The contrasting role of auxin in submergence-induced petiole elongation in two species from frequently flooded wetlands

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The involvement of auxin in the submergence-induced petiole elongation has been investigated in *Rumex palustris* and *Ranunculus sceleratus*. Both wetland species are capable of enhanced petiole elongation upon submergence or treatment with exogenous ethylene (5 μL L⁻¹). Treatment of intact *Rumex palustris* plants with 1-naphthalene acetic acid (NAA) at 10⁻⁴ M enhanced petiole elongation, while treatment with N-1-naphthylphthalamic acid (NPA) had no effect on petiole elongation. The elongation response after NAA or NPA treatment was comparable for plants in both submerged and drained conditions. Pre-ageing of detached petioles of *Rumex palustris* for 3 h in light or in dark conditions had no effect on the submergence-induced elongation. In comparison to intact plants, detached petioles of *Rumex palustris*, with or without lamina, did not show significant differences in responsiveness to IAA between drained or submerged conditions. This was in contrast to *Ranunculus sceleratus* where submergence caused a clear increase in responsiveness towards IAA. Removal of the lamina, the putative source of auxin, or treatment with NPA did not hinder the submergence-induced elongation of detached *Rumex palustris* petioles, but severely inhibited elongation of detached *Ranunculus sceleratus* petioles. This inhibition could be restored by application of NAA, suggesting the specific involvement of auxin in the submergence response of *Ranunculus sceleratus*. It is concluded that, in contrast to *Ranunculus sceleratus*, auxin is probably not involved in the submergence-induced petiole elongation of *Rumex palustris*.

Key words – Auxin, ethylene, *Ranunculus sceleratus*, *Rumex palustris*, submergence.

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Introduction

One of the adaptative features of plants to submergence is the stimulated elongation of petioles, stems or internodes, enabling plant structures to emerge from the water. This elongation is causally related to increased levels of endogenous ethylene found under submerged conditions (Ku et al. 1970, Musgrave et al. 1972, Voesenek et al. 1993) and leads to a relief of oxygen deficits, thus ensuring survival and sexual reproduction (Van der Sman et al. 1991, Voesenek and Van der Veen 1994). Enhanced petiole elongation and its relation to ethylene has been well described for the flooding resistant rosette plant *Rumex palustris*, occurring at low-elevation, frequently-flooded sites in river floodplains (Blom et al. 1994). However, the mechanism by which ethylene stimulates shoot elongation in *Rumex palustris* is still largely unknown. For a number of aquatic and semi-aquatic species (*Regnellidium diphyllum*, Walters and Osborne 1979; *Ranunculus sceleratus*, Horton and Samarakoön 1982; *Nymphoides peltata*, Malone and Ridge 1983) the presence of auxin has been shown to be prerequisite for optimal submergence-induced elongation. The similarity between *Rumex palustris* and *Ranunculus sceleratus* in submergence response, growth form (both are dicotyledonous rosette species) and habitat (frequently flooded sites) led to the assumption that auxin would also play a role in *Rumex palustris*. Therefore, the overall aim of this study was to determine whether auxin was also involved in submergence-induced petiole growth of...
**Rumex palustris**. In order to compare and validate our results, both *Rumex palustris* and *Ranunculus sceleratus* were used in similar experiments, since for the latter species auxin involvement has been clearly demonstrated (Musgrave and Walters 1973, Cookson and Osborne 1978, Horton and Samaraskoon 1982, Smulders and Horton 1991). The main objectives were to determine whether (1) the submergence-induced petiole elongation of *Rumex palustris* can be arrested by application of an auxin transport inhibitor, ageing of petiole tissue or removal of the lamina, (2) submergence results in a change in petiole responsiveness of *Rumex palustris* towards auxin, and (3) the involvement of auxin in the submergence response of *Rumex palustris* and *Ranunculus sceleratus* is comparable.

