

Current primary pedogenesis on Devonian sandstone in southern Estonia

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Abstract. A profile of Eutric Cambisol, formed during 50–55 years on the cross-bedded whitish-grey Devonian sandstone of the Burtnieki Stage, was studied on the excavated outcrop on the bank of the Helme River at Helme, southern Estonia. The topsoil of 10 cm, being the A–Bw subsection, represents the result of humus accumulation, pedogenetic weathering of the sand fractions, and the accumulation *in situ* of silt and clay in the initial sand column of 6.18 cm. Considerable loosening, accompanied with the upward expansion of the solum, has taken place at an annual rate of 0.7–0.8 mm. As a consequence of interfraction changes, 260–290 g m⁻² yr⁻¹ of fine textural particles and 100–120 g m⁻² yr⁻¹ of chemical constituents have accumulated in the thin solum, while some air deposition and surface lateral inflow cannot be excluded. The annual input of organic carbon at a rate of 25–28 g m⁻² has operated as the driving force for rapid and intensive soil formation of accumulative origin. Fulvic acids have played a great role in the progress of initial pedogenesis, while the formed fulvates bound with alkaline earths favour the development of accumulative phenomena. Increase in base exchange capacity of the solum several times is the direct result of accumulation of humus, silt, and clay.

Key words: primary pedogenesis, Devonian sandstone, accumulative soil processes, Cambisol.

INTRODUCTION

Soil formation on outcropping rocks and deposits of any origin has always been in the focus of studies of primary ecosystem processes. The use of archaeological objects has made it possible to ascertain the rate of soil formation during centennia and/or millennia (Aleksandrovski 1983; Holliday 1985; Reintam 1994, 2001b; Reintam & Lang 1999). Detritus and tailings of opencast quarries represent suitable objects for the study of initial soil formation during several years and/or decades (Schafer et al. 1980; Roberts et al. 1988a, 1988b; Fedorets et al. 1998; Reintam & Kaar 1999; Tribis 2000; Reintam et al. 2001). Special experiments have also been performed and studied having this aim in view (Gagarina &

Tsyplenkov 1974; Graham et al. 1995; Reintam 1995, 1997; Tice et al. 1996; Beschow et al. 2000; Romanya et al. 2000). All these investigations demonstrate the relative rapidity of contemporary soil formation.

At the end of the 1990s we carried out recurrent studies in some areas of the International Biological Programme to establish contemporary pedogenetic changes in several soil types (Reintam 1999, 2000, 2001a). Modern argillization as well as rapid argilluviation were found to have occurred within two–three decades. Clear 33-year contemporary podzolization was evident in pure sand overgrown with green mosses and covered with the newly formed forest floor during this time (Reintam 2001a). Similar phenomena have developed under afforested dunes in San Francisco (Amundson & Tremback 1989) and in the Falkland Islands (Wilson 2001). Thus primary pedogenesis in limestone bedrock and Quaternary deposits as well as contemporary genetic changes in particular soils have been a rather frequent object of research and discussion. However, as exposures of Devonian sandstones are rather rare, we had no data about initial soil formation in these materials. Still, some information has been available for corresponding buried palaeosols (Williams et al. 1996).

This study focuses on soil formation on light-coloured Middle Devonian sandstone in the bottom of a former opencast within about 50–55 years. The object was discovered occasionally on the route of a soil expedition to southern Estonia in 1998.

MATERIAL AND METHODS

Location, site, soil, and sampling

Devonian sandstone of the Burtneki Stage (Kleesment & Mark-Kurik 1997) crops out at Helme, southern Estonia (58°01' N, 25°53' E) at the northern verge of the valley of the Helme River. Cross-bedded slightly cemented whitish-grey (10YR7/1) and slightly yellowish-grey (10YR7/6) Middle Devonian sandstone is traversed by reddish-brown stripes (5YR5/8) which are partly inclined and partly intersect the former (Fig. 1). According to oral information obtained from local citizens, sand was excavated here in the late 1930s. In 1939–42, the vegetation was absent and pure light-coloured sandstone was exposed. Although its excavation was already prohibited, digging still continued in places and two outcrops formed. One is located about some tens of metres north of the village road at the level of the inter-valley plateau, the other above the waterlogged bottom of the river valley, right beside the road (Fig. 2). The flat area between these outcrops is covered with herbs and grasses and some grey alder (*Alnus incana*) bushes. Two elderly local inhabitants suggested that the vegetation had formed not earlier than 55 years and not later than 45–50 years ago. As far as they could remember, the first traces of soil were evident about 50–55 years ago. Considering these suggestions, we used 50–55 years as time in our calculations, because it was impossible to find more exact documented data about the spontaneous appearance and development of both the vegetation and the soil.



