whatman.com, 0.2 μm pore size), ensuring that the soil and roots remained undisturbed during removal of the front window. The membrane did not have any influence on the planar optode imaging - this was confirmed by comparing calibration curves for the planar optode with and without the membrane (data not shown). Subsequently, the rhizotrons were gently filled with tap water and kept anaerobic for two weeks prior to transplanting the rice seedlings.

Rice seeds, Oryza sativa (Indica cv. RIL 46) were germinated for ten days between moist tissue paper before transplanting the seedlings into the anaerobic soil. At the time of transplantation the roots and shoots were ~10 and 8 mm long, respectively. Seedlings were planted close to the front window of the rhizotrons (Fig. 1b), which were kept at an inclination of 45° to ensure that roots developed along the front window of the rhizotron.

Plants were grown at 22±2°C and an irradiance of ~150–200 μmol photons m⁻² s⁻¹ with a 14:10 h light:dark cycle for three weeks prior to the three weeks
long experimental period. The soil was kept in fully water saturated and anaerobic conditions. However, during the experimental period, the water height inside the rhizotron was varied to simulate both emerged and submerged conditions for the rice plants. The water level was maintained by replacing the evaporated water. No nutrients were added to the system.

In the following sections we use the term “emerged” for plants growing with leaves exposed to the air, but with a water height of 8–10 cm inside the rhizotron, covering the majority of the stem (Fig. 1 a, b). For “submerged” plants the entire plants were covered by water. During plant growth and the experimental period, the lower part of the rhizotron was kept in darkness to avoid exposing the roots and soil to light, which could have affected the root development and induce the growth of microphytes at the front plate.

Before planar optode imaging, the front window of the rhizotron was removed and the rhizotron was transferred into a larger water reservoir, with the root and soil surface facing the front of the reservoir (Fig. 1b). To ensure that the soil experienced minimal O2 exposure as the front window was removed, the procedure was carried out in a deoxygenated water basin. During the above procedure the rhizotron was kept at an inclination below 45°, to ensure minimal displacement of the soil and root. The rhizotron was fastened against the wall of the reservoir chamber by a number of braces. The transparent front window of the reservoir chamber had previously been equipped with an O2 sensitive planar optode foil (see later). The replacement of the front window did not lead to any disturbance or oxygenation of the soil as evaluated from the planar O2 sensor. Furthermore, the soil and plants were left to stabilize for a minimum of 24 h to ensure that all O2 gradients had reestablished before the experimental work was initiated.

The water level in the reservoir was maintained below the top of the rhizotron (Fig. 1a). The water reservoir with rhizotron was then placed in front of the camera and LED setup. During O2 imaging, the water reservoir including the rhizotron was kept at a 45° inclination with the O2 imaging camera positioned perpendicularly to the optode. The long term stability of planar optodes sensors, similar to the ones used in this study, has been reported to be in the order of months (Precht et al. 2004) and we likewise observed no drift in the signal. No toxic effects of planar optodes have been reported.

Optode sensor fabrication

The O2 quenchable luminophore PtTFPP (Platinum(II)-5,10,15,20-tetrakis-(2,3,4,5,6-pentafluorophenyl)-porphyrin (frontiersci.com), was chosen as O2 indicator due to its excellent photostability and relatively long luminescent lifetime (Borisov and Klimant 2007). Coumarin C545 (SigmaAldrich.com) was used as antenna dye in order to increase the brightness of the indicator (Mayr et al. 2009; Larsen et al. 2011). PtTFPP and Coumarin C545 were mixed in a 1/2 % (wt/wt) ratio, respectively, and dissolved in 4 % (wt/wt) polystyrene to form a sensor cocktail, using chloroform as solvent. The sensor cocktail was coated onto a 125 μm thick transparent polyester foil (Goodfellow.com) using a homemade knife-coating device. The final thickness of the highly transparent O2 sensing layer was ~5 μm. The size of the optode was 8×15 cm and covered the entire rhizosphere of the rice plant (Fig. 1a).

