Flooding regimes and life-history characteristics of short-lived species in river forelands

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Summary

1. A large-scale outdoor experiment was carried out to study the effect of flooding on growth, survival and reproduction in relation to developmental stage in three short-lived species. Several cohorts of *Chenopodium rubrum*, *Rumex maritimus* and *R. palustris* were raised in accordance with a flooding regime which was based on the average flooding conditions of their natural habitats in river forelands.

2. Survival of submergence in the pre-reproductive phase was high in both species of *Rumex*, but relatively low in *C. rubrum*. Biomass reduction following flooding depended on plant size before flooding and on the mean temperature of the flood-water.

3. Both *Rumex* spp. were less reduced in size during flooding, and better capable of regeneration afterwards, than *C. rubrum*.

4. Later-raised cohorts of the short-day species *C. rubrum* started to flower after a shorter time and at an earlier developmental stage than earlier cohorts. Flooding had little effect on probability of flowering and caused only a slight delay in flowering in this species. In both long-day species of *Rumex*, flowering stems were not initiated until a minimal leaf number was formed. As a result later cohorts remained vegetative, while flooding delayed flowering until later in the season or even until the following year.

5. All flooded plants of *C. rubrum* and most flooded plants in early cohorts of *R. maritimus* attempted to reproduce in the first growing season. Seed number was severely reduced in *C. rubrum* due to both reduction of plant biomass and destruction of flowers during flooding. In flooded plants of *R. maritimus*, seed output was correlated with plant size at the time of bolting. Seed number was maximized at the expense of seed size in *R. maritimus*.

6. These results suggest that there is a trade-off between survival of flooding and early reproduction in these species, and that there are complex interrelationships between highly variable environmental factors and developmental factors which determine the survival and reproductive success of short-lived species in river forelands.

Key-words: *Chenopodium rubrum*, flowering, growth, *Rumex maritimus*, *Rumex palustris*, seed production, survival

Introduction

Flooded zones along rivers in the Netherlands are bounded by the main dykes at either side of the river. These ribbon-like zones, which are up to several kilometres wide, are referred to as river forelands. Differences in elevation, both natural and man-made, combined with strongly fluctuating water levels resulting from unpredictable changes in river-water discharge, lead to highly dynamic hydrological regimes in space and in time in these areas (Van de Steeg 1984; Brock, Van der Velde & Van de Steeg 1987; Blom 1990; Blom et al. 1990). The occurrence of several *Rumex* species within restricted zones along elevational gradients in these forelands suggests that these species differ in flooding tolerance.

Comparative studies concerning different aspects of flooding tolerance in *Rumex* have revealed a dichotomy between species from high elevations...
on the one hand and species from low and inter-
mediate elevations on the other (e.g. Laan et al. 
1989; Voesenek & Blom 1989; Voesenek, Blom & 
These studies were mostly performed on young 
vegetative plants grown under controlled conditions 
where flooding was treated as a stress factor (sensu 
Grime 1979). In the lowest areas of river forelands, 
flooding will primarily be experienced as a dis-
turbance by short-lived ruderal species (Menges & 
Waller 1983). Floods may occur at several stages of 
the life cycle and these stages should all be included 
in studies of flooding tolerance. For example, Rumex 
species from frequently flooded areas have persistent 
seed banks, whereas species from higher elevations 
have transient ones (Voesenek & Blom 1992). A 
distinction between avoidance strategies, such as 
survival through periods of adverse conditions by 
means of persistent seeds, and tolerance strategies, 
such as metabolic adaptations, is extremely useful in 
this respect (Crawley 1986).

