Vegetative Propagation of Five Northern Forest Understory Plant Species from Either Rhizome or Stem Sections

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Abstract. Five understory plants, Asarum canadense L., Caulophyllum thalictroides (L.) Michx., Sanguinaria canadensis L., Trillium grandiflorum (Michx.) Salisb., of the Northeastern hardwood forests, and Oplopanax horridus (Sm.) Miq., of Western North American temperate forests, are of particular interest for horticultural and natural medicinal products industries. A rapid and efficient propagation method was needed to reproduce these plants vegetatively. To achieve this, the effect of 1000 mg L−1 (for herbaceous species) or 3000 mg L−1 (for O. horridus) auxin (IBA) and/or cytokinin (kinetin) on the growth of rhizome (for A. canadense, C. thalictroides, S. canadensis, T. grandiflorum) or stem sections (for O. horridus) was investigated. Nontreated controls were included for each species and an additional control with an intact apical bud was included for A. canadense, S. canadensis, and T. grandiflorum. No vegetative propagation was obtained for T. grandiflorum. Survival of O. horridus and C. thalictroides propagating units was 60% to 80% and 90% to 100%, respectively, and both rooted well even in absence of growth regulators. Asarum canadense produced two times and S. canadensis three to four times more roots when treated with the IBA + kinetin or the IBA treatment, respectively, compared with the control rhizome sections without an apical bud. For these two species, the presence of an apical bud enhanced survival and/or shoot emergence and those rhizome sections produced on average more biomass than the other treatments. Our results showed that either rhizome or stem sections can provide an efficient method to propagate A. canadense, C. thalictroides, O. horridus, and S. canadensis and thus reduce pressure on wild populations.

The market for medicinal plants has been growing steadily with an annual increase of 5% to 15% and an estimated value of $60 billion U.S. worldwide (Kartal, 2007). Understory shrubs and herbaceous species with medicinal values have traditionally been harvested from the wild. In addition to collection pressure on understory plants with medicinal properties, demand for indigenous woodland species that have an ornamental value is growing in the North American horticultural market. The lack of information on the most suitable cultivation methods for these ornamental and medicinal species, coupled with the fact that their growth is usually slow (Lamoureux and Nantel, 1999), has generally discouraged nurseries to grow them commercially.

Collecting plants from the wild reduces the ability of the exploited population to regenerate and also increases variability in the quality of the product. Four herbaceous understory species traditionally harvested from northeast North America, Asarum canadense (wild ginger), Caulophyllum thalictroides (blue cohosh), Sanguinaria canadensis (bloodroot), and Trillium grandiflorum (large-flowered trillium), and one woody shrub from northwest North America, Oplopanax horridus (devil’s club), were identified as species presenting an economical interest but no commercial method of propagation published to date. The commercial interest for these species, except T. grandiflorum, is recent. Although propagation from seeds is an interesting approach because it can lead to the production of many individuals from a single mother plant, its success is limited by the fact that most understory plant species require complex stratification treatments and that the growth of their seedlings is slow (Baskin and Baskin, 1998; Luna, 2001). On the other hand, vegetative propagation could lead to mature individuals after a single year of cultivation.

Several methods for the vegetative propagation of Trillium spp. have been described, all being based on the same principles. Briefly, they consist of reducing or eliminating apical dominance to induce the production of new buds along the rhizome or to stimulate existing dormant buds, both of which result in the formation of small offsets. After the flowering season, the terminal bud can be either girdled or removed and the rhizome replanted. The rhizome can also be cut transversally in two parts, leaving one section with the terminal bud and one without, both sections being replanted. In certain cases, buds along the rhizome are initiated, sprout, and even bloom the next year (Blanchette, 1998; Case and Case, 1997; Edgren, 1993). However, as reported by Case and Case (1997), researchers are still working on efficient propagating techniques that could be applied on a commercial basis for trillium species. The most detailed technique requires the application of a lanolin paste containing cytokinin on the cut surface of the rhizome, then once a callus is formed, a regular application of gibberellin on this callus until it differentiates into a shoot (Edgren, 1993). Such technique can be used by experienced gardeners but is difficult to upscale to commercial production.

The other herbaceous species selected for the present study naturally undergo vegetative propagation by rhizome growth (Lamoureux, 2002; Muir, 1995), whereas lateral branches of O. horridus form adventitious roots when in contact with soil (Lantz and Antos, 2002). Rhizome pieces or sections of stems are, therefore, likely to be promising avenues of investigation for the vegetative propagation of these species as recently shown for S. canadensis (Onofrietti, 2007). Vegetative propagation from stem tip cuttings has also been described for O. horridus (Luna, 2004).

Auxin, more specifically indole-3-butyric acid (IBA), is commonly used by commercial nurseries for induction of adventitious root formation on stem and leaf cuttings (McMahon et al., 2007; Srivastava, 2002), which is a prerequisite for successful propagation. The

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impact of auxin on rhizome sections, however, has not been tested to any great extent so far.