**Abbreviations** — NAA, 1-naphthaleneacetic acid; NPA, N-1-naphthylphthalamic acid.

**Materials and methods**

**Chemicals**

Indoleacetic acid (IAA) and 1-naphthaleneacetic acid (NAA) were purchased from Sigma Chemical Company. N-1-naphthylphthalamic acid (NPA) was synthesized in the Dept of Organic Biochemistry at the Univ. of Nijmegen. IAA, NAA and NPA were first dissolved in a few drops of ethanol (96%). Controls received an equal amount of ethanol. Ethylene (Hoekloos, mixture in medical air) was applied at a growth saturating concentration of 5 µl l⁻¹ (Voesenek and Blom 1989).

**Plant material**

Seeds (achenes) of *Rumex palustris*, collected in river floodplains, were germinated on moistened filter paper in Petri dishes for 10 days in a germination cabinet (16-h photoperiod, day/night temperature of 25/10°C, photosynthetic photon flux density, PPFD, of 30 µmol m⁻² s⁻¹ from fluorescent lamps, Philips TL 8W/33). Seedlings were planted in plastic pots (volume 60 ml) filled with a mixture of sand and potting compost (1:1, v/v). Seedlings of *Ranunculus sceleratus* were planted in plastic pots (volume 150 ml) filled with the same substrate. Both plant species were grown for 4–5 weeks in a growth chamber (16-h photoperiod, day/night temperature of 21/15°C, PPFD of 95 µmol m⁻² s⁻¹ from fluorescent lamps, Philips TLD 58W/84). Plants were selected for experiments on the basis of homogeneity of developmental stage of the 5th leaf.

**Submergence and ethylene response**

Intact *Rumex palustris* and *Ranunculus sceleratus* plants were submerged in 200-l tanks filled with tapwater (22°C) placed in the greenhouse (16-h photoperiod, day/night temperature of 22/16 ± 2°C) with additional illumination (150 µmol m⁻² s⁻¹ at water surface). The plants were approximately 50 cm below the water surface. Exogenous ethylene was applied in an air-tight growth cabinet, located in the greenhouse, at ambient temperature and with additional illumination (80 µmol m⁻² s⁻¹ from fluorescent lamps, Philips TL 8W/33, 16-h photoperiod, day/night temperature of 23/17 ± 2°C). Enough pure ethylene was initially injected into the cabinet to give a final concentration of 5 µl l⁻¹ ethylene in air. Subsequently, ethylene at a concentration of 5 µl l⁻¹ in air was then flushed through the system at a rate of 20 l h⁻¹. Drained plants were maintained in the greenhouse. Petiole elongation was measured after 48 h in leaves of three different developmental stages: (1) the youngest leaf with still furled lamina, (2) an intermediate stage with partly unfurled lamina, and (3) an old leaf with fully-expanded lamina.

**Auxin and NPA dose response experiments**

Intact *Rumex palustris* plants were submerged in 10 mM K-phosphate buffer (pH 6.0, tapwater) or in buffered NAA (10⁻⁵–10⁻⁴ M) or NPA (10⁻⁴–10⁻³ M) solutions in 20-l glass containers. Drained plants were sprayed until runoff once every 24 h with the same solutions used for submergence. Both submerged and drained plants were kept in the growth chamber mentioned in Plant material. Elongation of petioles of leaves in the intermediate stage was measured 48 h after the onset of experiments with a ruler to the nearest 0.5 mm.

**Experiments with detached petioles**

*Ranunculus sceleratus* petioles were detached according to Samaraskoon and Horton (1983) by removing a leaf with fully expanded lamina and cutting off the petiole at 25 mm from the lamina. For drained conditions of detached petioles with lamina attached, petioles were inserted through Paraflim, spread over a 100-nil beaker, into the test solution, thus leaving the lamina in air (cf. Musgrave and Walters 1973). Petioles without lamina were floated as another drained control. For submerged conditions, petioles (with or without lamina) were weighted by gently clamping the petiole in a latex serum cap and submerged in 1/l test solution in a 1-l glass beaker. For *Rumex palustris* a different method was followed. Preliminary experiments showed that floating or partly submerged petioles also displayed enhanced elongation and that removal of the root system had no effect on the enhanced petiole elongation at submergence (results not shown). Detached *Rumex palustris* petioles were obtained by removal of the leaves 1–4 (petiole and lamina), leaf 6 and the apex. The root system was cut off approximately 10 mm below the rosette node and laterals on the remaining part of the tap root were removed. This left the 5th petiole and lamina attached to the rosette node and ca 10 mm tap root. The tap root remainders of the plantlets were put in latex serum caps. These serum caps were mounted in a PVC plate and placed in a 1-l
glass beaker. The beaker was filled to root level for drained experimental conditions (ca 200 ml) or with 1 l of test solution for submergence.

