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1 Vegetation influence on soil formation rate in a proglacial chronosequence (Lys Glacier, NW

# 2 Italian Alps)

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7

# 8 Abstract

- 9 Climate change has huge impacts on alpine ecosystems . One of the most visible effects in the Alps
- 10 is glacier retreat since the end of the Little Ice Age (LIA), which caused the exposure of previously
- glaciated surfaces. These surfaces are open-air laboratories, verifying theories regarding ecosystem
   and soil development.
- 13 In order to increase our knowledge on the effects of time and vegetation primary succession on soil
- 14 development in proglacial areas, we sampled soils and surveyed plant communities on stable points
- 15 in the proglacial area of the Lys glacier, in the Italian north-western Alps (Valle d'Aosta Region).
- 16 The sampling points were located on dated sites (based on literature and/or historical photographs).
- 17 Glacial till is attacked by weathering processes immediately after deposition and stabilization, with
- 18 a consequent loss of soluble compounds, decrease of pH and primary mineral weathering. The
- 19 speed of these processes was largely increased after the establishment of a continuous vegetation
- 20 cover, thanks to the organic matter accumulation caused by litter input and root decomposition21 below the soil surface.
- 22 On sialic glacial tills, below timberline and under a quasi-climax Larch Rhododendron forest, a
- 23 fast and steady decrease in pH values, increase in organic matter content and horizon differentiation
- 24 was observed. In particular, genetic eluvial horizons formed in approximately 60 years, while
- 25 diagnostic albic horizons were developed after ca. 90 years, pointing to an early start of the
- 26 podzolization processes. Cheluviation of Fe and, secondarily, Al were analytically verified.
- 27 However, illuviation of Fe, Al and organic matter in incipient B horizons were not sufficient to
- 28 obtain diagnostic spodic horizons on LIA materials.
- 29 Under grazed grassland below timberline and alpine prairie above timberline, acidification and
- weathering were slightly slower, and no redistribution with depth of Fe and Al oxi-hydroxides wasobserved. A cambic Bw horizon developed on the oldest LIA moraines.
- 32 Therefore it seemed that this fast onset of the podzolization process in comparison to other
- 33 proglacial chronosequences in the Alps was mainly driven by vegetation properties rather than by
- 34 specific climatic conditions.
- 35

# 36 Keywords

- 37 Soil formation; proglacial area; soil chronosequence; podzolization; subalpine soils; alpine soils
- 38
- 39 Highlights
- Plant succession and soil development were slower above than below timberline;
- Under grassland below and above timberline, Cambisols were found on old moraines;
- Under typical subalpine forest-shrub vegetation, podzolization was very fast;
- E horizons and "real" Podzols appeared faster than in other forefields in the Alps;

• Vegetation seemed very important in the early podzolization onset.

45

### 46 1. Introduction

47 Climate change is having huge impacts on mountain ecosystems. One of the most visible effect is 48 the glacier retreat, which continued with few interruptions since the end of the Little Ice Age (LIA, 49 between ca. years 1300 and 1821/1861, Ivy-Ochs et al., 2009). The bare surfaces left by retreating 50 glaciers offer the opportunity to observe the early stages of soil development and the primary 51 ecogenetic succession, thus validating existing theories about ecosystem evolution and allowing to 52 determine the speed of soil forming processes. A large number of soil chronosequences in proglacial 53 areas has been studied on mountain ranges worldwide and in arctic and antarctic environments (e.g. 54 Ugolini, 1966, Burt and Alexander, 1996, Egli et al., 2001, Dümig et al., 2011, Mahaney et al., 55 2009, Hodkinson et al., 2003). According to most of these studies, soil chronosequences under 56 boreal/subalpine forests frequently end up with Podzols, but the rate of development of a podzolic 57 morphology and chemistry widely differ in several studies considering different environments. For 58 example, in superhumid basins in New Zealand, with an average annual rainfall higher than 10000 59 mm (Tonkin and Basher, 2001), Podzols were formed in about 1000 years, despite the extremely 60 fast weathering of the gravel and sand fractions and the consequent increase in silt and clay contents 61 in surface horizons. In Alaska, high rainfall and humidity leads to the formation of weakly 62 developed E and Bs horizons after only 70 years, and "real" Podzols after 230 years (Ugolini, 1966, 63 Burt and Alexander, 1996). In these areas, E horizons immediately appeared after the establishment 64 of spruce in the forest succession. This happened despite the presence of abundant carbonatic 65 materials in the parent till. Similar fast rates of Podzol formation (less than 230 years) were 66 observed on marine terraces in Finland (Mokma et al., 2004). 67 Focusing on the European Alps, many studies have been performed on soil and ecosystem

