Exchange of Radioactive Phosphorus ³²P between the Components of an Artificial Plant Community*

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Abstract

In artificial plant communities (wooden cases with soil) the following plants were grown together for one year: case No. 1: Betula verrucosa Ehrh. (2 specimens) and Carex pilosa Scop.; sases No. 2 and 3: Alnus incana (L.) Mnch., Fraxinus excelsior L., Padus avium Mill., Aegopodium podagraria L., Eupatorium cannabinum L. and others.

Into the stalks or stems of one to three plants of every case ³²P was introduced. After 3 months all the plants in every case contained radiactive phosphorus. There were great differences in ³²P concentration (up to ten-fold) between plants grown in the same case. The concentration of ³²P in plants into which it had not been introduced was of the range of 0.1%.

Preliminary investigations by the authors (Domnicz, Zarzycki 1970) showed that tomatoes in short-term water and soil cultures discharged into the substrate about $0.5^{\circ}/_{\circ}$ of the ^{32}P introduced into these plants. A significant part of the phosphorus discharged (up to $70^{\circ}/_{\circ}$) was then absorbed from the substrate by the roots of other specimens of the same species. Fober and Giertych (1970) observed weak emission of ^{32}P in Poa annua L., which was markedly stronger in seedlings of Picea abies (L.) Karsten.

The purpose of the next experiment was to follow the course of radioactive phosphorus exchange between components of a simplified plant community in nearly natural conditions.

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MATERIAL AND INVESTIGATION METHODS

In spring 1969 six $55 \times 40 \times 35$ cm wooden cases were filled with sandy-loamy, slightly humic garden soil. In three cases young specimens of Betula verrucosa 60-120 cm tall, and several specimens of Carex pilosa were planted; in this way a simplified model of forest community was created. In the remaining three cases young trees and herbaceous plants were planted, belonging to the composition of Alnetum incanae (Fig. 1). For 1 year the cases were left in the garden, the plants were watered during the dry summer period and protected against frost in winter. After one year, when the plants had well developed roots, 3 cases with abundantly and uniformly developed vegetation were selected for the proper investigations.

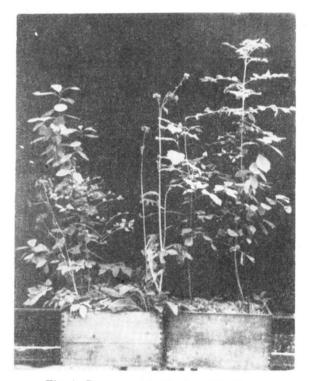


Fig. 1. Cases used in the investigations

The vegetation of case No. 1 consisted of three specimens of birch and numerous specimens of Carex pilosa. In case No. 2 there grew (one specimen each) Alnus incana, Padus avium and Fraxinus excelsior, as well as aboundant herbaceous plants: Eupatorium cannabinum, Asarum europaeum L., Carex silvatica Huds., Chelidonium majus L., Brachypodium silvaticum (Huds.) Roem. et Schult. and Aegopodium podagraria. The vegetation of case No. 3 consisted of young trees of Alnus incana, Padus avium and

Fraxinus excelsior, as well as the herbaceous plants Eupatorium cannabinum, Carex silvatica, Brachypodium silvaticum and Aegopodium podagraria.

An injection of ³²P in the form of NaH₂PO₄ was given on May 18th, 1970, during exuberant growth of the plants. In case No. 1 the radioactive substance was introduced into the stems of two birch trees; the holes of c. 2 mm diameter were directed at a slant downwards to a depth of c. 10 mm, and c. 1 mci of ³²P was introduced into each (Fig. 2). The places of injection were protected against ambient contamination by wood-wool and sealing tape. In case No. 2 about 2 mci of ³²P were introduced into the stalk of *Aegopodium podagraria*, and in case No. 3 about 1 mci of the same into 3 shoots of the same plant.