Rhizosphere imaging of O2 and plaque

The planar optode system used in this study is similar to what has been presented previously (Holst et al. 1998; Frederiksen and Glud 2006) and will only be described briefly. The luminescent lifetime of the O2 quenchable luminophore was recorded with a fast gateable 12-bit Charged Coupled Device camera (SensiCam, PCO.de) through a 590 nm longpass filter (UQGoptics.com). Excitation light was delivered from four blue high power LEDs (LXHL-LR3C, Luxeon.com) equipped with a 470 nm short pass filter (UQGoptics.com). The operation of camera and LED’s were synchronized via a custom made PC-controlled triggerbox by the software Look@Molli (Holst and Grunwald 2001). The luminescent lifetime was inferred from two well defined time frames configured by the software Look@Molli (Holst and Grunwald 2001). The luminescent lifetime was inferred from two well defined time frames configured by the software. Recorded images were calibrated using the luminescent lifetime for two known O2 levels (saturation in the overlying water and 0 % air sat. in anoxic soil), using a modified Stern-Volmer equation (Eq. 1) (Klimant et al. 1995; Glud et al. 1996)

$$\frac{\tau}{\tau_0} = \left[ \alpha + \left(1-\alpha\right) \left(\frac{1}{1 + K_{SV} \cdot C}\right) \right]$$

(1)

Where $\tau_0$ is the luminescent lifetime in the absence of O2 and $\tau$ is the lifetime in the presence of O2 at levels (C), $K_{SV}$ the Stern-Volmer quenching constant and $\alpha$ is
the non-quenchable fraction of the luminescent signal, the latter experientially determined to 0.12. All images were recorded in darkness, i.e., for “light-images” ambient light was briefly turned off, to avoid interference from ambient light. For each O₂ image a corresponding black and white (BW) image was recorded. The maximum theoretical spatial resolution that could be achieved with the given optical configuration was ~120 × 120 μm, however, we estimate that the effective spatial resolution is two-three times lower than the maximum theoretical resolution due to light guidance in sensor foil and acrylic walls (Fischer and Wenzhofer 2010). This, however, will not affect any of the qualitative or quantitative conclusions made in this study.

The diffusive O₂ uptake (DOU) just below the SWI was calculated from extracted vertical O₂ profiles, using Fick’s first law of diffusion (Jørgensen and Revsbech 1985): DOU = Φ × Ds × ∂C/∂z, where Φ represents the soil porosity, Ds the molecular diffusion coefficient of O₂ in the soil at the given temperature and ∂C/∂z represents the O₂ gradient. The average volume-specific O₂ consumption was calculated as DOU/O₂ penetration depth (Glud 2008). In the following sections we define the rhizosphere oxic area as areas with O₂ levels above the detection limit of the optode sensor ≥ 1 % air sat. (2.3 μmol L⁻¹). The detection limit is quantified as three times the standard deviation of the measured concentrations at anoxia for an area of 5 × 5 cm. We predominantly use the “oxic area” as a parameter for describing the rhizosphere oxygenation as it allow for a direct comparison between plants and treatments without converting the values to a more complex geometry. However, the oxic area is directly related to the ROL. As only one planar optode imaging system was available, not all plants were imaged continuously during the 3 week experimental period; hence, the longest time series recorded is 6 days. The two-dimensional distribution of plaque formation in the rhizosphere was imaged in a parallel experiment. Due to overlapping absorption spectra of blue light by the plaque and planar optode, it was not technically possible to perform simultaneously imaging and hence plaque imaging was performed on a separate and older root system. The imaging principle is based on blue light absorption of Fe- and Mn-oxides (Sherman and Waite 1985; Estapa et al. 2012), the dominating minerals of root plaque in aquatic environments (Christensen and Sand-Jensen 1998). The absorption was imaged using a modified Canon D1000 camera (Larsen et al. 2011) and a blue LED light source, as previously described for the planar optode imaging.

Experimental procedure

The O₂ distribution in the rhizosphere was investigated as function of irradiance and O₂ levels in the overlying water and air, respectively, for both submerged and emerged plants. The O₂ saturation was regulated by mixing N₂ and atmospheric air by means of two 5850S mass flow controllers (BrooksInstruments.com) controlled by a type 0154 digital control/readout unit (BrooksInstrument.com). The O₂ saturation in the rhizotron above the soil level was monitored in parallel by a Microx TX3 micro-optode O₂ meter (PreSens.de). Levels of irradiance were manipulated by varying the height of the growth lamp (400 W HQI) relative to the plant, the absolute irradiance was measured with a Li-Cor 250 light meter equipped with a planar LI-190 Quantum Sensor (Licor.com). In total three plants were investigated.