Cytokinins are another class of growth regulators that could be useful by stimulating shoot development of cuttings. A high cytokinin to auxin ratio has been repeatedly shown to promote bud formation and shoot development in plant tissue cultures (Srivastava, 2002). Application of kinetin or benzyladenine, two synthetic cytokinins, to buds induces axillary shoot development by releasing auxillary buds from the inhibition caused by apical dominance (Greene and Autio, 1989; Lyons and Hale, 1987). The effect of the application of synthetic cytokinins on underground tissues is less obvious and deserves more studies.

The present study was designed to evaluate the capacity of either rhizome or stem sections from A. canadense, C. thalictroides, O. horridus, S. canadensis, and T. grandiflorum to sprout with or without the application of root and/or shoot-inducing growth regulators. The results should help identify the treatments and propagating unit characteristics needed to enhance emergence and subsequent growth from rhizome or stem sections of these five North American understory species during the first year of their establishment.

### Materials and Methods

Manipulation of rhizome and stem sections and treatments applied. From the date of collection or purchase (Table 1), all materials were maintained at 4 °C under moist conditions until the onset of the experiment in Jan. 2003. This cold storage period was required to break the natural dormancy of the buds because materials were harvested in fall after plants became dormant (Case and Case, 1997; Gracie et al., 2000; Onofrietti, 2007). Rhizome or stem pieces were soaked (15 min) in a 1.2 g l⁻¹ solution of the systemic benzimidazole fungicide (C₁₇H₁₆N₂O₇; Benlate, DuPont, LA) before cutting them in different sections (Table 1) using surface-sterilized tools. The rhizomes and stems were treated with IBA (Sigma-Aldrich Canada Ltd., Oakville, Canada), kinetin (6-furfurylaminopurine; Sigma-Aldrich Canada Ltd.), a mix of both (IBA + kinetin) following the procedure described in Table 1, or left as untreated controls (C). Both IBA and kinetin were first dissolved in a limited volume of ethanol, then the solution was diluted with distilled water to reach final concentrations. A second control called Control + AB was tested for some of the species (Table 1), in which apical buds (AB) were present to examine the importance of apical dominance on the subsequent development of the plant from rhizome sections and to compare shoot size of already formed buds with shoot size of induced buds.

A second propagation technique, based on the methods described by Edgren (1993), Case and Case (1997), and Blanchette (1998), was tested for T. grandiflorum. Briefly, whole rhizomes (≈3 to 4 cm in length) bearing an apical bud were used. Two incisions 5 mm in length and 2 mm deep were made on the top surface of each rhizome to induce the production of offsets. The distal part of the rhizomes (i.e., the oldest part, opposite to the position of the apical bud) was subsequently cut to remove decaying material, and then the cut end of the rhizome was treated with growth regulators as previously described.

Once treated, the stem or rhizome sections were weighed and placed horizontally in 15-cm plastic pots containing: 3:1 Planting Mix (Fafard & Frères ltee, St.-Benoît (Quebec, Canada)), vermiculite, and perlite [3:1:1 (v/v)]. The pots were transferred to a greenhouse, maintained at 24 °C day/18 °C night (16-h photoperiod, natural sunlight supplemented by HPS 400-W lamps), and watered as needed, once every ≈2 d. Light in the greenhouse was reduced by the presence of a 50% shadecloth set up above the supplemental lighting. Mean photosynthetic photon flux density (PPFD) at plant level was 200 μmol m⁻² s⁻¹. After 3 months of growth, the plants were fertilized twice, at a 2-week interval, with a 1 g l⁻¹ solution of 20N–8.8P–16.6K (Plant-Prod-Plus; Plant-Prod Quebec, Laval, Canada). Planting occurred from 14 to 18 Jan. 2003 and harvesting time depended on species (Table 1).

**Experimental design.** The experimental unit comprised a single pot containing a single propagation unit. Each experimental unit was replicated 20 times for each treatment, giving a total of either 80 or 100 (for species with Control + AB) plants/species. Because light and temperature were variable across the greenhouse unit, species were put at the appropriate place based on what is known of their light and heat requirements (Cullina, 2000; Lamoureux, 2002; Small and Catling, 1999) and experimental units were randomized within each species. For example, A. canadense and C. thalictroides were put in shadier sections of the greenhouse, whereas O. horridus was placed in the sunniest area.