68 development in proglacial areas in subalpine forest habitats (e.g. Egli et al., 2003, Egli et al., 2006,

69 Dümig et al., 2011), while a smaller number was devoted to alpine (i.e. above timberline) habitats,

70 with most of them considering mainly primary plant succession (e.g., Andreis et al., 2001,

71 Caccianiga et al., 2006, Raffl et al., 2006).

72 In the fifties some works described Podzol-like soils on recently deglaciated surfaces also in the

73 Alps: for example, Jenny (1958) showed a 315 years old soil which had a 11 cm thick light gray

74 horizon overlying a 14 cm thick brownish one, under subalpine ericaceous shrubs in the Rhone

- 75 Gletscher forefield. However, in the alpine range, much longer periods since deglaciation were
- required for the establishment of the podzolization process than in other boreal habitats.

77 Under subalpine vegetation, the youngest soils were usually Lithic and Skeletic Leptosols; after 120

- 78 years soil evolution led to the development of Humi-Skeletic Leptosols or, locally, to weakly
- 79 developed Dystric Cambisols (Righi et al., 1999, Egli et al., 2001, Egli et al., 2006). A continuous

80 humus layer appeared in the oldest soils, together with weak signs of chemical weathering

- 81 (formation of Fe and Al oxyhydroxides). Oxalate extractable Fe and Al decreased with depth, and
- 82 no cheluviation was visible (Egli et al., 2006). In the Alps, Dystric Cambisols are normally found

83 on 250-300 years old surfaces, while Podzols appear after 600 years in very humid sites (1800

84 mm/y, Zech and Wilcke, 1977), but usually more than 1300 years are needed for the development of

85 E horizons (Egli et al., 2003). In central Switzerland, in areas characterized by mean annual

86 precipitation equal to ~2000 mm/y, Podzols are found in 3300 years old sites (Egli et al., 2001,

87 Dümig et al., 2011), but they are normally weakly developed (Righi et al., 1999).

88 The establishment of vegetation on the bare deglaciated areas initiates gradients in many soil

89 properties. The primary succession of vegetation in several glacier forefields in the European Alps

90 include pioneer stages dominated by Oxyrietum digynae and Epilobietum fleischeri (Braun-

91 Blanquet 1948), lasting around 30 years, followed by the establishment of an open forest. *Larix* 

92 *decidua*, *Rhododendron ferrugineum* and other Ericaceae start colonizing regolith surfaces after

93 about 20-30 years since deglaciation, but *Rhododendron* generally shows low cover values in most

94 Swiss chronosequences, where the LIA primary succession was normally dominated by *Alnus* 

95 viridis (e.g., Burga, 1999, Föllmi et al., 2009, Burga et al., 2010) (except in the Damma glacier

96 forefield, Bernasconi et al., 2011).