Results

Soil and root O₂ dynamics

The soil in the rhizotron had a porosity of 0.58. The O₂ penetration depth at the soil water interface (SWI) was 2.7 mm (SD ± 0.2, n = 6) at 100 % air sat. (273 μmol L⁻¹), as derived from the planar optode images. The diffusive O₂ uptake (DOU) just below the SWI was 17.5 mmol m⁻² d⁻¹ (SD ± 2.9, n = 6). The DOU represents the soil O₂ uptake mediated by diffusion (Glud 2008). The average volume-specific O₂ consumption amounted to 0.07 nmol cm⁻³ s⁻¹ (SD ± 0.015, n = 6). This value was used to estimate the integrated rhizosphere related O₂ consumption rates (see later). The O₂ penetration depth and volume-specific O₂ consumption rates are in agreement with values measured in natural rice paddy soil, ranging between 0.8 to 2.0 mm and ~0.02–0.14 nmol cm⁻³ s⁻¹, respectively (Damgaard et al. 1998; Reim et al. 2012).

Root system architecture and the corresponding O₂ images were investigated for three rhizotrons each containing one 3–6 week old plant; a typical example is depicted in Fig. 1c, d. Typically, a zone of high O₂ levels was observed at the base of the plant (later referred to as ‘basal root zone’) that was usually connected to the O₂
Light availability affected the O₂ level in the rhizosphere for both emerged and submerged plants. For an emerged plant the onset of darkness lead to a rapid decrease in the oxic rhizosphere area of 19.2 % (Fig. 2a) from a stable value of 1080 mm² at ambient O₂ levels of 100 % air sat. and irradiance of ~200 umol photons m⁻² s⁻¹ throughout the experiential period.

Most roots were leaking O₂ however 5–10 % of the actively growing did not show any O₂ leakage. There was no apparent correlation between depth in the soil and the intensity of O₂ leakage.

Irradiance induced O₂ dynamics in the rhizosphere

For an emerged plant a step-wise increase in irradiance from 0 to ~300 µmol photons m⁻² s⁻¹ caused a gradual increase in rhizosphere oxic area of 64.0 % (from 364 to 597 mm²), the corresponding increase in the mean O₂ level within the oxic area was 57.7 % (Fig. 2b). As for the day-night transition roots and root tips showed smaller change in O₂ level, relative to the basal root zone when levels of irradiance was changed. During these experiments the O₂ level in the air above the emerged plant was maintained at 100 % air sat. Changing the irradiance level for a submerged plant gave similar results to those of emerged plants (Fig. 2b). It should be noted that the oxic area of the submerged plant was larger compared to the emerged plant due to difference in plant size, the trend and magnitude of the increase in oxic area is however comparable between the plants.

Even though irradiance had considerable effect on the O₂ levels and dynamics in the rhizosphere, the most dramatic changes were induced by regulating the O₂ level in the air or water surrounding the aboveground plant (Fig. 3a, b and d). For an emerged plant exposed to 100 % air sat., the root oxic area was 1012 mm². However as the O₂ level was stepwise lowered, the anoxic area gradually increased and the oxic area only accounted for 142 mm² at ambient O₂ levels of 21 % air sat. (Fig 3d), corresponding to a decrease in the oxic area of 870 mm². The maximum O₂ level in the oxic area decreased from 88.2 to 18.9 % air sat., whereas the mean O₂ level in the oxic area decreased from 34.8 to 12.7 % air sat. Extrapolation of the observed trends in oxic area and average O₂ levels indicated that the rhizosphere would become completely anoxic at ambient O₂ levels of ~13 % air sat. The root tips appeared to be most sensitive to declining ambient O₂ levels; here anoxia was reached at ambient levels of 39.4 % air sat. (±SD=10.7, n=5). In comparison anoxia was reached at ambient O₂ levels of 19.7 % and 13.7 % (±SD=10.6, n=4), for the basal root zone and the middle section of long roots, respectively.