**Plant measurements.** Shoot production was measured at emergence and leaf area was measured either at the outset of leaf senescence or shortly before harvesting if senescence of the aboveground tissues had not occurred during the 4-month growth period. Allometric relations between leaf area, and leaf length and width were established for all herbaceous species using measurements obtained from natural populations (data not shown). These relations were used to estimate leaf area from leaf length and width measured directly on the plant before leaf senescence. The number of roots and buds produced during the 4-month growth period were determined at harvesting. Roots were harvested and divided into new (i.e., paler) and old (i.e., present at the time of planting) roots, dried at 70 °C for 48 h, and weighed. Visual observations indicated that the old roots did not elongate during the 4-month period; their biomass was thus a safe estimate of the initial root biomass at planting. We counted the number of new roots including both new roots that developed on the rhizomes and their lateral roots. Dry biomass values were also obtained for the rhizome or stem sections and for the shoots (70 °C for 48 h). Rhizome growth was estimated using two methods: the “loss or gain of rhizome biomass” [(dry rhizome biomass)ₕₖₕ – (dry rhizome biomass)ₖᵢₕ] and the “percentage of rhizome growth” [(|(dry rhizome biomass)ₕₖₕ – (dry rhizome biomass)ₖᵢₕ|)/(dry rhizome biomass)ₖᵢₕ]. The initial dry biomass of the rhizome or stem was calculated using a dry weight/fresh weight ratio that was established using 12 rhizome or stem sections of each species at the end of the experiment.

**Data analysis.** The percent emergence was calculated as the number of plants that produced at least one viable shoot out of the 20 plants included in each treatment. The percent survival was measured at harvest as the number of rhizome or stem sections that produced at least one new structure (shoot, root, or bud) during the experimental period. Percent survival and percent emergence were compared between treatments using a χ² test. Only plants considered to be alive at harvest were used for all other statistical analyses. The impact of the IBA and/or kinetin treatments on the different plant variables (i.e., production of new roots, shoots, and buds and rhizome growth) was investigated using analyses of covariance (ANCOVAs) with initial rhizome biomass (or initial stem biomass in the case of O. horridus), shoot biomass (used for one variable in O. horridus), and initial root biomass (when present) as covariates. When one of the covariates was significant (P < 0.05), results of ANCOVA were presented; otherwise, results of analysis of variance (ANOVA) were presented (P values) in the different tables. Pearson’s correlation tests were run to identify possible relations between the different initial plant variables (i.e., initial root biomass or initial rhizome biomass) and subsequent growth of the propagating units. When strong relations were found for a given species, t tests or ANOVAs were used to more fully characterize those relations. All statistical analyses were performed using Statistix 8.0 for Windows package (Analytical Software, Tallahassee, FL).

### Results

**Asarum canadense.** After a period of ≈6 weeks in the greenhouse, 61% of A. canadense rhizome sections sprouted (aboveground emergence; data not shown). At harvest, 56% of the planted rhizomes had developed new roots and shoots, whereas 5% developed only shoots, ≈1% developed only roots, and 7% developed only buds on the rhizome section (data not shown). Survival and all growth variables of A. canadense, except the number of new buds, were influenced to some extent by treatments (Table 2). Rhizome sections with apical bud (C + AB) had the greatest survival rate...
### Table 1. Characteristics of species and some aspects of treatments applied.