97 Despite these well-established patterns of soil and plant community chronosequences, in many 98 glacier forefields of the Aosta Valley (North-western Italian Alps), both plant community succession 99 (particularly at the subalpine belt) and pedogenesis seem faster than in most other proglacial areas 100 in the Alps. For example, in the proglacial area considered in this study, Rhododendron ferrugineum 101 dominates the understory vegetation already after few decades from glacial retreat (Treter et al., 102 2002). During many soil survey campaigns in Aosta Valley, a well-defined bleached E horizon was 103 commonly recognized in soils developed on 130 years old moraines, associated with weakly 104 developed BCs horizons, slightly enriched in pedogenic oxyhydroxides; this pattern was recorded 105 also in the Miage glacier forefield in the Mont Blanc Massif and in the Verra Grande glacier 106 forefield in the Monte Rosa Group. In these areas, above the treeline, ecosystem and soil 107 development had a similar rate as in other similar alpine environments (Andreis et al., 2001), and 108 "climax" soils and plant communities are reached in longer periods than at lower elevation. 109 Thanks to the well-known chronology of deglaciation of the Lys glacier (e.g., Monterin, 1932), we 110 chose its proglacial area to investigate the rates of the soil forming processes in the Alps. The Lys 111 forefield includes habitats both below (subalpine) and above timberline (alpine), thus evidencing 112 multidirectional trends in pedogenesis and vegetation succession (Huggett, 1998), hence

113 contributing to the detection of the main driving factors of these processes.

114 The comparison between contrasting subalpine and alpine primary vegetation successions and soil 115 chronosequences provides useful information to evaluate the importance of pedoclimatic and 116 vegetation spatial variability in driving the direction of pedogenic processes.

117

### 118 2 Material and Methods

# 119 **2.1 Study area** 120

121 The proglacial area of the Lys Glacier is located in the upper Lys Valley (Aosta Valley, Pennine 122 Alps, Italy, Figure 1). The morainic systems left by several glacier fluctuations during the Holocene 123 were usually erased by the larger advance of the Little Ice Age (LIA), between ca. years 1350 and 124 1850 (Joerin et al., 2006). The maximum LIA glacier advance was reached in 1821; a secondary 125 advance ended in 1861, when the glacier approached the moraine deposited in 1821, leaving no 126 frontal morainic arc because of river erosion. Since 1861, approximately 1.8 km were left free of 127 ice. A minor advance (1915-1921) left a small morainic arc about 800 m from present day glacier 128 terminus.

129 We sampled a recent soil (P1, ca. 7-10 years old) as representative of fresh, raw till (starting point of 130 soil development and vegetation succession for both alpine and subalpine chronosequences). Above 131 timberline ("alpine" chronosequence), we sampled soils formed at 4 different sites under common 132 vegetation covers (Table 1). Below timberline (subalpine chronosequence), we sampled soils at 5 133 different sites (Table 1); if two different vegetation types occurred on same-age surfaces, we opened 134 a soil pit under each type (i.e. S3, S4, S6 were observed under larch forest with Rhododendron 135 ferrugineum, SG2, SG5, SG7 below grazed grassland). S10 and A6 are on late-glacial (Younger 136 Dryas) till (more than 10000-11000 years old, according to Pelfini et al., 1997). This age is 137 attributed thanks to similarities with most other proglacial areas in the Alps (Ivy-Ochs et al., 2009). 138 We excluded areas visibly disturbed by erosion or deposition processes. All soil profiles were 139 chosen as representative among a much larger number of soil observations by soil coring. Since 140 small scale topographic differences have strong effects on soil development and vegetation 141 succession dynamics (Burga et al. 2010), sites with similar surface rockiness and slope were 142 chosen.

143 Unfortunately, below the present-day glacier front the slope is steep and eroded ("Rocce di
144 Salzen"), and it becomes reasonably stable only 200 m below, where the material was left by the
145 glacier around year 1945. Therefore, no soil has been sampled and analyzed with ages ranging
146 between 5/10 and 60/70 years.