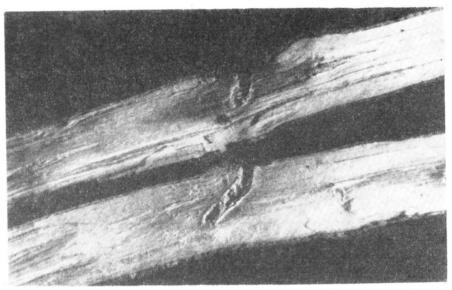


Fig. 2. Cross-section of young birch stem with a hole for introducing 32P

The rate of distribution of radioactive phosphorus in plants into which it had been introduced was checked in the leaves of birch trees in the upper, central and lower part of the crown. In the case of *Aegopodium podagraria* circles of 5-mm diameter were cut from young, older and intermediate leaves. Collection of the material was begun as early as 1/2 hour after the injection.

Three months after introduction of ³²P (i.e. on August 18th, 1970) the plants from the individual cases were harvested, divided into leaves and stalks, or, in the case of trees, into leaves, branches and stems, and then dried, weighed and burned. In the ash total phosphorus content was colorimetrically determined (Domnicz, Gorlach 1970), and total radioactivity in the manner described in the preceding paper (Domnicz, Zarzycki 1970).

Considerable methodical difficulties were encountered in the separation of small roots and root hairs from the soil. Therefore, determination of the ³²P content in underground part of plant and soil was omitted.

INVESTIGATION RESULTS AND DISCUSSION

The radioactive phosphorus from the stems or stalks of herbaceous plants was very quickly distributed within all the plant. Ten hours after injection the ³²P content in birch leaves reached its maximal level (Fig. 3).

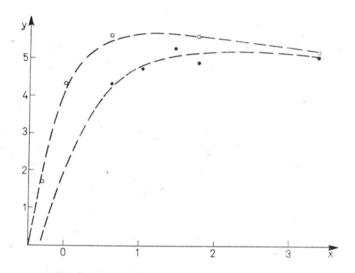


Fig. 3. The rate of distribution of ³²P to leaves from upper and lower part of crown of *Betula verrucosa* into which stem it had been introduced Y-axis — log activity in imp./min./100 mg dry matter of leaves; X-axis — log time in hours after injection of ³²P

After 3 months from the injection ³²P radioactive phosphorus was found in all plants growing together in one case. If in plants which had been injected with ³²P (plants "R") activity reached, or even surpassed, 10⁶ imp./min./100 g dry matter, then in adjacent recipient plants this activity did not exceed as a rule 10³ imp./min./100 g dry matter. If the ³²P content in leaves of "R" plants was assumed at 100⁹/₉, the content of this element in leaves of "O" plants was between 0.1 and 1.4⁹/₉. The high activity found in one of the specimens of Aegopodium podagraria from case No. 2 (plant No. 6) was probably caused by an underground connection with the "R' plant. The amounts of ³²P in other species do not show such wide differences when compared with one another (Figs 4, 5 and 6).

The authors' invesigations show that the components of one plant community exchange between each other, although to an insignificant

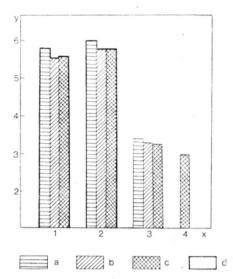


Fig. 4. Radioactivity of plants (case No. 1) months after the injection of ³²P into two stems of young briches

Y-axis — log activity in imp./min. per mg P; a — stems, b — branches, c — leaves, d — plants which had been injected with $^{32}{\rm P}$

X-axis — plant species: 1, 2 — Betula verrucosa (plants injected with 32P), 3 — Betula verrucosa (without injection), 4 — Carex pilosa

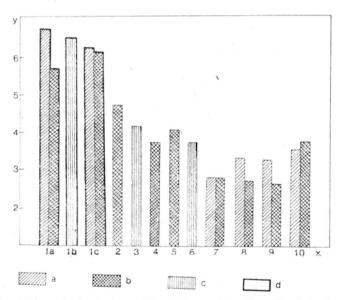


Fig. 5. Radioactivity of plants (case No. 3) 3 months after the injection of ³²P into 3 stalks of Aegopodium podagraria

Y-axis — log activity in imp./min./mg P; X-axis — plant species:

1a, 1b, 1c — Aegopodium podagraria (plants injected with ³²P), 2, 3 — Aegopodium podagraria (without injection), 4 — Brachypodium silvaticum, 5 — Carex silvatica, 6 — Eupatorium cannabinum, 7, 8 — Fraxinus excelsior, 9 — Alnus incana, 10 — Padus avium;

a — stalks, branches, stems, b — leaves, c — all plant, d — plants injected with 32 P