| Species | Growth habits | Collection and propagules used | Application of IBA and kinetin | Status of apical bud (AB) | Harvesting and senescence |
|---------|----------------|---------------------------------|---------------------------------|--------------------------|---------------------------|
| *Asarum canadense* | Produces long rhizomes that spread horizontally just a few centimeters below the soil surface; produces a limited number of roots; lateral buds along the rhizome first develop into new rhizomes before they can develop into terminal bud at the end of the season. | Collected in an *A. saccharum* dominated forest near Joliette (Quebec, Canada) in Oct. 2002; long segments of rhizomes, with a limited number of terminal buds, were harvested in natural populations; cut in 6-cm long sections at the time of planting. | The distal part (1 cm) of the rhizome sections was soaked in a solution of 1000 mg L⁻¹ during 15 s | AB was either removed or absent from rhizome sections except in Control + AB treatment | Plants had started to senesce when harvested from 20 to 30 May 2003 |
| *Oplopanax horridus* | Shrub species that produces low branches; branches root when they come in contact with the forest floor and the terminal end of the branch curves upward to produce a new ramet by layering. | Horizontal stem sections were harvested in early Fall 2002 and supplied by Pacific Rim Native Plants (Chilliwack, B.C., Canada); they were cut with an electrical saw to produce 7-cm long stem sections at the time of planting. | The distal part (1 cm) of the stem sections was soaked in a solution of 3000 mg L⁻¹ during 15 s | Most sections lacked a terminal bud; Control + AB not tested in this species. | Plants had not started to senesce when harvested from 22 to 29 May 2003; we decided to harvest after 4 months to mimic the length of a natural growing season. |
| *Sanguinaria canadensis* | Produces long rhizomes that spread horizontally just a few centimeters below the soil surface; produces a limited number of roots; lateral buds along the rhizome first develop into new rhizomes before they can develop into terminal bud at the end of the season. | Collected in an *A. saccharum* dominated forest near Joliette (Quebec, Canada) in Oct. 2002; long segments of rhizomes, with a limited number of terminal buds, were harvested in natural populations; cut in 2- to 3-cm long sections at the time of planting. | The distal part (1 cm) of the rhizome sections was soaked in a solution of 1000 mg L⁻¹ during 15 s | AB was either removed or absent from rhizome sections except in Control + AB treatment | Plants were harvested late—from 25 June to 7 July 2003—in the hope that shoots might emerge (see “Results”). |
| *Caulophyllum thalictroides* | Produces a compact rhizomatous system; crown of each individual plant contains many intertwined and branched rhizomes from which many shoots develop each year; plants usually extensively rooted. | Collected in an *A. saccharum* dominated forest near Joliette (Quebec, Canada) in Oct. 2002; a number of crowns were harvested, and then rhizome segments were detached from them and cut in 10-cm long sections at the time of planting. | Root tips (2 to 3 cm) were soaked in a solution of 1000 mg L⁻¹ during 15 s | There were numerous buds along the rhizomes and all were left in place on each section; Control + AB not tested in this species. | Plants harvested when completely senesced from 2 to 13 May 2003 |
| *Trillium grandiflorum* | Not a clonal species; produces a thick and short rhizome (≈3 to 6 cm in length for fully mature plants) with generally a single large bud at the growing end; there are many roots along the rhizome and new roots are produced each year near the bud. | Collected in an *A. saccharum* dominated forest near Joliette (Quebec, Canada) in Oct. 2002; rhizomes of mature plants were harvested and 1) cut in 2- to 3-cm long sections or 2) wound on the top surface of the rhizome (see text for more details). | 1) and 2) The distal part (1 cm) of the rhizome sections was soaked in a solution of 1000 mg L⁻¹ during 15 s | 1) AB was either removed or absent from rhizome sections except in Control + AB treatment; 2) all were whole rhizomes with AB. | Plants harvested when completely senesced; 1) plants harvested between 6 and 13 May 2003; 2) plants harvested between 28 Apr. and 17 May 2003 |

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1 Concentration used was based on Luna (2001) and McMahon et al. (2007).
2 As described for *Maianthemum dilatatum* (Wood) Nell & Macbr. (Lezberg et al., 2001) and *Podophyllum peltatum* L. (Geber et al., 1997), two other woodland herbs.
3 The distal end is the one opposite to the growing end (either a rhizome apex or a differentiated bud). It is the equivalent of the basal end of aboveground shoots.
4 Lantz and Antos, 2002.
5 Only sections with at least one small bud were used. Buds longer than 1 cm were removed to retain only small dormant buds on each section.
6 As described in Hanzawa and Kalisz (1993) and Lamoureux (2002), although Case and Case (1997) reported vegetative propagation of this species in some natural populations.

IBA = indole-3-butyric acid.
Table 2. Effect of indole-3-butryic acid (IBA), kinetin (Kin), or both (IBA + Kin) as well as the presence of apical bud (Control + AB) on the following variables: survival rate, total leaf area, number of new roots, new root, shoot, and change in rhizome biomass, rhizome growth, and bud number for *Asarum canadense* rhizome sections.

| State of rhizome sections at planting | Control | IBA | Kin | IBA + Kin | Control + AB | df | error | F | P |
|-------------------------------------|---------|-----|-----|-----------|--------------|----|-------|---|---|
| Absence of roots (N = 46)            | 75      | 65  | 65  | 65         | 65           |    |       |   |   |
| Presence of roots (N = 23)           | 100     | 65  | 65  | 65         | 75           |    |       |   |   |

Number of new roots

| Number of new roots                  | 5.1 c   | 8.8 bc | 6.0 c | 6.0 c | 15.0 a | 63  | 7.44 | <0.001 |

New root dry biomass

| New root dry biomass (g)             | 0.30 c  | 0.52 abc | 0.31 bc | 0.65 a | 0.56 ab | 61  | 3.05 | 0.023 |

Shoot dry biomass

| Shoot dry biomass (g)                | 0.19 c  | 0.33 c | 0.27 bc | 0.45 ab | 0.58 a | 61  | 5.62 | <0.001 |

Change in rhizome biomass

| Change in rhizome biomass (g)        | -0.03 c | 0.15 bc | -0.06 c | 0.32 a | 0.23 ab | 62  | 7.34 | <0.001 |

Rhizome growth

| Rhizome growth (%)                  | -8.0 c  | 28.8 bc | -12.6 c | 79.6 a | 54.5 ab | 62  | 7.17 | <0.001 |

Number of new buds

| Number of new buds                  | 3.5     | 3.1    | 2.8    | 3.9    | 3.2     | 63  | 0.51 | 0.73  |

Table 3. Effect of presence of roots at planting on subsequent growth of rhizome and on production of new roots, shoots, and buds (mean ± se) and *P* values for the *t* tests for *Asarum canadense* rhizome sections.