147 The altitude of the proglacial area ranges from 1990 m a.s.l. to about 2480 m a.s.l.. The highest

- 148 morainic ridge (2480 m a.s.l.) was deposited in 1755 (Strada, 1988). The present-day glacier tongue
- 149 is almost 100 m above the natural timberline in the Lys valley, even though young individuals of

150 larch (*Larix decidua*) are found up to 2400 m a.s.l., evidencing an ongoing increase in timberline

151 associated with recent climate change and/or reduction in cattle grazing pressure. The whole Lys

- 152 proglacial area is roughly exposed to the south and only the most ancient subalpine LIA soil profiles
- 153 are located on the northward slopes of the 1821 morainic arc.
- 154 The parent glacial till is made of granitic gneiss and paragneiss belonging to the Monte Rosa nappe,
- 155 with minor (ca. 10%) mafic and ultramafic inclusions derived from ophiolitic outcrops in the
- 156 southernmost portions of the glaciated part of the massif, belonging to the Piedmontese Ophiolitic
- 157 Units (Mattirolo et al., 1951). Based on the observation of the stone fraction, the lithological

158 composition of the glacial till was similar in every stage of the soil chronosequences. Only the pre-

- 159 LIA alpine site (A6) had a higher mafic-ultramafic content (ca. 30% in weight of rock fragments)
- 160 than the other sites.
- 161 The soil moisture regime is udic (Soil Survey Staff, 2010), with a mean yearly rainfall around 1200
- 162 mm (Figure 2) and no dry season (alpine subatlantic climate). The south-north direction of the Lys
- 163 Valley increases the advection of warm, moist Mediterranean air masses from the south, increasing
- summer rainfall, while the proximity to the main Alpine divide allows some spillover of
- precipitation also from the north during strong foehn wind events. The mean annual air temperature
  ranges between ca. 2°C at 1900 m a.s.l. and -1°C at 2400 m a.s.l., while winter average temperature
  is below -4°C (Mercalli, 2003).
- 168

# 169 2.2 Soil description vegetation survey and numerical analysis

- Floristic relevées were performed on 4X4 m square surfaces around the soil pits. The plant species
  were determined according to Pignatti (1992) and single species cover (%) was visually estimated.
- 172 Total vegetation cover (%), proxy for plant productivity, was estimated as well.
- 173 Field description of soil profiles was done according to FAO (2006). Approximately 0.5-1 kg of soil
- 174 material was collected from every horizon in the soil pits. In the field we were not able to obtain
- 175 samples for the calculation of bulk densities because of excessive stoniness and/or the abundant
- 176 presence of medium and/or large roots. The soil chemical and physical analyses were performed
- 177 according to standard methods (Ministero delle Politiche Agricole e Forestali, 2000).
- 178 All samples were air-dried and sieved to separate the fine earth (below 2 mm) from the coarse
- 179 fraction. pH was measured in water (soil:water=1:2.5); total carbon (corresponding to total organic
- 180 carbon, TOC, thanks to the absence of carbonates) and nitrogen (TN) were analyzed by dry

- 181 combustion with a CN elemental analyzer (CE Instruments NA2100, Rodano, Italy). The cation
- 182 exchange capacity (CEC) was measured with the ammonium acetate extraction (pH 7) method, in
- 183 order to classify soils according to the IUSS Working Group (2006). Exchangeable base content and
- 184 saturation, on the ammonium acetate extracts, were measured by AAS (flame atomic absorption
- 185 spectrometer, Analyst 400, Perkin Elmer, Waltham, MS, USA). The particle size distribution was
- 186 determined by the pipette method. In order to detect the spodic properties, the oxalate and
- 187 dithionite-extractable fractions of Fe and Al (Feo, Alo) were measured.
- 188 Chronofunctions of TOC concentration changes in surface horizons, of the Spodicity Index (later
- 189 on, IS, equation 1, Soil Survey Staff, 2010) in CB-BC-Bs horizons and the ratio between the IS in
- 190 subsurface CB-BC-Bs horizons and in surface A-AE-E ones (later on, ISratio, equation 2) in
- 191 subalpine soils were calculated with the *lm* function, included in the R software (R Development
- **192** Core Team, 2000).
- 193 IS = 0.5\*Feo +Alo (1)
- 194 ISratio =  $IS_{(B-BC)}/IS_{(E-AE)}$  (2)
- 195 As reported above, we could not calculate the soil carbon stock because no bulk density data have 196 been measured. However, we believe that the changes in concentration of the different soil 197 compounds can effectively show pedogenic trends in the studied soils, considering that the skeletal 198 fraction resulted quite constant, particularly in the first few hundred years of soil development. The best variable transformation (logarithmic or power) was chosen according to the R<sup>2</sup> and the 199 200 significance of the regression coefficients. The chronofunctions were only descriptive, as the 201 sampling site number was excessively small to obtain statistically significant data. Moreover no 202 data were available for the 260-11000 years BP time span, and the precise ages of pre-LIA S10 and 203 A6 were not available. Significant differences in many edaphic parameters between different plant 204 covers were also checked and displayed as boxplots, using the *multcomp* R package (Hothorn et al., 205 2008).
- 206