Fig. 1. A 50–55-year-old A–Bw–BC–C profile of Eutric Cambisol on light-coloured cross-bedded Devonian sandstone of the Burtneki Stage. Photo by Illar Lemetti.



Fig. 2. Two outcrops of light-coloured cross-bedded Devonian sandstone. Photo by Illar Lemetti.

A soil section has formed in the thin top of the opencast (Fig. 1). As a result of humus accumulation, the depth of the A-horizon (10YR3/1) is 8 cm; it is commonly rooted by fine and medium roots, and its boundary is gradually clear and smooth. The next, brown Bw-horizon (10YR5–6/4) with a slightly irregular boundary is silty, *in situ* enriched with humus and contains moderately few fine roots. Although the horizon is only 2–3 cm thick, it is clearly pedogenetic and can be qualified as cambic. The original cross-bedded lamination has disappeared and/or been disguised both in the A- and Bw-horizons during initial soil formation. The transitional BC-horizon (10YR7/3) is residually cross-bedded across the whole of its depth of 10–13 cm. Some channels of fine and medium roots are greyish-brown (10YR5/4) owing to accumulated *in situ* humus substances inside them. Abundant sparkling grains of muscovite are present through the entire section, indicating that the soil has formed on Devonian material. Deeper than 20–23 cm occurs practically unchanged light-coloured Devonian bedrock. Sporadic occurrence of such light varieties is characteristic of the “Old Red” area, especially at lagoon sites where the formation of iron films on fine quartz grains was inhibited.

Well-known methods of soil science were applied in soil sampling by formed genetic horizons as well as in their morphological description (Schoeneberger et al. 1998). Each sample was compiled as an average of ten individual samples from different points of the respective horizon and/or initial Devonian sandstone. The bulk density of the soil horizons and initial sandstone was determined in four replications using a steel cylinder of 50 cm³. Accuracy requirements for sampling, laboratory techniques, and measurements were satisfied as for any reference profile (Batjes & van Engelen 1997).

Analyses

Laboratory analyses of the soil were performed at the Institute of Soil Science and Agrochemistry, Estonian Agricultural University. Fine earth with particle size <1 mm was used. The grain-size fraction of 1–2 mm was absent. Samples for the determination of particle size were treated with sodium pyrophosphate to break down aggregates. Sands were sieved and fractions <0.05 mm were determined by pipette analysis (*Pipette Apparatus Table Model 7 Samples*). Total chemical analysis was carried out after alkaline fusion treatment. Iron and aluminium were detected with sulphosalicylic acid and aluminone, respectively, alkaline earths, potassium, and sodium by flame photometry. Carbonates were determined acidometrically for the recalculation of the measurement data. The obtained results were expressed for ignited noncalcareous material (Arinushkina 1970; van Ranst et al. 1999). The reliability of these techniques was repeatedly verified by atomic absorption spectrometry.

The group and fractional composition of humus was determined by alternate acid–alkaline treatment using the Tyurin–Ponomareva volumetric method (Ponomareva 1957). The results were expressed in the percentages of organic carbon.

Group composition represents humic acids, fulvic acids, and humins (insoluble residue). Fractions within the groups are the following: 1a – free fulvic acids (only in the group of fulvic acids); 1 – humic and fulvic acids bound with mobile R_2O_3 (Al_2O_3 and Fe_2O_3); 2 – humic and fulvic acids bound with alkaline earths (Ca, Mg); 3 – humic and fulvic acids bound with immobile R_2O_3 and clay material. The humic:fulvic ratio (HA:FA) represents the integral parameter of the maturity and mobility as well as humicity (>1 , according to some authors >0.7) and/or fulvicity (<0.7) of humus. The ratio of the 1st to the 2nd fraction characterizes the ratio of brown humic–fulvic complexes, bound with mobile sesquioxides, to grey (black) complexes bound with calcium and magnesium. Decalcification with 0.05 M sulphuric acid forms part of humus fractioning, which enables the determination of the second and the third fraction as well as the hydrolysate of 0.5 M sulphuric acid (Ponomareva 1957). Decalcinate contains metals released from humates and fulvates (salts of both humus acids).