| State of rhizome sections at planting | Control | IBA | Kin | IBA + Kin | Control + AB | df | error | F | P |
|-------------------------------------|---------|-----|-----|-----------|--------------|----|-------|---|---|
| Absence of roots (N = 46)            | 7.9 ± 1.1 | 13.1 ± 1.4 | 0.005 |
| Presence of roots (N = 23)           | 11.6 ± 9.7 | 66.7 ± 13.0 | 0.001 |

Number of new roots

| Number of new roots                  | 5.1 b    | 5.3 b | 3.9 b | 8.7 a | 55 | 5.07 | 0.004 |

New root dry biomass

| New root dry biomass (g)             | 0.37 ± 0.05 | 0.68 ± 0.08 | 0.001 |

Shoot dry biomass

| Shoot dry biomass (g)                | 0.30 ± 0.04 | 0.58 ± 0.08 | 0.002 |

Rhizome growth

| Rhizome growth (%)                  | 11.6 ± 9.7 | 66.7 ± 13.0 | 0.001 |

Number of new buds

| Number of new buds                  | 4.5 ± 0.3 | <0.001 |

Table 4. Effect of indole-3-butryic acid (IBA), kinetin (Kin), or both (IBA + Kin) on survival rate, number of new roots, new root, shoot, and change in stem biomass, stem growth, and bud number for *Oplopanax horridus* stem sections.

| State of stem sections at planting | Control | IBA | Kin | IBA + Kin | Control + AB | df | error | F | P |
|-----------------------------------|---------|-----|-----|-----------|--------------|----|-------|---|---|
| Absence of buds (N = 46)          | 80      | 60  | 80  | 75        | 75           |    |       |   |   |
| Presence of buds (N = 23)         | 80      | 60  | 80  | 75        | 75           |    |       |   |   |

Number of new buds

| Number of new buds                 | 8.6 a   | 7.8 | 4.7 b | 7.5 a | 5.4 | 3.01 | 0.038 |

Table 5. Effect of presence of roots at planting on subsequent growth of rhizome and on production of new roots, shoots, and buds (mean ± se) and *P* values for the *t* tests for *Asarum canadense* rhizome sections.

| State of rhizome sections at planting | Control | IBA | Kin | IBA + Kin | Control + AB | df | error | F | P |
|-------------------------------------|---------|-----|-----|-----------|--------------|----|-------|---|---|
| Absence of roots (N = 46)            | 75      | 65  | 65  | 65         | 65           |    |       |   |   |
| Presence of roots (N = 23)           | 100     | 65  | 65  | 65         | 75           |    |       |   |   |

Number of new roots

| Number of new roots                  | 5.1 c   | 8.8 bc | 6.0 c | 6.0 c | 15.0 a | 63  | 7.44 | <0.001 |

New root dry biomass

| New root dry biomass (g)             | 0.30 c  | 0.52 abc | 0.31 bc | 0.65 a | 0.56 ab | 61  | 3.05 | 0.023 |

Shoot dry biomass

| Shoot dry biomass (g)                | 0.19 c  | 0.33 c | 0.27 bc | 0.45 ab | 0.58 a | 61  | 5.62 | <0.001 |

Change in rhizome biomass

| Change in rhizome biomass (g)        | -0.03 c | 0.15 bc | -0.06 c | 0.32 a | 0.23 ab | 62  | 7.34 | <0.001 |

Rhizome growth

| Rhizome growth (%)                  | -8.0 c  | 28.8 bc | -12.6 c | 79.6 a | 54.5 ab | 62  | 7.17 | <0.001 |

Number of new buds

| Number of new buds                  | 3.5     | 3.1    | 2.8    | 3.9    | 3.2     | 63  | 0.51 | 0.73  |

The best growth responses were obtained for rhizome section with apical bud and for those treated with IBA, in particular when combined with kinetin. Control + AB and IBA + kinetin produced more than or nearly twice the number of roots, biomass of roots, and shoots and total leaf area compared with the control. Rhizome biomass increased slightly with application of IBA alone but decreased with kinetin application and in absence of growth regulators (Control). The highest increase was observed with IBA + kinetin or in presence of an apical bud (Control + AB). Rhizome sections from all groups produced a mean of approximately three new buds. Further analyses of the data indicated that rhizome sections with roots at the outset of the experiment had better subsequent growth than plants lacking roots at planting (Table 3). They produced more shoot and root biomass, more buds, and more rhizome mass than rhizomes devoid of roots at planting. Dry biomass of the new roots was strongly correlated with total leaf area (*r* = 0.90, *P* < 0.001; data not shown).