# 207 **3. Results**

- 208
- 209 3.1. Vegetation primary succession
- 210 3.1.1 Alpine primary succession
- 211
- Above timberline (Table 2, Figure 3), two different grassland communities were normally foundoutside the LIA glacial till, roughly corresponding to the lower alpine belt: flat, humid areas were

214 dominated by the *Carex curvula* association (*Caricetum curvulae*, Braun-Blanquet 1948), while

- 215 steeper and drier sites were characterized by *Festuca varia* (*Festucetum variae*, Braun-Blanquet,
- 216 1948); the observed pre-LIA site was colonized by a rather xerophilous community dominated by

**217** *Festuca varia* (A5, Table 2).

218 Immediately after moraine stabilization (5-7 years after glacier retreat, P1), the pioneer species of 219 the *Epilobietum fleischeri* (quite similar to the pioneer community described by Burga et al., 2010 220 in the Morteratsch forefield) began the colonization of the raw till. Some of these species, often 221 typical of base-rich soils (Pignatti 1992), were still present on 60 years old moraines (A2, A3). Mid 222 successional species (Table 1) were common in 60-190 years old sites (A2, A3, A4); these species 223 were characteristic of disturbed, rocky and eroded soils, only weakly acidified in the surface 224 horizon. Species typical of later stages of succession appeared on 60 years old moraines (e.g., Carex 225 curvula, Festuca varia, Festuca halleri), but became more common in later stages (A4, A5). The 226 vegetation growing on 260 years old materials (A5) and the pre-LIA site (A6) was almost 227 completely devoid of early and mid-successional species. The A5 site, in particular, was covered by 228 a hygrophilous facies of the *Caricetum curvulae* (Braun-Blanquet, 1948), rich in dwarf Ericaceae 229 such as Loiseleuria procumbens and Vaccinium uliginosum subsp. gaulterioides, probably because 230 of microclimatic conditions favoring a long-lasting snow cover (north-west aspect).

231

### 232 3.1.2 Subalpine primary succession

Below timberline (i.e., subalpine primary succession, Table 3, Figure 3), the steep surfaces 233 234 deglaciated between ca. 1950 and 1987 (not sampled) were mostly colonized by Salix spp., 235 accompanied by Rhododendron ferrugineum and young and scattered larch trees (Treter et al., 236 2002). An extreme vegetation patchiness existed on surfaces deglaciated between 1922 and 1950 237 (excluding stony mounds, not considered in this study). Under an open Larix decidua forest, 238 surfaces covered mostly by *Rhododendron ferrugineum* (S3, S4) were intermixed with more open, 239 grazed, grass-dominated ones (SG2, SG5), which still included a few species characteristic of 240 pioneer *Epilobietum fleischeri*. Mid-successional species, such as *Salix* spp., were locally common 241 but never dominating. Species typical of subalpine forest-heath communities already showed high 242 covering rate in grassland patches, such as Avenella flexuosa and Calamagrostis villosa. Other 243 common species were typical of subalpine prairie (e.g., Festuca varia, Nardus stricta, 244 Anthoxanthum odoratum). 245 Surfaces deglaciated between 1860 and ca. 1922 (S6) were covered by a quasi-climax subalpine