Total amounts of organic carbon and nitrogen were ascertained by the Tyurin and Kjeldahl methods, respectively. It is necessary to point out that the Anne method known in the Western countries (van Ranst et al. 1999) is equivalent to the Tyurin method used in this study. Tithionite-extractable (total nonsiliceous) iron (Fe_d), oxalate-extractable amorphous iron, aluminium, and silica (Fe_o , Al_o , Si_o) were measured after Coffin and Tamm, respectively; iron activity was calculated after Schwertmann (van Ranst et al. 1999). Base exchange capacity (BEC) and exchangeable bases were measured by the percolation of the samples with ammonium acetate at pH 7.0 and were expressed in $cmol\ kg^{-1}$. The pH of both water and 1 M KCl suspensions was measured potentiometrically with the pH-meter *Jenway 3071*. Exchange acidity ($H^+ + Al^{3+}$) was ascertained after Daikuhara–Sokolov in 1 M KCl (Kitse & Rooma 1984).

The quantitative origin of current changes was established by the material balance method (Targulian et al. 1974; Rozanov 1975; Reintam 1985, 1997; van Breemen & Buurman 1998). The weight of the soil horizons and their textural and chemical constituents as of 1998 were compared with those recorded for initial Devonian sandstone at depths of 30–40 cm. According to Targulian et al. (1974) and Rozanov (1975), the sand fractions >0.05 mm were regarded as an indicator of the initial status prior to pedogenesis. Changes in the thickness (depth) of the initial column of the soil horizons with an area of one square metre were calculated from the measurement data using formulas published elsewhere (Targulian et al. 1974; Rozanov 1975; Reintam 1997).

RESULTS AND DISCUSSION

Synchronously with the progress of the vegetative cover, organic matter accumulated in the thin top of initial sandstone. The total amount of organic carbon in the humus horizon (epipedon) with a thickness of 8 cm is about $1.4\ kg\ m^{-2}$ (Table 1). Average annual increment is $25\text{--}28\ g\ m^{-2}$ of organic carbon

and 2.6–2.9 g m⁻² of nitrogen against the background of a thickness increase of 1.4–1.6 mm yr⁻¹. The low C:N ratio indicates the presence of nitrogen bridges in polyphenolic molecules and a rather high maturity of humus. Except for thickness, all parameters of epipedon humus satisfy the criteria of mollic (Driessen et al. 2001). According to Romanya et al. (2000), afforestation of Mediterranean Eutric Cambisol on granodiorite resulted in about 10 g m⁻² yr⁻¹ increase in organic carbon in the topmost 5 cm, while in the ectoorganic layers the respective increase was similar to that observed in our object.

The situation is almost similar in the very thin but clearly developed cambic Bw-horizon. Although the contents of organic carbon and nitrogen in it are much smaller than even per cm of the epipedon, both organic constituents are

Table 1. Humus composition in the percentages of organic carbon

Characteristic	A-horizon	Bw-horizon
Thickness, cm	8	2
Bulk density, Mg m ⁻³ *	1.06	1.55
Colour	10YR3/1	10YR5–6/4
Weight of horizon, kg m ⁻²	84.8	31.0
Content of organic carbon, g kg ⁻¹	16.6	4.5
Pool of organic carbon, g m ⁻²	1 407.7	139.5
Content of total nitrogen, g kg ⁻¹	1.7	0.8
Pool of total nitrogen, g m ⁻²	144.2	24.8
C : N	9.8	5.6
Humic acids (HA) 1	1.8	2.2
2	6.6	11.1
3	2.4	2.3
Total	10.8	15.6
Fulvic acids (FA) 1a	2.4	8.9
1	16.3	6.7
2	31.9	28.9
3	7.8	8.9
Total	58.4	53.4
Hydrolsate of 0.5 M H ₂ SO ₄	9.6	11.1
Total soluble	78.8	80.1
Insoluble	21.2	19.9
Humic acids : fulvic acids	0.2	0.3
First fraction : second fraction	0.5	0.2
Decalcinate, g kg ⁻¹ Fe	0.4	0.4
Ca	6.6	2.4
Mg	6.0	2.6

