

RESEARCH NOTE

Release of phosphorus, aluminium and iron in fractionation of inorganic soil phosphorus

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Abstract. Release of phosphorus, aluminium and iron by a modified CHANG and JACKSON procedure was studied in five mineral soils. Quantities of aluminium and iron released during the procedure and extracted by acid ammonium oxalate were compared. The extractability of P, Al and Fe by 1 M NH_4Cl and that of Al and Fe by alkaline 0.5 M NH_4F was poor. Proportions of P extracted by 0.5 M NH_4F (0.2—10.4 mmol/kg soil) and 0.1 M NaOH (0.1—9.8 mmol/kg soil) were related to the molar ratio of oxalate-extractable iron and aluminium. P extracted by 0.25 M H_2SO_4 amounted to 2.1—12.2 mmol/kg soil. Al extracted by 0.1 M NaOH (7—174 mmol/kg soil) and 0.25 M H_2SO_4 (17—112 mmol/kg soil) amounted to 55—94 % and 16—245 % of oxalate-extractable Al, respectively. Fe released by 0.1 M NaOH (1—10 mmol/kg soil) and 0.25 M H_2SO_4 (30—196 mmol/kg soil) amounted to 1—13 % and 62—272 % of oxalate-extractable Fe, respectively. In total, 91—309 % of oxalate-extractable Al and 70—285 % of oxalate-extractable Fe were released by NaOH and H_2SO_4 .

Index words: phosphorus fractions, extractable aluminium and iron

Introduction

The fractionation procedure developed by CHANG and JACKSON (1957) is frequently used in estimation of inorganic soil phosphorus (e.g. KAILA 1964, HARTIKAINEN 1979). Phosphorus fractions bound by aluminium and iron oxides and phosphorus of calcium phosphates such as apatite are considered to be extracted successively in the procedure. The reagents used are known partly to extract soil aluminium and iron, but the amounts released

besides phosphorus are infrequently determined. The aim of this study was to examine the simultaneous release of phosphorus, aluminium and iron in fractionation and to compare the extractability of aluminium and iron with their extractability by acid ammonium oxalate.

Material and methods

The material consisted of five mineral soil samples from the Viikki Experimental Farm,

Table 1. Characteristics of experimental soils.

	Soil sample				
	1	2	3	4	5
Sampling depth, cm	0—20	20—40	20—40	0—20	20—40
pH(CaCl ₂)	5.1	4.6	4.8	5.3	5.0
Org. C, %	3.6	0.8	2.6	4.4	1.0
Particle-size distribution, %					
<0.002 mm	13	2	47	10	26
0.002—0.02 mm	20	1	30	7	2
0.02—0.06 mm	27	7	18	15	23
0.06—0.20 mm	31	35	5	61	42
0.20—2.00 mm	9	56	0	7	6
Oxalate-soluble Al mmol/kg	186	104	76	23	11
Oxalate-soluble Fe mmol/kg	53	32	224	140	11
Oxalate-soluble Fe/Al	0.3	0.3	2.9	6.1	1.0

University of Helsinki, (Nos. 2—4) and South Karelia (Imatra) (No. 1): Nos. 1 and 4 from plough layer (0—20 cm) and Nos. 3 and 5 from deeper layer (20—40 cm) of cultivated soils, No. 2 from deeper layer of virgin soil (Table 1). The samples were air-dried and ground to pass a 2-mm sieve. Soil pH was measured in soil-0.01 M CaCl₂ suspension (1:2.5 v/v) (RYTI 1965). The organic carbon content was determined by a modified (GRAHAM 1948) ALTEN wet combustion method. The particle-size distribution of the inorganic matter of soil was determined by the pipette method (ELONEN 1971). The amorphous aluminium and iron were extracted by acid ammonium oxalate (0.18 M ammonium oxalate, 0.10 M oxalic acid, pH 3.3, 1:20 w/v) (TAMM 1922) and determined by atomic absorption spectrophotometry.

The soils were extracted by a modified CHANG and JACKSON (1957) fractionation procedure using a slightly alkaline NH₄F (pH 8.5) as recommended by FIFE (1959). The extracts were analysed for phosphorus by a molybdenum blue method modified by KAILA (1955) and for aluminium and iron by atomic absorption spectrophotometry. Fractionation was carried out in triplicate.