In order to study the interaction between natural 
flooding regimes and reproductive behaviour of 
established plants, three co-occurring species from 
low, frequently flooded areas of the river forelands 
were selected. Chenopodium rubrum (L.), Rumex 
maritimus L. and R. palustris Sm. are short-lived 
species with different life histories. Both Rumex 
spp. are annual if they reach a mature vegetative 
size before a certain photoperiod has passed (Van 
der Sman, Blom & Van de Steeg 1992). This photo-
period is longer for R. palustris, which displays the 
biennial strategy more often in the river forelands 
than does R. maritimus. In contrast to both Rumex 
spp., the short days of late summer and autumn 
promote early flowering in the strictly annual C. 
rubrum. In a glasshouse experiment, R. maritimus 
appeared to be much more tolerant towards flooding 
than C. rubrum (Van der Sman, Van Tongeren & 
Blom 1988), but duration, time and frequency of 
flooding were all factors that considerably modified 
growth, the onset of flowering and seed production 
within one or both species. Plant developmental 
stage has been shown to play an important role in 
the rapid under-water elongation response of shoots 
in R. maritimus (Van der Sman et al. 1991).

A large-scale outdoor experiment was designed 
to enable the simultaneous comparison during one 
growing season of several realistic flooding con-
ditions. Several cohorts of plants of C. rubrum, 
R. maritimus and R. palustris were raised in accord-
ance with these flooding regimes to study the effect 
of flooding on growth, survival and reproduction 
in relation to plant developmental stage. Species 
differences in flooding tolerance are discussed in 
relation to differences in life-history characteristics.

Materials and methods

**SPECIES HABITAT DESCRIPTION AND PLANT MATERIAL**

Chenopodium rubrum is one of the character-species 
of the Chenopodium glauco-rubi Lohm. and both 
Rumex maritimus and R. palustris are character-
species of the Ranunculo-Rumicetum maritimi Siss. 
(Westhoff & Den Held 1975). C. rubrum occurs as 
companion species in the Ranunculo-Rumicetum 
association and, likewise, both Rumex spp. occur in 
the Chenopodietum. In the river forelands several 
plant communities exist which may be interpreted 
as incomplete stages of these associations or as 
transitional stages between them (H.M. van de 
Steeg, unpublished relevés). The Chenopodietum 
may occur on exposed clay soils as well as sandy 
soils, especially at flood-marks on sandy river beaches, 
while the Rumicetum is more characteristic of wet 
clay soils. Both Rumex spp. also occur in associations 
of marsh species (Oenanthon; Westhoff & Den 
Held 1975). Germination of plants of C. rubrum 
and both Rumex spp. occurs after the subsidence 
of floods from April onwards and multiple post-
flooding cohorts are commonly observed as de-
scribed for R. palustris and R. crispus in Voesenek & 
Blom (1992). Maximum germination in the lab-
oratory is achieved under conditions of light and 
fluctuating temperatures in all three species. Such 
patterns appear to be common in wetland species 
(Thompson & Grime 1983).

The flooding regime used in the outdoor exper-
iment was derived from water-level measurements 
which were made over the period 1970–87 and 
analysed in relation to the elevation of three study 
areas near Nijmegen, the Netherlands. These were 
a clay-pit, a sandy river beach and a former river 
bed. These sites provide examples of the habitats in 
which the above-described plant communities occur 
in river forelands. The clay-pit area was partly 
protected by a summer dyke, which contains a sluice 
that is closed after the subsidence of winter/spring 
floods to prevent further flooding during the growing 
season. In unprotected areas, flooding frequency 
was highly variable between years but in years 
and sites in which winter floods subsided early 
(April–May), the mean number of floods was 3–4 
per growing season. Mean duration ranged from 10 
days of flooding in the highest sites (clay-pit area) to 
16 in the lowest (river-bed area). Water depth for 
the majority of floods was less than 1 m above the 
soil surface. In the river-bed area, extremely severe 
floods, i.e. those with a duration of more than 1 
month and maximum water levels of several metres 
above the soil, occurred approximately once every 3 
years. Such floods are lethal to established plants in 
the growing season (Voesenek 1990) and for this 
reason a moderate regime of four 10-day flooding
periods with a constant depth of 40 cm was chosen for the experiment. Seeds of *C. rubrum*, *R. maritimus* and *R. palustris* were harvested in 1986 in the clay-pit area inside the summer dyke. The total seed output from five plants per species was mixed and dry-stored in the laboratory.