* The bulk density of the BC-horizon and Devonian sand (C-horizon) is 1.64 Mg m⁻³.

of *in situ* origin. Undoubtedly, such a weathered horizon, enriched with humus, represents the first product of primary pedogenesis. Against the background of the progressing accumulation of humus the development of the ochric epipedon takes place on the basis of the Bw-horizon. Gradual development of both humus accumulation and weathering appears to favour the penetration of both horizons into depth.

Humus is clearly fulvic in both horizons (Table 1), while Ca-fulvates are prevalent. As in any outcome of primary pedogenesis (Reintam 1995), a large proportion of humus can be attributed to fulvic acids bound with stable sesquioxides and secondary minerals (3rd fraction) as well as occurring in the interlayer structure of the latter (hydrolysate). The content of free fulvic acids (fraction 1a) shows considerable increase in the Bw-horizon, favouring there, through organic-mineral interactions *in situ*, an intensive development of cambic properties in depth. The amount of humic acids is significantly smaller, while about 60–70% of them are bound with alkaline earths (Table 1). However, the amount of humic acids is still sufficient to ensure the stability of humus, the loosening of the mineral stratum, and the progress of the initial epipedon. Aluminium is not yet mobilized, and humic-fulvic complexes of the first fraction are bound only with iron.

The pH is uniformly neutral across the entire profile (Table 2). The low exchange acidity of Devonian sandstone has completely disappeared from the formed solum. This can be explained by the intensive formation of the colloidal complex and by the rapid progress of BEC which is more than ten times higher in the A- and Bw-horizons than in initial sandstone at a depth of 30–40 cm. Exchangeable calcium accounts for 88–89% of total BEC in the soil horizons and 85–87% in parent material. Average annual increase in BEC is 0.15–0.16 and 0.10–0.11 cmol kg⁻¹ for the A- and Bw-horizons, respectively. Accumulativity of exchangeable bases within only some decades of primary pedogenesis is characteristic of any initial soil (Beschov et al. 2000), among them even sandy minipodzol under a forest vegetation (Reintam 2001a). Exchange absorption of bases which are bound biologically and released in the process of the transformation of organic residues is more intensive under grasses in the conditions of comparatively high pH values and alternating moistening and drying-up typical of sands.

Table 2. Exchange properties, cmol kg⁻¹

Horizon and depth, cm	Ca ²⁺	Mg ²⁺	K ⁺	Na ⁺	BEC	H _{5,6} *	pH _{H₂O}	pH _{KCl}
A 0–8	7.62	0.84	0.14	0.05	8.65	0	7.1	7.1
Bw 8–10	5.09	0.54	0.12	0.02	5.77	0	7.1	7.1
BC 10–20	1.51	0.12	0.06	0.01	1.70	0	7.2	6.9
C 20–30	1.09	0.11	0.06	0.01	1.27	0.31	7.2	6.5
C 30–40	0.47	0.03	0.03	0.01	0.54	0.10	7.3	6.8

* Exchange acidity.

Like decalciate, amorphous aluminium is practically lacking in the oxalate extraction (Table 3). Its mobilization appears to be inhibited in the conditions of neutral reaction and complete base saturation. Only some traces of it were found in the oxalate extraction. The accumulation of amorphous iron is also rather slow. About two-thirds of it are subjected to rapid crystallization while in the BC-horizon this process is many times more intensive. Possibly there occurs hydrothermal gradient for inhibition of the persistence of amorphous compounds. According to Wang et al. (1989), appreciable amounts of crystalline iron oxide may slow down any eluvial process in the soil. The same applies also to silica.