For *Rumex palustris*, ageing of detached petioles without lamina was performed by positioning the petioles upright with the root remainder in distilled water in 1-l glass containers covered with plastic foil or wrapped in aluminium foil for dark treatment. For *Ranunculus sceleratus*, petioles were floated on distilled water. In all experiments with detached petioles, with or without lamina attached, plant material was aged for 3 h.

Elongation of petiole segments of detached *Rumex palustris* and *Ranunculus sceleratus* petioles was measured by making marks with waterproof ink every 2 mm. Petioles were either kept drained or submerged in K-phosphate buffer (10 mM, pH 6.0) with and without lamina. Segment length was measured again after 48 h with a ruler to the nearest 0.5 mm.

For other experiments with detached petioles, test solutions were 10⁻²–10⁻⁵ M IAA in 10 mM K-phosphate buffer (pH 6.0) and 10⁻³ M NAA, 10⁻⁵ M NPA or a combination of NAA/NPA in 10 mM K-phosphate buffer (pH 6.0). Ethylene (5 μl l⁻¹) was applied continuously in a flow-through system at 1 l h⁻¹ in gas-tight 2-l containers with an in- and outlet port and a sealed port for gas analysis. Unless stated otherwise, all experiments lasted 48 h. Petiole length was measured with a ruler to the nearest 0.5 mm at the beginning and end of the experiments.

**Statistics**

For each experiment 10–12 replicates were used unless stated otherwise. Experiments were repeated at least once. Representative data are shown. Differences between the means of the treatments were assessed with Tukey and LSD multiple comparison procedures after ANOVA.

**Results**

**Submergence and ethylene**

Submergence and exposure to 5 μl l⁻¹ ethylene significantly enhanced elongation for nearly all developmental stages of both species (Fig. 1). In *Rumex palustris*, treatment with ethylene partly mimicked the submergence response. In *Ranunculus sceleratus* petiole elongation of all three developmental stages was significantly higher upon ethylene treatment compared to submergence (Fig. 1). In both species petiole elongation upon submergence was most pronounced in leaves at the intermediate developmental stage. Therefore, this developmental stage was used in subsequent experiments.

**Effect of NAA and NPA on petiole elongation of intact plants**

Intact *Rumex palustris* plants were submerged in buffered aqueous solutions of 10⁻⁶–10⁻⁴ M NAA while drained plants were sprayed daily with an identical range of NAA concentrations. There was no significant petiole elongation response to auxin at 10⁻⁴ and 10⁻⁵ M NAA over a 48-h period (Fig. 2A). Elongation was significantly stimulated by 10⁻⁴ M NAA and this effect was comparable in both submerged and drained conditions.

In *Rumex palustris* plants, application of the auxin transport inhibitor NPA did not affect growth at concentrations of 10⁻⁷ to 10⁻⁴ M, whether sprayed on drained plants or for plants submerged in NPA solution (Fig. 2B). Extra pretreatment with NPA 3 h before submergence in a NPA solution of the same concentration produced comparable results as shown in Fig. 2B (data not shown).

However, it can not be excluded that high endogenous auxin levels interfered with exogenous auxin application and concealed these results. Therefore, in subsequent experiments petioles were detached and aged for 3 h in an attempt to minimize endogenous auxin levels (cf. Malone and Ridge 1983).

**Effect of IAA, NAA and NPA on elongation of detached petioles**

Detachment of *Rumex palustris* petioles reduced the response towards ethylene and submergence when com-
Fig. 2. Effect of NAA (A) and NPA (B) on petiole elongation of submerged and drained intact *Rumex palustris* plants (n=10; ±SE) over a 48-h period. Plants were either sprayed with NAA or NPA in 10 mM P-buffer (pH 6.0) or submerged in the same solutions. All solutions, including controls contained 0.1% ethanol. Spraying was repeated after 24 h. Bars represent SE for each treatment (P<0.05).