**OUTDOOR FLOODING EXPERIMENT**

Uniform germination was achieved as described previously (e.g. Van der Sman *et al.* 1991). After 4 days, germinating seeds were transplanted into pots (16 cm in diameter × 50 cm high) filled with a 1:1 mixture (vol.) of river sand and peat. Seven seedlings were transplanted per pot and thinned to one seedling per pot after 2 weeks. Four cohorts, each of 50 pots species⁻¹, were raised in large basins of water (length × width × depth: 8 × 2 × 1 m). Four basins, of which two were flooded, were divided lengthwise and each cohort was split between two basin-halves with parallel plant arrangements in the one to be drained and the one to be flooded. The first cohort was sown on 14 May 1987, 32 days before the start of the first 10-day flooding period (Fig. 1). The following cohorts were sown at 6-week intervals, immediately after subsequent flooding periods, on 25 June, 6 August and 17 September, respectively. Drained conditions were provided by a water level of 5–10 cm in the basins. To simulate flooding according to the regime shown in Fig. 1, basins were filled with tap-water until the water level reached an overflow approximately 40 cm above the soil level in the pots. At this point the water depth in the basins was 90 cm. After 10 days water was pumped out to restore the water level to that provided by the drained conditions. Filling and emptying the basins both took approximately 8 h.

**Measurements**

Irradiation and temperature were continuously monitored during the study period. These parameters are presented for each cohort as radiation and temperature sums (with 10°C as base temperature; Van der Sman, Blom & Van de Steeg 1992) over the growth period up to first flooding. During flooding periods, water temperatures were measured twice daily, at approximately 08.00 and 17.00 hours. Before and after each flooding period (Fig. 1) the following parameters were measured for all plants: the cumulative number of leaves on the main shoot that were ≥1 cm in length (see Van der Sman, Blom & Van de Steeg 1992), the number of dead leaves on the main shoot, the number of axillary shoots, the length, width and petiole length of the largest leaf, and the length of the main stem. Before and after the first flooding period of each cohort, three plants per species and treatment were harvested to determine the dry weight of shoots, the maximum diameter (r) of tap roots, and the length (h) of this root if it was ≥1 mm in diameter. The volumes of tap roots were estimated from these parameters as 0.5πr²h. Of the remaining 20 plants per species, cohort and treatment, five randomly selected plants were measured every 10 or 11 days (Fig. 1) with respect to the above-mentioned non-destructive parameters, as well as the number of living leaves on axillary shoots (≥1 cm) and the lengths of axillary stems. Twice a week, the developmental stage of all plants was recorded, distinguishing the following stages: vegetative, bolting (*Rumex* spp.), flower buds visible, flowering, ripe fruits (*R. maritimus*; golden brown; *R. palustris*; dark red brown; *C. rubrum*, black). Ripe seeds (including perianths) were collected in November and weighed after sieving and cleaning several times. This was done for all reproductive plants of both *Rumex* spp. and for a maximum of 10 plants per cohort and treatment of *C. rubrum*. For each species, cohort and treatment, five samples of 100 seeds were taken from each of three plants. After removal of perianths, seeds were counted and weighed to determine mean seed weights. The empty perianths were also weighed.
Flooding and life history in river forelands

Germination characteristics

Seeds were incubated under the following conditions: constant temperatures of 10°C and 20°C, and fluctuating temperatures of 12 h at 20°C and 12 h at 10°C, both in the light (15 μmol m⁻² s⁻¹ PAR during the 12-h day) and in the dark. Three samples of 50 seeds each from a mixture of five plants per species, cohort and treatment were incubated in each of the above regimes in Petri dishes on filter paper (Schleicher & Schüll 595) watered with de-mineralized water. As soon as radicles appeared, germinated seeds were removed. After 3 weeks, ungerminated seeds were placed into the regime with fluctuating temperatures and light, which normally induced near 100% germination (Van der Sman, Blom & Van de Steeg 1992). Ungerminated seeds were stained with tetrazolium chloride (1% weight/vol., Sigma Chemical Co., St Louis, Miss., USA) which colours live tissue red (Moore 1972).