Changes in the granulometric composition are typical of weathering *in situ* under the impact of organic substances and hydrothermal agents. Although sand fractions are of slightly different distribution in cross-bedded Devonian layers, their sum is rather similar (Table 4). Clay is lacking; only brown-coloured internal stripes and lines contain some clay. During the initial soil formation of 50–55 years, silt and clay have accumulated as a consequence of the disintegration of fine sand, while coarse and medium sands appear to have been residual in cross-bedded structure where current soil formation took place. The distribution of sand and silt + clay is completely balanced deeper than 10 cm. This phenomenon indicates that soil processes have affected the exchangeable complex and

Table 3. The content and pool of nonsiliceous substances

Horizon and depth, cm	Content, g kg ⁻¹				Fe _o /Fe _d	Pool, g hor ⁻¹ m ⁻²		
	Fe _d	Fe _o	Al _o	Si _o		Fe _d	Fe _o	Si _o
A 0–8	3.6	1.3	traces	0.6	36	305	110	51
Bw 8–10	2.7	1.0	traces	0.2	37	84	31	6
BC 10–20	4.3	0.4	traces	0.0	9	705	66	6
C 20–30	1.0	0.4	traces	0.1	40	164	66	16
C 30–40	1.6	0.3	traces	0.1	19	262	49	16

Table 4. Grain-size fractions (in mm) and their pool

Horizon and depth, cm	Content, g kg ⁻¹					Pool, g m ⁻² hor ⁻¹		
	1–0.5	0.5–0.25	0.25–0.05	0.05–0.002	<0.002	Sand	Silt	Clay
A 0–8	34	200	618	146	2	72 250	12 381	170
Bw 8–10	20	205	554	166	55	24 149	5 146	1 705
BC 10–20	6	114	801	77	2	151 044	12 628	328
C 20–30	6	127	805	62	0	153 832	10 168	0
C 30–40	1	22	928	49	0	155 964	8 036	0
Brown line	1	19	916	32	32	Impossible to detect		
Grey stripe	2	66	874	58	0	because of small thickness		
C 50–60	4	53	883	60	0	154 160	9 840	0

nonsiliceous substances but have not had any impact either on bulk density (uniformly 1.64 Mg m^{-3} in the BC-horizon and in the upper layers of initial Devonian sandstone) or on the translocation of the fractions. Only a balanced accumulation of silt and clay at the expense of disintegrated sand has taken place in the BC-horizon (Table 4).

Pedogenetic changes in granulometric fractions are obvious in the two upper horizons. In spite of the contemporary accumulation of silt and clay, considerable decrease in the content of sand fractions has resulted in the 31% loss of the weight of initial sandstone (Tables 4 and 5), which makes $87\text{--}96 \text{ g m}^{-2} \text{ yr}^{-1} \text{ cm}^{-1}$ of the solum. In fact, judging by changes in bulk density and by the accumulation of organic matter (Table 1), a loosening and an upward expansion of the solum have taken place. According to mineral indicators (Targulian et al. 1974; Rozanov 1975; Reintam 1997, 1999), these horizons have been formed on initial sandstone which was 3.82 cm thinner 50–55 years ago than the current solum (Table 5). The average annual rate of upward expansion has been extremely high ($69\text{--}76 \mu\text{m}$), probably not only due to the common root system and accumulated humus but also because of the activity of fauna, as suggested by Kristiansen & Amelung (2001) in their study on the effect of the activity of ants on soil status.

After recalculations of the depths of sand for the A- and Bw-horizons (4.63 and 1.55 cm, respectively), the solum proved accumulative at the expense of silt in the A-horizon and silt and clay in the Bw-horizon altogether by 14% of initial sandstone (Table 5). The average annual increase in soil mass is $263\text{--}289 \text{ g m}^{-2}$, of which $161\text{--}177 \text{ g m}^{-2}$ can be ascribed to the A-horizon. Partial input of silt by air deposition and/or surface flow cannot be excluded here. However, 38–39% of annual increase in the content of silt and clay can be attributed to the cambic Bw-horizon where only some lateral removal of clay could have taken place. Although orientated clayskins are still lacking, it is possible that the Bw-horizon has been formed at the initial site of the brown stripe originally enriched with clay.