*Oplopanax horridus*. Seventy-four percent of *O. horridus* stem sections developed at least one shoot and from those, 86% rooted as well (data not shown). For most stem sections, sprouting started 2 weeks after the outset of the experiment. The original stem section remained unchanged but new shoots were produced from dormant buds located along this original stem section. At the end of the 4-month experiment, stem sections had developed at least one vertical stem with four to five leaves. Plants treated with the IBA + kinetin produced more roots than the other treatments (Table 4), but because this number was highly variable between plants, root biomass was probably a better estimation of the capacity of these plants to absorb water and nutrients. Kinetin application significantly reduced root biomass along with shoot biomass. Buds were produced at two locations on the vertical stems: in a crown around the base of the stem and at the point of attachment of each petiole. As a result of the close correlation between the number of buds and the size of the aerial stems, shoot dry biomass was tested as a covariable in the ANCOVA for the number of new buds. When this covariable was included in the statistical model, treatments no longer differed in terms of bud number (*P* = 0.93; data not shown). However, when the initial dry biomass of the stem section was included as a covariable, bud production was reduced by kinetin application, most probably through its effect on shoot biomass (Table 4). The initial dry biomass of *O. horridus* stem section positively influenced the dry biomass of new roots and shoots and the production of buds (Table 4; covariable significant). Sprouting capacity was also influenced by the initial biomass of the stem section; stem sections that did not sprout had an initial fresh weight (FW) of 16.7 ± 2.0 g, whereas those that produced at least one shoot had an initial FW of 26.6 ± 2.4 g. The initial stem section did not grow, thus explaining the lower final biomass measured for this variable regardless of the treatment.
roots. Rhizomes with an apical bud and those treated with IBA produced the highest root biomass, whereas the number of new roots was at least three times higher for IBA, IBA + kinetin, and Control + AB than for kinetin alone or the control (Table 5). Because of the low percent emergence, evaluating the effect of IBA and kinetin on shoot biomass was impossible. Rhizome sections with an apical bud increased their total rhizome mass by 166%, whereas all other groups exhibited a mean decrease of 31%. The number of new rhizome sections rotted was 1.5 on average, independently of the treatment (Table 5), but almost half of the rhizome sections treated with IBA failed to produce any buds (data not shown). Furthermore, the number of buds produced was influenced by the initial rhizome and root biomass because the inclusion of these variables in the statistical model failed to reveal differences between treatments (P = 0.12). Rhizome sections with an apical bud had more roots at planting than the other treatment groups (P < 0.001; data not shown).

Caulophyllum thalictroides. Ninety-six percent of the C. thalictroides rhizome sections produced new structures and 81% sprouted (data not shown). All rhizome sections that sprouted developed new roots, only 9% of rhizome sections rooted without producing shoots, and 6% only formed new buds on the rhizome sections (data not shown).

Most of the shoots emerged 1 week after planting. Neither sprouting nor other measures variables responded to the treatments (Table 6). At the end of the experiment, the rhizomes had lost an average of 2 g or 20% of their initial biomass and produced an average of 46 new roots (for a dry biomass of 1 g), two new stems (with a total leaf area of 582 cm²), and 13 new buds/rhizome.

The initial biomass of roots at planting was correlated (P < 0.001, data not shown) with new root dry biomass production (r = 0.52), shoot dry biomass (r = 0.79), and total leaf area (r = 0.68). Initial rhizome section biomass was negatively correlated with the percent increase in rhizome biomass (r = -0.43, P < 0.001). Furthermore, bigger rhizomes produced smaller shoot biomass (r = -0.60, P < 0.001) and total leaf area (r = -0.64, P < 0.001) when both were expressed on a per gram of initial rhizome section biomass.

Trillium grandiflorum. Only 21% of the T. grandiflorum rhizome sections without an apical bud produced new roots or buds on the rhizome section, and none sprouted (data not shown). Although only 3% of the rhizome sections rotted, the rhizome sections that did not produce any new structures (79%) were considered dead. On the other hand, all rhizome sections with an apical bud produced a shoot and some new roots except for one rhizome section that rotted. The limited data for rhizome sections with new structures thus prevented comparisons between treatments. Cutting rhizome sections of T. grandiflorum into two pieces failed as a propagation method, even when combined with IBA, kinetin, or both treatments, because in most cases, only the section with the apical bud survived.

Of those T. grandiflorum rhizomes that had incisions in the top half (wounded), 96% produced at least one new structure and 91% sprouted (Table 7). Neither emergence nor growth variables responded to treatments. On average, rhizomes produced seven new roots for a mean dry biomass of 0.09 g, developed 0.35 g of new shoot tissue, and their rhizome biomass increased by 0.38 g or 25%. Only two rhizome sections (one control and one IBA-treated) produced new buds at the site of the incisions. For all the other rhizomes, the wounded areas had healed over and the only bud that develops on these rhizomes was in the apical position. Overall, a high initial
Rhizome biomass, coupled with a high root biomass at planting, enhanced subsequent shoot production (data not shown).