Tab. 1. The effect of submergence and ethylene (5 µl l⁻¹) on petiole elongation of intact and detached *Rumex palustris* petioles (n=12). Elongation was recorded after 48 h. Different letters indicate significantly different means (ANOVA with Tukey's post test, P<0.05).

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Detached petiole (mm ± SE)</th>
<th>Intact plant (mm ± SE)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Drained</td>
<td>4.6±0.4a</td>
<td>4.9±0.6a</td>
</tr>
<tr>
<td>Ethylene (5 µl l⁻¹)</td>
<td>11.5±0.6b</td>
<td>22.2±0.9d</td>
</tr>
<tr>
<td>Submerged</td>
<td>16.3±1.2c</td>
<td>23.8±1.0d</td>
</tr>
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Tab. 2. The effect of ageing under light and dark conditions on petiole elongation of detached *Rumex palustris* petioles (n=10). Ageing was performed by standing petioles upright for 3 h with root remainder in distilled water in 1-l glass containers covered with plastic foil or wrapped in aluminium foil. Petiole length was measured before and after 48 h submergence. Different letters indicate significantly different means (ANOVA with Tukey's post test, P<0.05).

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Petiole elongation (mm ± SE)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Drained</td>
<td></td>
</tr>
<tr>
<td>Not aged</td>
<td>5.0±0.4a</td>
</tr>
<tr>
<td>Aged in light</td>
<td>3.3±0.6a</td>
</tr>
<tr>
<td>Aged in dark</td>
<td>2.6±0.6a</td>
</tr>
<tr>
<td>Submerged</td>
<td></td>
</tr>
<tr>
<td>Not aged</td>
<td>9.4±0.5b</td>
</tr>
<tr>
<td>Aged in light</td>
<td>9.2±0.9b</td>
</tr>
<tr>
<td>Aged in dark</td>
<td>8.6±0.4b</td>
</tr>
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equal, while *Ranunculus sceleratus* segments closest to the lamina elongated significantly more than basal segments. Removal of the lamina did not affect this pattern in petiole elongation for either species, but drastically reduced overall elongation in *Ranunculus sceleratus*. This in contrast to *Rumex palustris*, where petiole elongation response only was slightly reduced (Fig. 3).

Detached petioles of *Rumex palustris* were less responsive to applied IAA than those of *Ranunculus sceleratus* (Fig. 4). Submergence increased the responsive-
petioles with lamina also suppressed petiole elongation significantly. This growth reduction could be overruled by application of $10^{-5} M$ NAA, thus showing the specific involvement of auxin in the submergence response of *Ranunculus sceleratus*.

**Discussion**

*Rumex palustris* and *Ranunculus sceleratus* are both dicotyledonous rosette plants occurring in river floodplains at frequently flooded sites. Both species are adapted to flooding by their ability to initiate rapid petiole elongation, especially those of the younger leaves. This elongation response can be mimicked to a large extent by exogenous application of ethylene and these results are in agreement with earlier reports (Musgrave and Walters 1973, Ridge 1985, Voesenek and Blom 1989).

While the morphological response upon submergence is similar for *Rumex palustris* and *Ranunculus sceleratus*, the results of the present study demonstrate that the involvement of auxin clearly differs in these two species. In *Rumex palustris* the responsiveness of petioles to exogenously applied auxin did not differ between drained or submerged conditions and the response was
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elongation in either intact plants or detached petioles. Sceleratus petioles or enough active auxin remains in the tissue after detachment and ageing. Floating of petioles on 10^{-4} M buffered NAA or IAA solution caused a 10- and 25-fold increase, respectively, in ethylene production compared to the control (results not shown). Furthermore, submergence of petioles in NAA or IAA caused a swelling of the petiole tissue that showed a strong linear relationship with the applied NAA or IAA concentration (results not shown). Both phenomena demonstrate that auxin is perceived by the petiole tissue even at low exogenous concentrations. Since intact plants and detached petioles were completely submerged in NAA or IAA solution, assumption (1) can probably be neglected. For assumption (2) no definite conclusion could be made without determination of endogenous auxin. Although, in an effort to delineate petiole tissue of endogenous hormones, petioles were aged in both light and dark without significant effect on submergence response (Tab. 2), rejecting assumption (2) would be disputable. Consequently, regarding elongation, Rumex palustris petioles are either less susceptible towards auxin than Ranunculus sceleratus petioles or enough active auxin remains in the tissue after detachment and ageing.