Extracted O₂ profiles from a typical root (Fig 3e), demonstrated that the O₂ level at the root surface decreased from 26.2 to 5.14 % air sat., when the O₂ level in the ambient air was reduced from 100 to 21 % air sat.
Fig. 2 a Temporal development of the mean O₂ level (black circle) and the oxic area (white circle) in the rhizosphere of a rice plant during a light/dark transition for an emerged plant (see text for explanation). The oxic area was defined as the area with O₂ levels >1 % air sat. Light grey color represent the dark period. The O₂ levels represent the mean value within the corresponding oxic area and are based on measurements of 60,763 to 76,388 pixels. b Development of the oxic area in the rhizosphere as function of irradiance for an emerged (black bars) and submerged plant (white bars). The difference in oxic area for the two plants is due to variation in plant size. Error bars represent SD, calculated form three individual images at each irradiance level. Each image is based on measurements of 25,294 to 41,502 pixels.

Fig. 3 a Planar optode O₂ images of the rhizosphere of emerged rice (Oryza sativa L.) as function of O₂ level in the air above the plant. The O₂ level in the air is 21 and 100 % air sat., for (a) and (b) respectively. c Close-up of a single root at 100 % air sat. in the air above the plant. The dashed lines represents the position of extracted profiles, presented in (e) and (f). Images are scaled only to show O₂ levels from 0 to 70 % air sat., for better visualization of rhizosphere O₂ dynamics. d Oxic area of the rhizosphere as function of O₂ level in the air and water above the plant for a submerged and emerged plant. e Variability of O₂ distribution across a single root from one profile, as function of O₂ level in the air above the plant. The position of the extracted profiles is equivalent to the “0 mm” profile (furthest away from the tip on panel e). f Spatial variability in root O₂ distribution as function of distance from base of the root towards the tip, the position of the profiles are indicated on (e), with profile “20 mm” closest to the root tip. The points in panel e and f represent individual pixels, with a spatial resolution of 120×120 μm.
Consequently, the width of the oxic zone decreased from 1.3 to 0.6 mm. The extent of the oxygenated area varied between neighbouring roots and along individual roots. A transect of O$_2$ profiles perpendicular to a single root is presented in Fig. 3f, showing that the width of the oxic zone decreased from 2.6 to 0.3 mm when approaching the root tip at 100 % air sat., and irradiance of ~200 μmol photons m$^{-2}$ s$^{-1}$. The general pattern as shown in Fig. 3e and f were observed for all roots on the plant and for all plants investigated.

Changing the O$_2$ level in the ambient water for submerged plants gave very similar results as for emerge plants. In fact we found that the root oxic area disappears when an O$_2$ level in the water was reduced to ~13 % air sat., this was the same threshold as estimated for the emerged plant (Fig. 3d).

Temporal O$_2$ dynamics

The spatial extent of the oxygenated rhizosphere zone was dynamic, as the rhizosphere evolved; older roots ceased to leak O$_2$, while young, newly grown roots exhibited high O$_2$ release. We observed root growth rates of up to 2.6 mm d$^{-1}$. One example of the temporal development is depicted in Fig. 4, where an emerging root begins leaking O$_2$ into previously anoxic soil (root ‘a’ in Fig. 4), while concurrently the O$_2$ levels around a neighbouring root drastically diminish (root ‘b’). Interestingly it could be observed that as O$_2$ leakage from the main part of root ‘a’ diminishes with time, leakage from its laterals roots could be observed (Fig. 5). In contrast, O$_2$ release from the neighbouring roots remained virtually unchanged during this ~5.5 day period.

Plaque distribution

The distribution of metal plaques was highly patchy and variable between different roots – no roots were observed to have plaque formation along the entire root (Fig. 6). The total visual metal plaque-containing area in the rhizosphere was found to be 739 mm$^2$, 5.9 % of the visible area. Plaque density was highest in a zone 41.1–66.1 mm below the soil surface. Below this zone, plaque density decreased gradually. The soil area covered by plaque did on average corresponded to ~30.4 % of the visible root surface area.

Discussion

O$_2$ dynamics in the rhizosphere studied by planar optodes

We present novel high resolution two-dimensional images of the O$_2$ dynamics of the rhizosphere of lowland rice. We demonstrate that the spatio-temporal O$_2$ distribution in the rhizosphere is highly dynamic and affected by abiotic factors (light and ambient O$_2$ level) and different responses are observed for different sections of the rhizosphere. The use of high resolution planar optodes greatly improves the possibility of understanding and mapping these dynamics and furthermore...
underpin that single point or single profile measurements, cannot be extrapolated to the entire rhizosphere.