Data analysis

All tests were carried out with the SAS statistical package (SAS Institute Inc. 1985). Correlation analysis was performed using the Pearson rank correlation test. The General Linear Models procedure was used for all unbalanced analyses of variance. Significance of differences between means were tested using Student’s t-test. In C. rubrum and R. maritimus, between-plant differences in mean seed size were tested as a nested effect within the effects of cohort and flooding treatment. In C. rubrum, the effects of cohort and flooding treatment were not tested separately but as one main treatment effect because too many cells were empty in the case of flooded plants, due to plant death.

Results

Flooding tolerance in the pre-reproductive phase

Plant biomass is correlated with the product of the number of living leaves and maximum leaf length in these species (Van der Sman, Van Tongeren & Blom 1988; Voesenek 1990). These parameters were therefore chosen to represent plant growth in relation to radiation and temperature sums up to the first flooding period per cohort (Table 1). Growth rates appeared to depend on temperature during the larger part of the growing season (see also Van der Sman, Blom & Van de Steeg 1992). Radiation was probably only limiting in the last, September cohort (Table 1).

The initiation of leaves stopped during the period of first flooding for all cohorts, whereas leaf senescence on the main shoot was high only in the first three cohorts (Fig. 2). In addition, most axillary shoots died. In both Rumex species, the oldest leaves of the main shoot died first. In C. rubrum, old as well as young leaves died. In the June and August cohorts, mean above-ground dry weight was reduced immediately after the first flooding period by 90% in R. palustris, 95% in R. maritimus and 98% in C. rubrum, compared to drained plants of the same age.

<table>
<thead>
<tr>
<th>Cohort</th>
<th>May</th>
<th>June</th>
<th>August</th>
<th>September</th>
</tr>
</thead>
<tbody>
<tr>
<td>Temperature sum (°days)</td>
<td>103</td>
<td>254</td>
<td>238</td>
<td>92</td>
</tr>
<tr>
<td>Radiation sum (kJ cm⁻²)</td>
<td>40</td>
<td>52</td>
<td>35</td>
<td>22</td>
</tr>
<tr>
<td>Leaf number (≥1 cm)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cr</td>
<td>6.0 ± 0.0</td>
<td>32.8 ± 2.5</td>
<td>23.0 ± 2.4</td>
<td>0.0 ± 0.0</td>
</tr>
<tr>
<td>Rm</td>
<td>4.0 ± 0.0</td>
<td>21.2 ± 1.3</td>
<td>21.6 ± 0.8</td>
<td>2.0 ± 0.0</td>
</tr>
<tr>
<td>Rp</td>
<td>3.8 ± 0.2</td>
<td>23.6 ± 1.2</td>
<td>18.4 ± 0.4</td>
<td>2.0 ± 0.0</td>
</tr>
<tr>
<td>Leaf length (cm)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cr</td>
<td>2.0 ± 0.2</td>
<td>9.3 ± 0.6</td>
<td>5.2 ± 0.3</td>
<td>0.2 ± 0.1</td>
</tr>
<tr>
<td>Rm</td>
<td>2.4 ± 0.1</td>
<td>11.6 ± 0.5</td>
<td>9.8 ± 0.4</td>
<td>1.1 ± 0.1</td>
</tr>
<tr>
<td>Rp</td>
<td>2.1 ± 0.2</td>
<td>11.6 ± 0.4</td>
<td>9.0 ± 0.2</td>
<td>1.4 ± 0.2</td>
</tr>
<tr>
<td>Survival (%)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cr</td>
<td>0</td>
<td>65</td>
<td>50</td>
<td>95</td>
</tr>
<tr>
<td>Rm</td>
<td>85</td>
<td>100</td>
<td>100</td>
<td>100</td>
</tr>
<tr>
<td>Rp</td>
<td>100</td>
<td>100</td>
<td>100</td>
<td>100</td>
</tr>
<tr>
<td>Mean water temperature (°C)</td>
<td>14.2</td>
<td>17.1</td>
<td>17.8</td>
<td>10.3</td>
</tr>
</tbody>
</table>

* Base temperature 10°C.
Fig. 2. Mean cumulative initiation (closed symbols) and loss (open symbols) of primary leaves during the course of the outdoor experiment (n = 5 unless otherwise indicated). The September cohort is omitted. Shaded areas indicate flooding periods. (●) Drained plants; (■) vegetative flooded plants; (●) reproductive flooded plants. Reproductive development of the main shoot is indicated with letters: S = stem elongation (Rumex spp.), B = appearance of buds, F = in full flower, O = out of flower, R = ripe fruits (R* = incompletely ripened).