- 246 Larix decidua open forest, with Rhododendron ferrugineum dominating the understory, together
- 247 with Vaccinium spp; grasses were already characteristic of a well-developed subalpine forest

- 248 (Avenella flexuosa, Calamagrostis villosa, Homogyne alpina). The easternmost part of this area has
- always been used for intensive cattle grazing and only scattered trees and shrubs were present
- 250 (SG7). Here, species typical of subalpine acidophilous grazed grassland were common, such as
- 251 *Nardus stricta, Festuca varia* and *Rhinanthus alectorolophus* (Table 3).
- 252 The 1821 moraine is steep and north-facing; a thick, long-lasting and creeping snow cover
- 253 characterizes the microclimate of this site, which is colonized mainly by green alder (Alnus viridis),
- which partly inhibits high covering values by *Vaccinium myrtillus* and *Rhododendron ferrugineum*,
- which are only locally dominant (S9).
- 256 The older surfaces, originated from glacial till abandoned during late glacial maxima (around 10000
- 257 years B.P., S10) were colonized by a climax open Larch forest, with an understory dominated by
- 258 *Rhododendron ferrugineum* and *Vaccinium* spp. The specific composition and covering values were
- similar to those characterizing most of the stable forest sites in the area older than 90 years.
- 260 The plant cover was already around 100% in 65 years old soils, while above timberline it reached
- the same value after 90 years.

### 262 **3.2** Soil properties along the chronosequences

- Soil development trends were well correlated with the primary succession of the vegetation cover
  and land use; three different pathways were observed under alpine prairie, subalpine grassland and
  typical subalpine forest-heath communities (Figure 4).
- 266

### 267 3.2.1. Alpine soil chronosequence

- 268 Above timberline (Table 4, Figure 4a), organic matter accumulation and acidification in the soil 269 surface horizons were the main pedogenetic processes in soils younger than 200 years. These 270 processes led to the formation of A horizons, with a maximum thickness and organic carbon content 271 in 260 years old soils (A5). The C/N ratio in upper alpine mineral soil horizons ranged between ca. 272 11 (A2) and 17 (A5, influenced by Ericaceae), while during the first stages of soil development 273 subsurface horizons showed much lower values. The difference between surface and subsurface 274 horizons decreased with increasing soil age, suggesting a more efficient organic matter 275 incorporation in the soil profile. 276 Weathering in subsurface horizons, releasing free Fe and Al oxi-hydroxides, led to the formation of
- weakly developed, brownish AB and BC horizons in 190 years old soils (A4). Younger soils had
- 278 only A, AC and C horizons while a "true" brown, structured Bw appeared in 260 years (A5). Well-
- 279 developed pre-LIA soils were characterized by thick and well developed brown Bw horizons with
- 280 strongly acidic pH values, particularly in the A horizon.

- 281 Amorphous and crystalline Fe and Al oxi-hydroxides were weakly redistributed with depth
- 282 (increasing contents in subsurface horizon associated with a depletion in surface ones) in the 260
- 283 years old and in the "late glacial" soils (A5 and A6, Table 5).
- 284 Initial soils (P1) had near-neutral pH values and high Base Status, thanks to the abundance of
- 285 freshly ground, highly reactive primary minerals. Acidification and desaturation proceeded quite
- 286 fast, particularly in surface A horizons, together with the strong increase in organic matter. Base
- status below 50% appeared after more than 65 years.
- 288 According to the WRB soil classification (IUSS Working Group, 2006) the soils up to 65 years in
- the forefield (P1, A2, A3) were classified as Haplic Regosol (Eutric, Skeletic). After 190 years (A4),
- 290 weakly developed but already acidified soils were classified as Haplic Regosol (Dystric, Skeletic).
- 291 On the oldest LIA morainic arc and on late glacial materials (A5 and A6) Haplic Cambisol (Dystric,
- 292 Skeletic) were found, thanks to the presence of well developed Bw horizons (Table 4).
- 293 RhizoMODER humus forms characterized most of the soils older than 60 years, while A5, with an
- ericaceous cover, had a hemiMODER (Zanella et al., 2011).
- 295

### 296 3.2.2. Subalpine soil chronosequences

297 Considering only stable sites below timberline, two different soil chronosequences were observed,298 associated with subalpine forest-shrub vegetation or with subalpine grazed grassland.