However, all planar optode imaging is performed along a wall, this will have implications for the measured concentration and distribution of O$_2$ (Wenzhofer and Glud 2004; Glud 2008; Meysman et al. 2010). As the wall does not consume O$_2$, the oxic zone surrounding the roots is wider than for roots only surrounded by soil. However, assuming that the oxic zone around roots along the wall can be approximated as half-spherical or half-cylinders and that the total oxic volume around roots are independent of the wall, the radial O$_2$ distribution for “undisturbed” roots can be calculated (Frederiksen and Glud 2006; Polerecky et al. 2006). The corrected width of the oxic zone is 29 to 41 % less (depending on the width of the oxic zone) than the width observed at the planar optodes. This correction is used in all the following calculations.

Influence of irradiance and ambient O$_2$ levels on rhizosphere O$_2$ dynamics

The O$_2$ release from the roots was clearly affected by changes in irradiance levels – suggesting that photosynthetic O$_2$ production is important for regulating radial O$_2$ loss (ROL). The observed mean increase of 57.7 % air sat. in the rhizosphere O$_2$ level during an increase in irradiance from 0 to 300 μmol photons m$^{-2}$ s$^{-1}$, is however lower than reported values from microsensors studies for submerged rice grown in artificial substrates. One study found an O$_2$ increase in rice root tissue 10 mm behind the root tip of ~500 % during a dark–light cycle with irradiance: 350–400 μmol photons m$^{-2}$ s$^{-1}$ (Colmer and Pedersen 2008). These values however represent measurements inside the tissue of plants grown in non-reducing agar medium. Likewise it have been reported that a dark–light transition (irradiance: ~900 μmol photons m$^{-2}$ s$^{-1}$), increased the O$_2$ in the root tissue by >300 %, for rice plants grown in nutrient solution (Waters et al. 1989). Only a few studies have, as the present one, investigated rhizosphere O$_2$ dynamics of plants growing in natural soil. One microsensor study, found no O$_2$ changes inside root tissues during a dark–light cycle (irradiance: 400–500 μmol photons m$^{-2}$ s$^{-1}$) for rice plants of similar age compared...
to the plants in our study (Revsbech et al. 1999). The explanation for this large discrepancy among studies is not clear, but it is apparent that the highest responses to light–dark shifts apparently are found for plants grown in artificial, non-reducing media and are largely based on single point measurements with O2 electrodes (Waters et al. 1989; Colmer and Pedersen 2008). One explanation is that comparison between studies is confounded by differences in plant morphology e.g., ROL barrier and experimental conditions of the respective studies. However, the variability of ROL observed in this study show that single-point measurements only give a limited insight in the true spatial heterogeneity, as large differences in the extent of rhizosphere oxygenation may exist within a single root system. In fact, analyzing the dark–light response (irradiance: 300 µmol photons m\(^{-2}\) s\(^{-1}\)) of individual pixels from a single root, we found an relative increase up to 195 %, while the average value across the oxygenated area of this individual root was only 31.2 %.

After the initial sharp drop in rhizosphere O2 at the onset of darkness, the subsequent steady increase in rhizosphere O2 levels and expanding oxic area (Fig. 2a) must be associated to I) concurrent reduction in root (and shoot) respiration and/or II) an alternative reduction in O2 respiration of microorganisms in the surrounding soil. The first point is supported by the evidence of reduced dark respiration in roots following substrate depletion in plant tissue after prolonged exposure to darkness (Waters et al. 1989; Colmer and Pedersen 2008). As the metabolic activity is highest in root tips, it is to be expected that shortage of substrates would occur there first. This is consistent with our finding that root tips and the basal root zone, hosting relatively large amounts of young metabolically active tissue, exhibit the fastest increase in O2 levels after the initial decline. Therefore, we suggest that reduced respiration in the root tissues, at least partly, could explain the observed increase of ROL during darkness. Likewise, reduced leakage of organic substrates from the root tip area would reduce soil metabolism and increase the ambient O2 availability during nighttime.

We cannot quantitatively discriminate between the importance of the two processes, but we suggest that the resolved dynamic is related to reduced O2 consumption in both the root and the surrounding soil. Both factors are presumably being regulated by the availability of plant derived exudates that in some cases have been demonstrated to decrease during dark periods (Watt and Evans 1999; Badri et al. 2010). The reduction in soil/root respiration, estimated from extracted O2 profiles, during a day-night transition amounted to 32 %, from 0.081 nmol cm\(^{-3}\) s\(^{-1}\) (±SD=0.5×10\(^{-2}\), n=6) to 0.064 nmol cm\(^{-3}\) s\(^{-1}\) (±SD=0.96×10\(^{-2}\), n=12) (Fig. 7).