Vegetative growth in relation to the flooding regime

During submergence periods, the orientation of leaves and branches changed from horizontal to vertical within a day in all three species. In addition, all vegetative plants of both Rumex species exhibited fast extension of the petioles of young leaves (Fig. 3). The water surface was not reached by any of the plants. The extended leaves were easily damaged and did not survive for more than a few days or, exceptionally, a few weeks after submergence. After submergence, plants of both Rumex spp. resumed normal growth patterns. Plants of C. rubrum recovered from submergence by developing new leaves from the apex and/or a few scattered leaf axils but they exhibited a stunted growth habit, quite different from that of drained plants. Plants of all three species which survived the first flooding period also survived subsequent flooding periods.

Reproductive development in relation to the flooding regime

Transition to the reproductive phase started with rapid stem elongation in the cauline species C. rubrum as well as in the two rosette species of Rumex (Fig. 4; see also Bernier, Kinet & Sachs 1981). All surviving plants in the first three cohorts of C. rubrum flowered regardless of treatment (Table 2). Some plants in the September cohort initiated flower buds but they were unable to produce seeds before they were killed by a short period of frost in December. Flowering was not much delayed by flooding in C. rubrum (Fig. 2).

Flowering occurred in the May and June cohorts of R. maritimus and in the May cohort of R. palustris (Table 2). Seed output was correlated with the number and size of axillary stems in drained plants.
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Fig. 3. Mean maximum leaf (closed symbols) and petiole (open symbols) lengths in the course of the outdoor experiment in both Rumex species (n = 5 unless otherwise indicated). The September cohort is omitted. Shaded areas indicate flooding periods. (●) Drained plants; (●) vegetative flooded plants; (●) reproductive flooded plants.

of both species (Table 3). Flooded plants of R. palustris all remained vegetative, while percentage of plants flowering was reduced in flooded plants of R. maritimus (Table 2). Flowering occurred after approximately 12 primary leaves were initiated in drained as well as flooded plants of R. maritimus (see also Van der Sman, Blom & Van de Steeg 1992). At this stage, flooded plants possessed fewer (≤4) axillary leaves than drained plants (c. 45), and they did not produce axillary stems. Plants of both Rumex spp. which remained vegetative either reached the 12-leaf stage, at which flowering could commence, too late in the season, or did not reach this stage at all before the onset of the winter (September cohort).

Submergence reduced the final size of reproductive plants compared to drained conditions. Flowering plants of R. maritimus produced approximately the same total number of primary leaves under flooded as under drained conditions (Fig. 2), but stems of flooded plants remained significantly smaller (Fig. 4), whilst plants of C. rubrum were reduced in both respects after flooding. In addition, submergence of flowering plants resulted in severe loss of flowers and only a small number of seeds was produced by flooded plants (Table 4). Only two plants of C. rubrum in the August cohort, which was in full flower just before the start of the last flooding period, produced a few large seeds. In R. maritimus, however, new flowering shoots developed from axillary nodes of stem leaves after flooding. This resulted in a relatively constant seed output which was correlated with plant size (biomass estimated from the product of the number of living leaves and maximum leaf length) at the time of bolting (Pearson rank correlation coefficient 0.78, n = 22, P < 0.001).