Table 5. Changes in the pool of grain-size fractions

Horizon and thickness, cm	Pool of natural soil, $\pm \text{g m}^{-2}$				CTS, cm	CPS, g m^{-2}	Corrected pool, $\pm \text{g m}^{-2}$			
	Sand	Silt	Clay	Total			Sand	Silt	Clay	Total
A 8	-52 521	+5 952	+170	-46 399	4.63	75 932	+39	+8 660	+170	+8 869
Bw 2	-7 044	+3 539	+1 705	-1 800	1.55	25 420	-26	+3 901	+1 705	+5 580
BC 10	-4 920	+4 592	+328	0			No changes or corrections			
Total	-64 485	+14 083	+2 203	-48 199	6.18	101 352	+13	12 561	1 875	+14 449

CTS – corrected thickness of initial sand; CPS – corrected pool of initial sand.
For the pools of the fractions see Table 4.

Owing to the different distribution of seams with iron films, aluminosilicates, and micas, the chemical composition of cross-bedded Devonian sandstone is rather variable regarding iron, aluminium, magnesium, and potassium (Table 6). Chemical poverty tends to increase with depth. Nevertheless, differences in the sum of the elements between the horizons do not exceed 3–4% (Table 7). The high chemical variability noted in tailings of opencast quarries by Fedorets et al.

Table 6. Chemical composition and pools of the elements

Element	Content in horizons, g kg ⁻¹					Pool in horizons, g m ⁻² hor ⁻¹				
	A 0–8	Bw 8–10	BC 10–20	C 20–30	C 30–40	A 0–8	Bw 8–10	BC 10–20	C 20–30	C 30–40
Si	414.8	413.6	422.7	403.6	438.7	35 175	12 822	69 323	66 190	71 947
Fe	7.5	7.1	5.4	10.3	3.3	636	220	886	1 689	541
Al	27.1	33.2	31.8	47.3	18.5	2 298	1 029	5 215	7 757	3 034
P	5.8	3.0	1.1	1.5	0.7	492	93	180	246	115
Ca	1.6	4.1	6.1	8.9	8.6	136	127	1 000	1 460	1 410
Mg	7.2	3.5	2.0	4.0	1.6	611	108	328	656	262
K	14.2	15.1	13.6	22.2	10.7	1 204	468	2 230	3 641	1 755
Na	1.8	1.3	0.9	1.3	1.3	153	40	148	213	213
Total	480.0	480.9	483.6	499.1	483.4	40 705	14 907	79 310	81 852	79 277

Table 7. Changes in the pool of chemical elements during 50–55 years in comparison with the corrected thickness of initial Devonian sand, ±g m⁻² hor⁻¹

Element	A+Bw 10 cm	C* 6.18 cm	Difference ±g m ⁻² hor ⁻¹	C** 6.18 cm	Difference ±g m ⁻² hor ⁻¹	C for BC 10 cm**	Difference ±g m ⁻² hor ⁻¹
Si	47 997	44 463	+3 534	42 689	+5 308	69 069	+254
Fe	856	334	+522	689	+167	1 115	-229
Al	3 327	1 875	+1 452	3 943	-616	5 395	-180
P	585	71	+514	112	+473	180	0
Ca	263	872	-609	892	-629	1 435	-435
Mg	719	162	+557	294	+425	459	-131
K	1 672	1 084	+588	1 672	0	2 698	-468
Na	193	132	+61	132	+61	213	-65
Total	55 612	48 993	+6 619	50 423	+5 189	80 564	-1 254
Annual			+120–132		+94–104		-22–25

* The characteristics of initial sand were calculated on the basis of the data from a depth of 30–40 cm, presented in Table 6.

** The characteristics of initial sand were calculated on the basis of the average data from the depths of 20–30 and 30–40 cm, presented in Table 6. These data were also used for the comparison of the BC-horizon.

(1998) is lacking in primary pedogenesis on Devonian sandstone. Obviously, soil formation favours accumulation of iron, aluminium, phosphorus, and magnesium but loss of calcium. However, initial stratification can play some role in this case. Against the background of balance in the textural constituents of the BC-horizon (Tables 4 and 5), a slight removal of elements ($2.2\text{--}2.5\text{ g m}^{-2}\text{ yr}^{-1}\text{ cm}^{-1}$) has taken place there (Tables 6 and 7). Since the annual losses of merely calcium and potassium amounted to $8\text{--}9\text{ g m}^{-2}$ per 10 cm, being about two to three times lower for the other elements, general chemical relationships can be interpreted either as slightly changed or practically unchanged in the BC-horizon. This is not valid for the sum of the A- and Bw-horizons within the upper 10 cm.