299 Under subalpine grazed grassland (Figure 4b), soil processes and horizon formation were similar,

300 but faster, to the ones observed above timberline. TOC accumulation in A horizons increased with

301 age and pH values decreased to  $\sim$ 5.5 (SG7); in the meantime, weakly structured, brownish Bw

302 horizons formed (in 130 years old prairie sites, SG7). Where the vegetation was dominated by

303 grassland species (SG5, SG7) or by Alnus viridis (S8), the C/N values were below 14 (Figure 5a).

304 Under grassland vegetation, the albic horizon did not form, and the Fe and Al oxi-hydroxides did305 not redistribute with depth.

Under subalpine larch forest with Rhododendron understory (Figure 4c), great morphological and chemical changes characterized soils in the first 60-65 years since deglaciation (S3). Up to 6 cm of litter accumulated on the soil surface, below which 3-6 cm thick, dark grayish A horizons developed and weathering created yellowish BC ones (Table 6). These A horizons were characterized by the absence of structure and by the juxtaposition of mineral and organic particles, creating a "salt and pepper" appearance. pH values dropped from 6.5 to 5.0 in the upper horizons, while the C/N values were already close to 20. The C/N values were significantly higher than under other vegetation

- 313 types (Figure 5a) while the pH values were lower (Figure 5b). Also the thickness of all O horizons
- 314 was significantly higher under forest than under grassland (Figure 5c).

315 In the following 30 years (S4, on the 1921 moraine), the A horizons were substituted by thin, visibly 316 bleached E horizons, characterized by a decrease in pedogenic Fe oxi-hydroxides (Table 7). A 317 further decrease in pH values (4.7 in the E horizon) was measured in ~130 years old soils (S6) and 318 the E horizons met the morphological requirements for the diagnostic albic horizons (Soil Survey 319 Staff, 2010 and IUSS Working Group, 2006). Under these albic horizons, weak Bs horizons formed 320 (here called CBs or BCs due to the lack of structure and the light yellowish colors; the 321 macromorphological requirements for the spodic horizon were not met, according to Soil Survey 322 Staff 2010, and IUSS Working Group, 2006). Fe redistribution was evidenced by yellowish colors 323 and by Feo and Fed depth trends (Figure 6). The E and BCs horizons further developed in 190 years 324 old soils (S8, S9), when the redistribution of Al (Alo) became measurable. Higher Feo 325 concentrations in subsurface horizons was verified by the ratio between Feo concentrations in B-326 BC-CB and in surface E-AE-A horizons (Figure 5d), which was significantly higher under forest-327 shrub vegetation also in young soils. The same increasing concentration in subsurface horizons was 328 observed for Fed (Figure 5e). Alo was extremely low in all LIA soils, and its redistribution with 329 depth was not significantly different under the considered vegetation types. Thus, the chemical 330 requirements (TOC > 0.5%, IS > 0.5%, ISratio >= 2) for the diagnostic spodic horizons were not 331 met. The ISratio resulted significantly higher under forest-shrub vegetation than under other land 332 covers (Figure 5f). 333 Pre-LIA climax soils were Podzols with an extremely well developed morphology, both under forest

333 Pre-LIA climax soils were Podzols with an extremely well developed morphology, both under fores
and under anthropogenic grassland (not shown). These Podzols were characterized by a strong
illuviation of organic matter and pedogenic Fe, Al and Si oxides in the spodic horizons. Deep
cemented horizons (ortstein) were generally developed below the Bs (S10).