Discussion

Percent survival of rhizome and stem sections. The use of rhizome or stem sections seems to be a satisfactory technique for the propagation of *O. horridus*, *S. canadensis*, and *C. thalictroides*. Shoot emergence of *O. horridus* was 74% and that of *C. thalictroides* was 81%. New roots were produced on 64% of *O. horridus* shoot sections and on 85% of *S. canadensis* and 90% of *C. thalictroides* rhizome sections. By contrast, only 61% of *A. canadense* rhizome sections produced a shoot and 57% produced new roots. Survival was not linked to initial rhizome biomass in this species (data not shown). However, the presence of roots at planting was positively correlated with new root biomass, shoot biomass, and percent rhizome growth. The initial presence of roots may have influenced rhizome section survival. Unfortunately, verifying this point was impossible because dead rhizomes decomposed before the end of the experiment. The irrigation regime may also have applied too much water for this species, which caused rotting of the rhizomes before sprouting, as noticed by G. Dostie (U. Sherbrooke, personal communication). *Asarum canadense* rhizome is apparently prone to rotting when poorly rooted.

The high survival rates obtained for stem sections of *O. horridus* were probably the result of the natural ability of this species to propagate by layering and by sprouting from basal stems (Lantz and Antos, 2002). We found that stem sections that weighed more than 16.7 g FW were more likely to survive than smaller ones, supporting results obtained from previous studies on woody and semiwoody species (Chalapathi et al., 2001; Palamisamy and Kumar, 1997). Total nonstructural carbohydrates have been shown to influence rooting by providing energy reserves and a carbon skeleton to support root initiation and subsequent growth (Haisig, 1986; Veierskov, 1988).

In the present study, all attempts to propagate *T. grandiflorum* failed. *T. grandiflorum* rhizome sections, without the application of growth regulators, remain the traditional propagation methods for this species (Case and Case, 1997), but the percentage of success using this technique is low. In contrast to the results obtained by Edgren (1993) and Blanchette (1998), superficial wounding applied on whole rhizomes, whether disbudded or not, failed to induce bud formation in the present study. The incisions may have been too shallow [Blanchette (1998) used 3- to 5-mm deep incisions] and perhaps healed over before inducing bud formation. Another possible reason underlying the poor results with this species may be related to intraspecific genetic variations within colonies or populations, which could result in a lower capacity of certain individuals to regenerate vegetatively. This possibility was advanced by Case and Case (1997), who observed high vegetative reproduction rates for certain of their own *T. grandiflorum* clones, whereas the cloning ability of this species was considered by some authors to be nonexistent (Hanzawa and Kalisz, 1993) or rare (Lamoureux, 2002). Therefore, studying the intraspecific genetic variation of *T. grandiflorum* with regard to cloning ability could be of considerable practical value for future commercial propagation of this species.

Treatment effects. For *A. canadense* and *S. canadensis*, treating rhizome sections with IBA or IBA + kinetin enhanced root production. The general effect of auxin on adventitious root production is well known (Loach, 1988; McMahon et al., 2007; Srivastava, 2002). For *A. canadense* rhizome sections, root biomass appears to influence both rhizome growth and shoot production. This indicates that better-rooted rhizome sections were able to take advantage of an increase in shoot biomass and leaf area, which likely increased the amount of carbon fixed and resulted in larger resources for rhizome growth compared with lightly rooted rhizome sections. Therefore, providing favorable conditions to promote good rooting appears important for the production of healthy *A. canadense* plants from rhizome sections, which in turn should enhance survival of these plants over time and may be of particular importance for species that are normally characterized by a limited number of superficial roots such as *A. canadense* and *S. canadensis*.

For *O. horridus*, IBA failed to affect root production on stem sections. This species undergoes natural layering (Lantz and Antos, 2002) and thus has a good inherent rooting capacity. In the present study, stem sections produced a large root biomass irrespective of treatment. Auxin (IBA) application also failed to improve the rooting capacity of *C. thalictroides*. The application of IBA to the apical end of roots in this species, contrary to an application to the distal part of the rhizome, may have impeded the expected root-promoting effect of the IBA treatment. By contrast, the initial biomass of roots on rhizome sections at planting was important and positively influenced both production of new roots and shoots. Although the present study was unable to show a positive effect of auxin on root formation, the effect of IBA on poorly rooted stem sections of *C. thalictroides* should be tested.

Interestingly, the species that produced the fewest roots (e.g., *A. canadense* and *S. canadensis*) were also those that were the most sensitive to IBA application. This situation is frequently observed and could be explained by the fact that the ability of a given species or cultivar to root seems to be related to its endogenous indole acetic acid (IAA) content (Srivastava, 2002). An exogenous auxin application would consequently correct for the lack of IAA in *A. canadense* and *S. canadensis*, but would have little effect on species such as *O. horridus*, which might already have a high level of endogenous IAA.