Results and discussion

In the fractionation procedure, the extractability of phosphorus, aluminium and iron

by 1 M NH₄Cl was poor (Table 2). According to KAILA (1964, 1965) and HARTIKAINEN (1979), the extractability of phosphorus by 1 M NH₄Cl is generally low in Finnish mineral soils. As an anion of a strong acid, Cl⁻ cannot participate in ligand exchange reactions with phosphate complexed by aluminium and iron oxides. The content of phosphate extractable by NH₄Cl is worth mentioning only when the sorption capacity of soil is covered with phosphate. In the experimental soils, however, the ratio of fractionated phosphorus to oxalate-soluble aluminium and iron was low.

According to KAILA (1964), the occurrence of phosphorus in the forms soluble in NH₄F and NaOH is to some extent regulated by the molar ratio of active aluminium and iron contents in Finnish soils. More phosphorus was extracted by 0.1 M NaOH than by 0.5 M NH₄F from soils No. 3 and 4 which contained more oxalate-extractable iron (mmol/kg soil) than aluminium (Table 2). In the other soils, the NH₄F-soluble fraction was greater than the NaOH-soluble one.

No aluminium was found in NH₄F extracts and iron was poorly soluble (Table 2). The fluoride-soluble iron in soils No. 1 and 2 amounted to 6—7 %, in the other soils to less than 2 % of the oxalate-soluble iron. Fluoride does not measurably complex ferric iron at a pH above 8.0 (FIFE 1959).

Table 2. Soil P, Al and Fe (mmol/kg soil) extracted successively by the CHANG and JACKSON procedure.

	Soil sample				
	1	2	3	4	5
P extracted by 1 M NH ₄ Cl	0.0	0.0	0.0	0.0	0.0
P extracted by 0.5 M NH ₄ F	10.4	2.3	0.8	0.7	0.2
P extracted by 0.1 M NaOH	3.9	0.9	9.8	6.0	0.1
P extracted by 0.25 M H ₂ SO ₄	2.1	4.8	5.1	6.6	12.2
NaOH-P/NH ₄ F-P	0.4	0.4	12.3	8.6	0.5
Al extracted by 1 M NH ₄ Cl	0	0	0	0	0
Al extracted by 0.5 M NH ₄ F	0	0	0	0	0
Al extracted by 0.1 M NaOH	174	78	42	21	7
Al extracted by 0.25 M H ₂ SO ₄	85	17	112	22	27
Fe extracted by 1 M NH ₄ Cl	0	0	0	0	2
Fe extracted by 0.5 M NH ₄ F	3	2	0	3	0
Fe extracted by 0.1 M NaOH	2	2	3	10	1
Fe extracted by 0.25 M H ₂ SO ₄	69	31	196	87	30

NaOH is frequently used in extraction of humic matter and oxides of aluminium and silica (JACKSON 1965). In the experimental soils, 0.1 M NaOH-soluble aluminium amounted to over 50 % (55—94 %) of oxalate-extractable aluminium, NaOH-extractable iron only to 1—13 % of oxalate-extractable iron. According to ALEKSANDROVA (1960), the solubility of iron in 0.1 M NaOH is low. Because humic matter is extracted by NaOH and alkaline NH₄F, it is possible that the iron released by these reagents is derived from humic complexes.

The proportion of H₂SO₄-soluble phosphorus was high in soils No. 2, 4 and 5 which were predominantly coarse (Table 2). About half of the fractionated phosphorus in soils No. 2 and 4 was extracted by H₂SO₄. In soil No. 5 of low oxalate-soluble aluminium and iron content, phosphorus was mainly H₂SO₄-soluble. In soils No. 1 and 3 of high aluminium and iron content, H₂SO₄ extracted 13 and 33 % of the fractionated phosphorus, respectively.

Amplify of aluminium and iron was extracted by 0.25 M H₂SO₄ (Table 2) which is an effective extractant of iron oxides (HSU 1964). H₂SO₄-soluble aluminium in soils No. 1 and 2 amounted to 46 and 16 %, respectively, and in soils No. 3—5 to 96—245 % of oxalate-extractable aluminium. In the fractionation

procedure, iron was mainly released by H₂SO₄, 62 % in soil No. 4 and 97—272 % of oxalate-extractable iron in the other soils.