In R. maritimus, mean seed weight was significantly lower in flooded plants than in drained plants (Table 4). Fruits of flooded plants appeared to be incompletely ripened (Fig. 2) and possessed seeds of a lighter colour than drained plants. Variation in seed weight was largely determined by flooding

<table>
<thead>
<tr>
<th>Table 2. Percentage flowering per cohort and treatment in Chenopodium rubrum, Rumex maritimus and R. palustris (n = number of plants which survived up to the flowering stage)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>C. rubrum</strong></td>
</tr>
<tr>
<td></td>
</tr>
<tr>
<td><strong>Cohort</strong></td>
</tr>
<tr>
<td>May</td>
</tr>
<tr>
<td>June</td>
</tr>
<tr>
<td>August</td>
</tr>
<tr>
<td>September</td>
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</table>
treatment (73%) and inter-plant differences (20%) in *R. maritimus*; less than 1% of the total sum of squares was attributable to cohort effect. In *C. rubrum*, only half the number of the flooded plants in the June cohort produced many seeds (Table 4). Mean seed weight was significantly higher in the flooded plants but variation in seed weight was only partly determined by overall treatment effect (51%) and inter-plant differences (2%) in *C. rubrum*.

Under fluctuating temperatures in the light, maximum germination was reduced in seeds from flooded plants (≥80%) compared to seeds from drained plants (≥98%) in both *C. rubrum* and *R. maritimus*. Ungerminated seeds appeared to be dead. Under all other conditions, seeds from flooded plants showed higher germination percentages than those of drained plants. At constant temperatures in the light, seeds from flooded plants showed 10–30% germination and from drained plants 0–10%. In the dark, seeds from flooded plants showed approximately the same germination percentages as in the light under all temperature

Table 3. Correlations (Pearson rank correlation coefficient) between stem parameters and total seed weight per plant in drained plants of *Rumex* spp. grown under drained conditions.

<table>
<thead>
<tr>
<th></th>
<th><em>R. palustris</em></th>
<th><em>R. maritimus</em></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td><em>(n = 19)</em></td>
<td><em>(n = 40)</em></td>
</tr>
<tr>
<td>Axillary stem number</td>
<td>0-73***</td>
<td>0-81***</td>
</tr>
<tr>
<td>Σ Axillary stem length</td>
<td>0-65**</td>
<td>0-88***</td>
</tr>
<tr>
<td>Primary stem length</td>
<td>0-39</td>
<td>-0-16</td>
</tr>
<tr>
<td>Σ Total stem length</td>
<td>0-70***</td>
<td>0-89***</td>
</tr>
</tbody>
</table>

** p<0-01, *** p<0-001.

Table 4. Mean (±SE) estimated seed number per plant and individual seed weight per cohort and treatment of *Chenopodium rubrum*, *Rumex maritimus* and *R. palustris*. The nested model explained 54% of the variation in seed weight in *C. rubrum* and 93% in *R. maritimus* (see text). Values followed by the same superscript letter are not significantly different.

<table>
<thead>
<tr>
<th>Cohort</th>
<th>Treatment</th>
<th>n</th>
<th>Estimated seed number</th>
<th>Individual seed weight (μg)</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>C. rubrum</em></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>May</td>
<td>drained</td>
<td>10</td>
<td>53 000 ± 2000a</td>
<td>61 ± 3b</td>
</tr>
<tr>
<td></td>
<td>flooded</td>
<td>0</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>June</td>
<td>drained</td>
<td>10</td>
<td>44 000 ± 1000b</td>
<td>65 ± 2b</td>
</tr>
<tr>
<td></td>
<td>flooded</td>
<td>6</td>
<td>990 ± 200c</td>
<td>85 ± 4a</td>
</tr>
<tr>
<td>August</td>
<td>drained</td>
<td>10</td>
<td>27 000 ± 3000d</td>
<td>92 ± 4a</td>
</tr>
<tr>
<td></td>
<td>flooded</td>
<td>2</td>
<td>≈10</td>
<td>—</td>
</tr>
<tr>
<td><em>R. maritimus</em></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>May</td>
<td>drained</td>
<td>20</td>
<td>23 000 ± 2000p</td>
<td>217 ± 7a</td>
</tr>
<tr>
<td></td>
<td>flooded</td>
<td>15</td>
<td>850 ± 160a</td>
<td>120 ± 5b</td>
</tr>
<tr>
<td>June</td>
<td>drained</td>
<td>20</td>
<td>13 000 ± 2000b</td>
<td>211 ± 7a</td>
</tr>
<tr>
<td></td>
<td>flooded</td>
<td>7</td>
<td>950 ± 240c</td>
<td>108 ± 10c</td>
</tr>
<tr>
<td><em>R. palustris</em></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>May</td>
<td>drained</td>
<td>19</td>
<td>8 200 ± 400</td>
<td>593 ± 5</td>
</tr>
</tbody>
</table>