Kinetin had a negative effect on root, shoot, and bud production of *O. horridus*. Cytokinins, when in too high a concentration, are known to have an adverse effect on root formation (Srivastava, 2002), which may explain why fewer roots formed on kinetin-treated stem sections than on stem sections receiving the other treatments. This result in turn probably explains why kinetin-treated stem sections produced less shoot biomass than stem sections receiving the other treatments.

Kinetin treatment failed to stimulate formation of adventitious buds (reviewed by Cline, 1991) in all five species. In the case of *A. canadense*, each Control + AB rhizome section produced, on average, 3.2 new lateral buds, indicating that apical dominance is weak in this species and unlikely to influence its vegetative propagation. Similarly, Control + AB rhizome sections of *S. canadensis* produced 2.7 buds on average, indicating that at least two shoots per plant would sprout the next year. There are conflicting results in the literature about the effect of cytokinin on lateral bud development. The cytokinin benzyl adenine, when applied basally, favored lateral bud development in *Arabidopsis* cuttings, but when applied apically, they remained dormant (Chatfield et al., 2000). Srivastava (2002) also reported that the application of cytokinins, alone or with auxin, to the apical end of a decapitated stem fails to release lateral buds from inhibition and may even reinforce inhibition. Application of benzylaminopurine, a synthetic cytokinin, to the basal part of rootless shoot cuttings of *Solanum tuberosum* spp. *andigena* Hawkes induced lateral shoot development (Woolley and Wareing, 1972). On the other hand, cytokinins stimulated bud development when applied to the lateral buds of plants with an intact apex (Srivastava, 2002) or to decapitated plants to which inhibitory concentrations of apical auxin were applied (Mok, 1994). Thus, application of exogenous cytokinins to either lateral buds directly or to the basal part of the stem like in the present study has been shown to release lateral buds from inhibition, but timing of application or concentrations might explain the lack of effect on the species under study.

Rhizome growth. In the present study, experiments were stopped after 4 months and measurements were taken at shoot senescence for each species except *O. horridus*, which was harvested before senescence. Nevertheless, rhizome biomass decreased for three of the species. Such a loss was anticipated for rhizome sections of *S. canadensis*, which used existing reserves to produce new buds and roots without producing shoots. This result was supported by the fact that control plants of *S. canadensis* possessing an intact apical bud (Control + AB) or which nearly all sprouted, exhibited a significant increase in rhizome biomass during the growth season. With regard to *O. horridus*, examination of the stem sections revealed an old dehydrated stem section, suggesting that the new shoots became partially or totally
independent of the initial horizontal stem section. In the case of C. thalictroides, a reduction in rhizome biomass was also observed irrespective of treatment. The reason behind this reduction is unclear. The rhizomes were alive and lacked decay, and the shoots, which had undergone a 4-month growth period, had all senesced naturally. Carbon reserves should have been translocated to the rhizome before harvesting. The fact that bigger rhizomes produced proportionally less shoot biomass and leaf area than smaller rhizomes might explain the negative correlation between rhizome size and growth observed for this species. Larger rhizomes would produce insufficient leaf area to allow rhizome growth either through production of new tissues or accumulation or reserves. However, the factor that limits shoot production by larger rhizomes must still be explained.

In conclusion, the present study has shown that untreated rhizome sections of C. thalictroides and stem sections of O. horridus, and IBA-treated rhizome sections of S. canadensis and A. canadensis, are efficient methods for vegetative propagation of these species. Rhizomes of C. thalictroides responded well to cutting provided that the rhizome sections weighed less than 20 g FW and were well furnished with roots at the time of planting. Stem sections of O. horridus (initial fresh weight greater than 16.7 g) rapidly established new roots and shoots. Rhizome sections of S. canadensis survived at high rate and IBA enhanced formation of new roots. The new buds produced during the first growing season would be expected to emerge the next year (Onofrietti, 2007). However, Pythium can attack S. canadensis shoots if the substrate is too moist (Greenfield et al., 2006). Rhizome sections of A. canadense generally responded well when used as propagating units but are susceptible to rotting when poorly rooted. Auxin application is recommended for vegetative propagation by rhizome sections for this species. The cytokinin kinetin was ineffective in stimulating either adventitious or axillary buds in any of the five species in contradiction to its effectiveness for this purpose in tissue culture propagation systems. None of the vegetative propagation treatments resulted in established plants of T. grandiflorum. Deeper incision of the rhizome, clonal differences in their ability to propagate vegetatively (Case and Case, 1997), in vitro propagation techniques (Pence and Soukop, 1993), and production from seeds (Solt, 2002) are among possible avenues of research for this species. The techniques used in the present study can be easily used in commercial nurseries as well as to set up plantations in the understory of hardwood forests. Other plant growth regulators and/or concentrations need to be tested in the future to further improve rooting and bud production. Harvesting mother plant material at other times during the year could also influence survival and sprouting in some of these species.

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