After making corrections with respect to the mineral indicators (Tables 4 and 5), it was found that soil formation began 50–55 years ago on top of Devonian sandstone. The thickness of the upper part of the sandstone was only 6.18 cm versus contemporary 10 cm for the two above mentioned horizons. The initial data for the corrected weights of the elements in Devonian sandstone with a thickness of 6.18 cm were taken from Table 6 and are presented, together with the differences between the soil ($A + Bw = 10$) and parent sand (C^* and C^{**}), in Table 7. The total range of profile accumulativity (6–8% of the initial amount) appears to be about twice as low as for textural constituents. This situation can be explained by the existence of chemical constituents in the form of oxides and silicates, containing oxygen, but not in the form of pure elements. As mentioned above, surface and air deposition of silty material can accumulate after the breakdown and removal of calcareous components without respective changes in its chemical status. The appreciable accumulation of phosphorus appears to be of anthropogenic origin and is associated with deposition of silt from the surrounding fields. On the other hand, the average annual gain of about $100\text{--}120\text{ g m}^{-2}$ per solum with a thickness of 10 cm seems to be insignificant and much closer to the desired analytical accuracy.

CONCLUSIONS

Contemporary initial soil formation on light-coloured Devonian sandstone was first detected in Estonia. A well-diagnosable A–Bw–BC profile with a total depth of 20 cm has developed during 50–55 years. As a consequence of the humus accumulation, weathering *in situ*, and loosening of the initial sand column of 6.18 cm, upward expansion progressed at a rate of $0.7\text{--}0.8\text{ mm yr}^{-1}$, forming the A–Bw subsection. The formed epipedon is still ochric but already rather close to mollic. The cambic horizon is thin but clearly expressed. Its presence allows qualification of this pedogenetic outcome as Eutric Cambisol. Ca-fulvic humus has been the driving force for the primary pedogenesis of accumulative origin. The thin A–Bw subsection is enriched with silt and clay as well as with the main chemical constituents. A considerable increase in base exchange capacity and in the crystalline form of nonsiliceous iron is characteristic of initial soil formation on Devonian sandstone.

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Nüüdisaegne primaarne mullateke Devoni liivakivil Lõuna-Eestis

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Esmakordselt Eestis uuriti primaarset mullateket Devoni liivakivil Helmes Helme jõe vasemkaldal liivakivi kaevandamisest tekkinud terrassikujulisel avamusel. Küllastunud liivmulla (Eutric Cambisol) A–Bw–BC-profiil oli kujunenud 50–55 aasta jooksul Kesk-Devoni Burtnieki lademe põimjaskihilisel, punakaspruunide vahetripudega valkjashallil liivakivil. A- ja Bw-horisontidest koosnev 10 cm tüsedune muld oli arenenud algselt 6,18 cm tüsedusel liivakivi kihil liivafraktsioonide murenemise ning huumuse, tolmu ja ibe kohapeal kogunemise teel. Lähtematerjali kobestumisel kasvas A–Bw-profiil keskmise kiirusega 0,7–0,8 mm aastas. Jämedamate fraktsioonide muundumine neis horisontides viis peente osakeste ja keemiliste koostekomponentide kuhjumisele vastavalt 260–290 ning 100–120 g m⁻² aastas. Selle kõrval pole välistatud ka mõningane tolmu ja peenliiva juurdekanne tuule või pindmise külgvoolu teel. Orgaaniline süsinik keskmise aastavooga 25–28 g m⁻² on kiire ja hooga akumulatiivse mullatekke liikumapanevaks jõuks. Fulvohapetele kuulub oluline osa primaarse mullatekke arengus, kusjuures mullatekkes moodustunud Ca-fulvaadid soodustavad akumulatiivsete protsesside järjepidevust. Liivakivist tekkinud mullahorisontide neelamis- mahutavuse mitmekordne suurenemine on põhjustatud huumuse, tolmu ja ibe kogunemisest mullatekkes.