* The remaining seven flowering plants produced ≤10 seeds.
† The remaining eight flowering plants did not produce seeds.
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Discussion

GROWTH AND SURVIVAL IN RELATION TO FLOODING

Upon submergence, plants of all three species showed the response of reorientation of leaves and branches from prostrate to vertical which is commonly observed in aquatic plants (Ridge 1987) and important in shallow water (Van der Sman, Van Tongeren & Blom 1988). In addition, rosette plants of both Rumex spp. exhibited petiole extension of younger leaves (Fig. 3). Survival of longer periods of submergence in the growing season depends on the restoration of shoot-air contact by means of rapid under-water shoot extension in several flooding-tolerant Rumex spp. (Voesenek & Blom 1989; Laan et al. 1990; Van der Sman et al. 1991). In C. rubrum, young a well as old leaves died during submergence, whereas in both Rumex spp., old leaves died first. The latter is of advantage with respect to growth of younger leaves and regeneration of shoots afterwards (Throught & Drew 1980; Jackson & Drew 1984). The higher overall flooding tolerance in R. palustris may result from a more efficient use of stored assimilates and/or a relatively high rate of alcoholic fermentation compared to R. maritimus (Laan & Blom 1990; Voesenek 1990).

The flooding tolerance of plants of both Rumex spp. and C. rubrum was highest in the September cohort (Table 1). The mean temperature of the flood-water was near the base temperature for growth of these species during the last flooding period (Van der Sman, Blom & Van de Steeg 1992) and this illustrates the important role of metabolic rates in the survival of submergence.

REPRODUCTION IN RELATION TO FLOODING

Plants of C. rubrum started to flower at an earlier stage and completed their life cycle in a shorter time in later cohorts (Fig. 2; see also Van der Sman, Blom & Van de Steeg 1992). The inability of plants of C. rubrum in the September cohort to produce seeds was probably due to low irradiance and/or insufficient photosynthesis (Kinet, Sachs & Bernier 1985). Flooding hardly reduced the probability of flowering in C. rubrum (Table 2). The onset of flowering, however, was not completely independent of size because large plants of C. rubrum flowered earlier than small ones (Fig. 2; see also Cumming 1969; Cumming & Seabrook 1985). A large size requirement in long days, gradually changing to a small one in short days probably results in a fine-tuned trade-off between growth and reproduction in C. rubrum.

In both Rumex spp. the minimum leaf number which must be achieved for flowering to occur remained constant in later cohorts (Fig. 2, see also Van der Sman, Blom & Van de Steeg 1992). Flooding delayed reaching this stage in both long-day species (Fig. 2) and as a result all flooded plants of R. palustris remained vegetative. Flooded plants of R. maritimus started to flower after they had initiated approximately the same number of primary leaves as drained plants, although their actual size (number and size of living leaves) was much smaller. The age-vs. size-dependency of the onset of flowering is the subject of many studies (e.g. Werner 1975; Lacey 1986; Klinkhamer, De Jong & Meelis 1987; Blom 1988). In most cases some measure of actual plant size is considered to represent developmental stage. In R. maritimus, stage and size are not equivalent in this respect. Flowering in this species is therefore probably not determined by the amount of stored reserves in the tap root, as was suggested for (short-lived) perennials with a size requirement for flowering (Werner 1975; Lacey 1986; Silvertown 1987).