Regulation of rhizosphere O2 dynamics

Variation in the O2 levels in the ambient air/water surrounding the plant imposed the largest O2 dynamics in the rhizosphere. For the plants investigated it apparently made little difference in rhizosphere O2 levels if plants were submerged or emerged. We do not know if the rice plants in our study had leaf gas film, previous studies have shown that this is genotypic dependent. However, the existence of such gas film could explain the apparent tolerance to submergence, as it would facilitate an efficient gas exchange between plant and water (Pedersen et al. 2009).

The net downward O2 transport is passive, and the degree of ROL is mainly controlled by the plants ability to adapt root morphology e.g., by elevated levels of suberin in the exodermis, lignin in the sclerenchyma and increased aerenchyma volume (Kotula et al. 2009). The rapid decrease in root O2 leakage from the main root axis and subsequently O2 leakage from the fine lateral roots, under otherwise constant environmental conditions (Figs. 4 and 5), most likely reflect a shift in root morphology and formation of a barrier to ROL. Such rapid development of a barrier to ROL have previously only been observed for roots transferred from oxic to anoxic conditions (Insalud et al. 2006). To our knowledge, the transition in O2 leakage from root axis to fine lateral roots has not previously been observed. However, capturing such transition is only achievable applying high resolution imaging approaches. From the BW images it was not possible to conclude if laterals were present already at Fig. 5a “0 h” or if laterals were formed during the image sequence. It have been proposed that fine lateral roots are the dominant surface for nutrient uptake (and remain permeable to O2) (Kirk 2003), thus it is plausible that the observed transition also reflects a change in root functioning.

The present study shows that ROL of root tips was particular sensitive to the ambient O2 level experienced by the aboveground plant, presumably as a result of intensified metabolism, longer transport distance from the soil surface and a large surface-to-volume ratio.
Rhizosphere O2 budget

Quantitative ROL rates can be calculated as: $J = \phi \times R_s \times L \times (L/2 \times A + 1)$, where $\phi$ is the soil porosity, $R_s$ the soil volume specific O2 uptake, $L$ equals half the width of the oxic zone (corrected for radial geometry, see above) and $A$ is the root diameter (Fenchel 1996). We used an average root diameter of ~0.32 mm as assessed from 6 images and 4 different roots and the value for $R_s$ calculated from the soil-water interface (see above). The calculated ROL exhibited considerable spatial variability. For instance did the ROL, as calculated from the profiles in Fig. 3c, f, range spatially from 10.6 to 0.46 mmol m$^{-2}$ d$^{-1}$, with an average of 5.22 mmol m$^{-2}$ d$^{-1}$. Extrapolating this procedure to all observed single roots of the image in Fig. 3, provided an ROL of 7.7 mmol m$^{-2}$ d$^{-1}$ (n=259 profiles) for all visible roots in Fig. 3b. Although not all roots are accessible for analysis (as some roots must grow in the interior of the rhizotron) a plant specific ROL of 0.27 mmol d$^{-1}$ can be calculated using previously estimated plant specific root surface area of 3.5×10$^{-2}$ m$^{-2}$ plant$^{-1}$ (Miyamoto et al. 2001). These findings are comparable to previously reported ROL values of 0.24 to 1.11 mmol d$^{-1}$ for 3 to 6 weeks old rice plants grown in soil (Revsbech et al. 1999; Cheng et al. 2014), but are substantially higher than rates derived from rice plants grown in stagnant hydroponics (Deng et al. 2010; Wu et al. 2012) with ROL rates of 1.3×10$^{-2}$ and 2.3×10$^{-2}$ mmol d$^{-1}$, respectively. It should be noted that calculations by other authors (Revsbech et al. 1999) and our present study, assume that oxic zones of individual roots do not merge. As can be observed on Figs. 1 and 3, that is not the case at the central root base and the assessments should thus be regarded as a maximum value.