In the course of the fractionation procedure, large amounts of aluminium and iron were released besides phosphorus by 0.1 M NaOH and 0.25 M H₂SO₄, release of the latter being particularly drastic. In total, these two reagents extracted aluminium and iron in amounts corresponding to 91—309 % and 70—285 % of oxalate-extractable aluminium and iron, respectively.

CHANG and JACKSON (1957) developed their procedure for fractionation of soil phosphorus into discrete chemical forms using aluminium, iron and calcium phosphate minerals variscite, strengite and apatite as controls. However, variscite and strengite are not likely to occur in normal agricultural soil. The solubility product of variscite controls the phosphorus concentration in solution only when the pH of the equilibrium solution is below 3.1 (BACHE 1963). At higher pH values, variscite dissolves incongruently, whereby a more basic solid phase of aluminium hydroxyphosphate is formed (TAYLOR and GURNEY 1962 a, b, 1964). Strengite is never likely to be in equilibrium with any soil solution (BACHE 1963).

According to the modern concept, adsorbed phosphate is more important in soil than dis-

crete phosphate compounds. In acid soil, phosphate is largely adsorbed through ligand exchange onto surfaces of aluminium and iron oxides. From this viewpoint, the soil phosphorus available is best extracted by solutions which release phosphate through ligand exchange without dissolution of aluminium or iron from the oxide surface. In the fractionation procedure the alkaline ammonium fluoride was the extractant best fulfilling these pre-assumptions. The oxides of aluminium were largely dissolved by NaOH and H₂SO₄, the latter dissolving effectively also iron oxides. According to KHANNA and ULRICH (1967), in

acid soils, the H₂SO₄-soluble phosphates cannot be designated solely as calcium phosphates. They may also include acid-soluble portions of occluded phosphates.

Although the selectivity of the extractants for different forms of phosphate is limited (BROMFIELD 1967 a, b, VAHTRAS and WIKLANDER 1970) and it varies in the original phosphate fraction during extraction (BROMFIELD 1970, RAJENDRAN and SUTTON 1970), the fractionation scheme of CHANG and JACKSON does, however, provide information on the general trends of phosphate transformation reactions.

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Fosforin, aluminiumin ja raudan uuttuminen fraktioitaessa maan epäorgaanista fosforia

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Fosforin, aluminiumin ja raudan uuttumista mukailussa Changin ja Jacksonin fraktioinnissa tutkittiin viidellä kivennäismaalla. Fraktioinnissa uuttuneiden aluminiumin ja raudan pitoisuuksia verrattiin happamalla ammoniumoksalaaatilla uuttuvan aluminiumin ja raudan pitoisuuksiin. Fosforin, aluminiumin ja raudan uuttuminen 1 M NH_4Cl :lla sekä aluminiumin ja raudan uuttuminen emäksisellä 0.5 M NH_4F :lla oli vähäistä. Fluoridiuttoisen fosforin (0.2—10.4 mmol/kg maata) ja 0.1 M NaOH:lla uuttuvan fosforin (0.1—9.8 mmol/kg maata) osuudet fraktioinnissa uuttuvasta fosforista riippuivat

oksalaaattiuttoisen raudan ja aluminiumin moolisuhteesta. H_2SO_4 :lla uuttui fosforia 2.1—12.2 mmol/kg maata. Aluminiumin uuttuminen 0.1 M NaOH:lla (7—174 mmol/kg maata) ja 0.25 M H_2SO_4 :lla (17—112 mmol/kg maata) vastasi 55—94 % ja 16—245 % oksalaattiuttoisesta aluminiumista. NaOH-uuttoinen rauta (1—10 mmol/kg maata) ja H_2SO_4 -uuttoinen rauta (30—196 mmol/kg maata) vastasivat 1—13 % ja 62—272 % oksalaattiuttoisesta raudasta. NaOH:lla ja H_2SO_4 :lla uuttui yhteensä 91—309 % oksalaattiuttoisesta aluminiumista ja 70—285 % oksalaattiuttoisesta raudasta.