More important, in Rumex palustris, the inhibition of auxin transport with NPA did not cause a reduction in elongation in either intact plants or detached petioles. NPA has been shown previously to be active in inhibiting polar auxin transport in Rumex palustris: next to petiole elongation, submergence or flooding also initiates the formation of adventitious roots, a process that is auxin mediated and arrested by application of NPA (Visser et al. 1995). Like NPA, removal of the lamina of detached petioles, the putative source of auxin, had no effect on petiole elongation under submersed conditions. In contrast, elongation in Ranunculus sceleratus during submergence was significantly inhibited by application of auxin transport inhibitors (Fig. 5: Horton and Samarakoon 1982) or by removal of the lamina (Fig. 5: Samarakoon and Horton 1983). This inhibiting effect on elongation could be counteracted by application of 10^{-6} M IAA or 10^{-5} M NAA, showing that the inhibition is auxin-specific and demonstrating the obligatory presence of auxin in submergence elicited petiole elongation. Similar results were reported for Regnellidium diphyllum (Walters and Osborne 1979) and Nymphoides peltata (Malone and Ridge 1983).

Under drained conditions, the growth of petiole segments was found to be evenly distributed over the petiole for both Rumex palustris and Ranunculus sceleratus. If auxin is involved in this growth pattern, two explanations seem possible: (1) a classical longitudinal gradient in auxin content exists (high in tip, low at basal parts, Sanchez-Bravo et al. 1993) with an opposing gradient in responsiveness (low at tip, high at basal part, as mentioned in Hoson et al. 1992) resulting in a similar growth rate along the petiole, or (2) both auxin content and tissue responsiveness are evenly distributed over the petiole. Upon submergence the growth pattern did not change for Rumex palustris, unlike in Ranunculus sceleratus, where segmental growth declined along the apical-basal gradient. A comparable gradient was found for epidermal cell length in submerged or ethylene treated Ranunculus repens petioles (Ridge 1985). It is difficult to explain this discrepancy due to several, mutually influencing factors like variations in endogenous auxin levels between different organs (high in young leaves, low in older leaves or vice versa; Ueda et al. 1991 and references therein) or possible changes in auxin transport velocity or auxin responsiveness. For Ranunculus sceleratus it was shown that treatment with 50 μl l^{-1} ethylene increased auxin transport velocity significantly (Musgrave and Walters 1973). This would mean that basal petiole parts receive more auxin and thus should show an enhanced growth compared to the apical petiole segments. However, our results show an opposing growth pattern even after removal of the lamina. Therefore, we hypothesise that in Ranunculus sceleratus submergence or ethylene causes an increase in responsiveness towards endogenous auxin of especially the apical segments and that a similar mechanism is lacking in Rumex palustris.

Together our results indicate that, in contrast to Ranunculus sceleratus, involvement of auxin in the submergence response of Rumex palustris is at least uncertain. This would be, as far as we know the first time an auxin independent ethylene-mediated elongation is reported for a terrestrial, dicotyledonous plant. Noteworthy is the phenomenon of enhanced petiole elongation of Rumex palustris at floating or partially submerged conditions, contrasted to Ranunculus sceleratus. Rumex palustris shares this feature with deep water rice. Over the past ten years, Kende and co-workers showed that growth of deepwater rice under submerged conditions is elicited by gibberellin (GA) via an ethylene mediated change in responsiveness towards GA (Raskin and Kende 1984) and an enhanced endogenous GA concentration (Hoffmann-Benning and Kende 1992). Further experiments will show whether or not GA is involved in the submergence response of Rumex palustris.

In conclusion, it is shown that auxin does not play a crucial role in the submergence response of Rumex palustris, while in Ranunculus sceleratus previous findings have been confirmed and extended with respect to the elongation pattern down the petiole. Therefore, with respect to auxin, the hormonal mechanism of submergence-induced petiole elongation in Rumex palustris...
appears to be different from that in *Ranunculus sceleratus*.

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**References**