Assuming densities of 40 to 100 plants m$^{-2}$ (as found in typical paddy rice fields (Xu et al. 2006)), root induced O2 leakage accounts for 10.8 to 27.0 mmol m$^{-2}$ d$^{-1}$, at 100 % air saturation in the ambient air/water and at an irradiance of ~200 μmol photons m$^{-2}$ s$^{-1}$. Corresponding night time values would be 8.7 to 21.8 mmol m$^{-2}$ d$^{-1}$. Calculated on a 24 h time scale (14:10 h, light–dark cycle) total soil O2 uptake in the rhizosphere amount to 144 % of the diffusive O2 uptake at the soil-water interface, indicating a considerable rhizosphere-mediated gas exchange in rice paddies.

Altogether, our rates compare well to previously reported values with daytime O2 uptake in the order of 30 mmol m$^{-2}$ d$^{-1}$ (125 plants m$^{-2}$) (Revsbech et al. 1999).

The oxic volume amounts to 14.9 to 18.7 cm$^3$ per plant, for night and daytime, respectively, which for natural plant densities scales (40 to 100 plants m$^{-2}$) to 748 to 1870 cm$^3$ m$^{-2}$ and 604 to 1511 cm$^3$ m$^{-2}$, for day- and nighttime, respectively. Rice paddies are usually exposed to higher light and temperature than applied...
in our study. Hence, the absolute values on ROL and O₂ consumption are therefore not directly comparable to most in situ conditions. Nevertheless, we argue that the conclusion on extensive temporal and spatial variations in ROL of rice plants still is valid. In fact, natural conditions with ever changing conditions in light and temperature would favor an even higher degree of variability.

Most roots showed ROL, and only 5–10 % of roots showed no traces of O₂ leakage – even at maximum light levels and 100 % ambient air saturation. The fact that most roots of young rice leaks O₂ aligns with previous observations (Armstrong 1994; Revsbech et al. 1999; Williams et al. 2014). It could be speculated that for young small plants with short roots, O₂ transport at the prevailing conditions is sufficient to ensure O₂ to the root-tips and that no morphological adaptations to restrain ROL is required or that short roots might not yet be able for form a barrier to ROL owing to the development stage for the tissue (Shiono et al. 2011).

Metal plaque distribution in the rhizosphere

Metal plaque images showed a high degree of spatial variation, most likely reflecting variations in O₂ availability (Fig. 6). Plaque formations mainly reflects Fe oxidation and have been considered to be predominately abiotically mediated (Mendelssohn et al. 1995), suggesting a direct link between O₂ leakage and plaque formation (Wu et al. 2012; Cheng et al. 2014). Our investigations showed that metal plaques were predominantly found in a millimeter wide zone around the roots, corresponding to the observed width of the oxic zone, but were heterogeneously distributed along the root in contrast to the observed continuous oxic zone along roots. However, as root metal plaques can persist in anoxic soil for months (Weiss et al. 2005) a direct temporal link between ROL and plaque distribution at any given time cannot be expected.

We did not measure metal plaque distribution and O₂ dynamics on the same plant, but the potential of combining real time imaging of O₂ and metal plaque distribution/dynamics could contribute to the understanding the heterogeneity in plaque dynamics and associated trace metal cycling (Deng et al. 2010; Seyfferth et al. 2010; Cheng et al. 2014).

By applying high resolution O₂ imaging we were able so resolve how irradiance, ambient O₂ levels and plant development affected the O₂ dynamics across the rhizosphere. The data demonstrated that the spatio-temporal dynamic in the rhizosphere of rice grown in submerged soil is substantially larger than previously anticipated. This underpins that extrapolations based on single point measurements within the rhizosphere, should be done with caution, as they likely provide an oversimplified view on the rhizosphere processes. This improved understanding of the magnitude and scale of O₂ dynamics can help to advance the understanding of the biogeochemical cycling the rice rhizosphere.

In the presented study we used a luminescent lifetime planar optode imaging approach but it should be noted that comparable data quality can be achieved using more simple and inexpensive imaging approaches (Zhu et al. 2006; Larsen et al. 2011; Tschiersch et al. 2011). With such systems high resolution measurements of O₂, pH and pCO₂ can be realized, greatly enhancing the possibilities to study the spatially complex rhizosphere. Furthermore, planar optode imaging can be combined with DGT measurements (Diffusive Gradient in Thin films) for simultaneously measurements of O₂/pH and trace metal fluxes (Stahl et al. 2012; Williams et al. 2014).

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