Seed number was maximized at the expense of seed size in flooded plants of R. maritimus (Table 4). In addition, the seeds from flooded plants of R. maritimus had a lighter colour and were less dormant than seeds from drained plants. This suggests that they lack a hard seed coat, which may be the result of both immaturity and development under wet conditions (Silvertown 1984). This appears to be disadvantageous as it may result in the generation of small seedlings under adverse conditions for germination and establishment. Flooded plants of C. rubrum hardly managed to produce seeds, but in contrast to R. maritimus, those which did possessed relatively large seeds compared to drained plants (Table 4). Potential seed number is determined early in C. rubrum (Cook 1976) and depends, together with seed size, on photoperiod during flower induction (Cook 1975). If the number of flowers is reduced afterwards, overfilling of the seeds is likely to occur (Harper, Lovell & Moore 1970). In C. rubrum, the larger seeds from flooded plants as well as from later cohorts of drained plants were less dormant than small seeds (see also Van der Sman, Blom & Van de Steeg 1992; Cumming 1969). Decreased dormancy of the larger seeds which are produced on plants grown in short days is also found in C. album and in this species is caused by thinner seed coats (Karssen 1970). A large part of the seed-size variation in C. rubrum was not explained by treatment effects (Table 4) and must be ascribed to within-plant seed size variation or seed polymorphism (Harper, Lovell & Moore 1970).
Flooding Tolerance in Relation to Life History

In unpredictable environments, r-selection should be favoured (Stearns 1976; Grime 1979). Advantages of the annual strategy are a high intrinsic rate of increase through fast growth and early reproduction, and survival of adverse conditions by means of persistent seeds in the soil. Disadvantages are the loss of competitive ability, which is probably not very important in the open habitats in river forelands, and the yearly need of establishment microsites and survival of adverse conditions by means of favoured (Stearns 1976; Grime 1979). Advantages

stage (Khan & Ungar 1986). Both whereas large seeds germinate early and have a small seeds are incorporated into a persistent seed control (Table 4). Within-plant seed size variation other structures in later cohorts (Van der Sman, Blom & Van de Steeg 1992). The remarkable plasticity with respect to reproductive parameters in this species was typically expressed between plants (Bradshaw 1965). Seed polymorphism in C. rubrum, however, was only partly regulated by environmental control (Table 4). Within-plant seed size variation may act as a kind of bet-hedging against unpredictable conditions in the habitat (Harper, Lovell & Moore 1970). For example, in Atriplex triangularis, small seeds are incorporated into a persistent seed bank and thus ensure long-term population survival, whereas large seeds germinate early and have a better chance of surviving until the reproductive stage (Khan & Ungar 1986). Both Rumex spp. allocated more resources into vegetative growth in later cohorts and instead of plasticity in reproductive parameters they showed partitioning of resources to different fitness components (Harper 1977; Venable 1984). The strictly annual C. rubrum allocated even more resources into seeds than into other structures in later cohorts (Van der Sman, Blom & Van de Steeg 1992). The remarkable plasticity with respect to reproductive parameters in this species was typically expressed between plants (Bradshaw 1965). Seed polymorphism in C. rubrum, however, was only partly regulated by environmental control (Table 4). Within-plant seed size variation may act as a kind of bet-hedging against unpredictable conditions in the habitat (Harper, Lovell & Moore 1970). For example, in Atriplex triangularis, small seeds are incorporated into a persistent seed bank and thus ensure long-term population survival, whereas large seeds germinate early and have a better chance of surviving until the reproductive stage (Khan & Ungar 1986). Both Rumex spp. allocated more resources into vegetative growth in later cohorts and instead of plasticity in reproductive parameters they showed within-plant plasticity with respect to traits promoting survival, such as the ability to produce different leaf types under flooded and drained conditions. R. maritimus flowered more rapidly and was more susceptible to flooding than R. palustris. The three species appear to have features corresponding to the opposing strategies of stress tolerators and ruderals which may be found in the most frequently flooded parts of river floodplains (Menges & Waller 1983). R. palustris is best capable of maintaining occupation of sites which probably explains its occurrence in associations of marsh species. This species may be described as a stress-tolerant ruderal (sensu Grime 1979), while R. maritimus and C. rubrum possess increasingly ruderal characteristics.

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