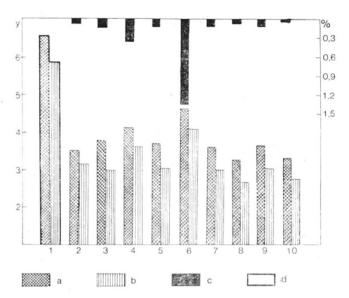


Fig. 6. Radioactivity of plants (case No. 2) 3 months after the injection of ³²P into one stalk of *Aegopodium podagraria*

Y-axis — log activity in imp./min. per mg P of leaves (a) or per 100 mg dry matter (b), c — percentage of radioactivity in non-injected plants (100% = radioactivity of plant injected with ^{32}P — d); X-axis — plant species:

1 — Aegopodium podagraria (plant injected with ³²P), 2 — Padus avium, 3 —
 Fraxinus excelsior, 4 — Eupatorium cannabinum, 5, 6 — Aegopodium podagraria,
 7 — Asarum europaeum, 8 — Chelidonium majus, 9 — Carex silvatica, 10 — Brachypodium silvaticum

degree only, phosphorus compounds *in vivo*. This exchange, in conditions of the experiment performed, is not of significant importance as regards plant nutrition but it indicates the existence of a feeding connection between such plants. Part of the phosphorus diffused in inorganic form is probably intercepted also by microorganisms.

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Wymiana radioaktywnego fosforu ³²P pomiędzy komponentami sztucznej asocjacji roślinnej

Streszczenie

Wstępne badania autorów wykazały (Domnicz, Zarzycki 1970), że pomidory w krótkotrwałych kulturach wodnych i glebowych wydzielały do podłoża około 0,5% wprowadzonego do nich radioaktywnego fosforu. Znaczna część wydzielonego 32P (do 70%) została następnie pobrana z podłoża przez inne osobniki tego samego gatunku. W następnym doświadczeniu prześledzono jak przebiega wymiana radioaktywnego fosforu pomiędzy komponentami uproszczonego zbiorowiska roślinnego w warunkach zbliżonych do naturalnych.

Do 6 skrzyń napełnionych glebą ogrodową wysadzono wiosną 1969 r. młode okazy drzew i roślin zielnych, które rosły w nich przez okres 1 roku (ryc. 1). Gdy rośliny dobrze się ukorzeniły wytypowano do właściwych badań 3 skrzynie. W skrzyni nr 1 rosły 2 młode brzozy brodawkowate i liczne okazy Carex pilosa (odpowiednik spotykanego czasami w przyrodzie lasu brzozowego, który powstał po wycięciu drzewostanu grądowego). W skrzyniach nr 2 i 3 uformowano model olszynki karpackiej (Alnetum incanae); rosły w nich m.in. Alnus incana, Fraxinus excelsior, Padus avium do 1,2 m wysokie oraz z roślin zielnych Aegopodium podagraria, Brachypodium silvaticum, Carex silvatica i in.

Do pni brzóz lub pędów podagrycznika w obrębie każdej skrzyni w dniu 18 maja 1970 r. wprowadzono ⁵²P (ryc. 2), który szybko został rozprowadzony po roślinie (ryc. 3). Po 3 miesiącach (18 sierpnia 1970) od wprowadzenia ⁵²P we wszystkich roślinach rosnących razem w jednej skrzyni znaleziono radioaktywny fosfor. Gdy w roślinach. do których wprowadzono ⁵²P drogą iniekcji (rośliny "R"), aktywność dochodziła lub nawet przekraczała 10⁶ imp./min./100 mg suchej masy, to w sąsiadujących z nimi roślinach (rośliny "O") nie przekraczała z reguły aktywności 10³ imp./min./100 mg suchej masy (ryc. 4, 5, 6).

Niniejsze badania wykazują, że komponenci jednego zbiorowiska roślinnego wymieniają między sobą, w niewielkim zresztą stopniu, związki fosforowe "na żywo. Wymiana ta, w warunkach przeprowadzonego doświadczenia, nie ma większego znaczenia dla odżywiania roślin, ale wskazuje na istnienie między nimi więzi pokarmowej. Część fosforu, który wydzielany jest w formie nieorganicznej, przechwytywana jest najprawdopodobniej także przez mikroorganizmy.

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