According to the WRB soil classification (IUSS Working Group, 2006), the soils under subalpine

forest/shrubs on the LIA materials, and soils under grassland up to 90 years old, were classified as
Haplic Regosol (Dystric, Skeletic). Subalpine "climax" soils were Ortsteinic Podzols (Skeletic)

340 (S10). Dystric Cambisols were found on 130 years old surfaces under grazed prairie (SG7).

341 Under subalpine grazed grassland, rhizoMODER humus forms were identified, characterized by the

342 presence of OF and, sometimes, OH horizons, overlying root-rich, single grain A ones. Under

343 Rhododendron-larch forest, the humus form was dysMODER in 60 years old soils (S3). It quickly

344 evolved towards MOR forms (hemiMOR in S4, S6 and S9, euMOR in S8); pre-LIA soils had

345 humiMOR forms (Zanella et al. 2011).

346

### 347 4. Discussion

349

### 348 4.1 Vegetation primary successions along the chronosequences

350 Microtopography governs the establishment of pioneer species and their turnover in the first few 351 hundred years during the primary succession (Burga et al., 2010). Considering only stable, fine 352 earth-enriched sites (as in this study), the trends in the vegetation succession are more linear. 353 As in other proglacial areas worldwide, the early stages of soil development and vegetation 354 succession were similar above and below timberline, as almost the same plant species colonized the 355 "young" morainic till, mostly belonging to the "Epilobietum fleischeri" (Braun Blanquet, 1948). 356 Above the treeline, the development of soils and the vegetation succession was slightly faster than 357 in other proglacial areas described in the Italian Alps (Andreis et al. 2001). The considered time 358 span (260 years) was enough for the establishment of a quasi-climax vegetation (Curvuletum or 359 *Festucetum variae*) while 500 years is the minimum time normally required for the establishment of 360 the climax Curvuletum in the Alps (Andreis et al., 2001). 361 The primary plant succession above timberline was strongly influenced by topography and 362 microclimate: for example, Festuca varia was common on 60 and 190 years old moraines (A3 and 363 A4) and in the climax, pre-LIA site (A6), while it was absent from the cooler and more humid 260 364 years old moraine (A5). It seems thus plausible that the further development of the vegetation cover 365 in the A3 and A4 sites will lead to the development of Festucetum variae, similar to "climax" A6, 366 and not towards a Curvuletum, similar to the one growing on the A5 site. 367 The similarity between primary vegetation successions in areas above and below timberline ends 368 soon, and already after about 60 years, larch trees (Larix decidua) have largely invaded the areas at 369 lower elevations, accompanied by ericaceous shrubs (e.g. Vaccinium spp., Calluna vulgaris and 370 Rhododendron ferrugineum) (similar to the situation described in Alaska by Boggs et al., 2010). 371 The establishment of a *Rhododendron* understory, accompanied by *Salix helvetica* and *Salix* appendiculata, was observed earlier than in other proglacial areas in the Alps (e.g., Bernasconi et 372

al., 2011). A stabilization of the species composition below timberline was observed after the

374 establishment of ericaceous shrubs, particularly *Rhododendron ferrugineum*, on ca. 90 years old

375 surfaces. *Rhododendron ferrugineum* is the main limiting factor to the survival of pioneer species,

376 because of soil shadowing and because of the thick litter layer created by this species (Pornon and

377 Doche, 1996). A quasi-climax vegetation structure and composition was generally reached after

378 100-120 years: a much faster vegetation succession is thus observed in this work, if compared to

379 other well studied chronosequences on the Alps (e.g., Burga et al., 2010, Dümig et al., 2011).

380 The early establishment of a quasi-climax vegetation below the timberline differs from what 381 reported by Burga (1999), Burga et al. (2010) and Föllmi et al. (2009) in the Morteratsch and the 382 Rhone proglacial areas respectively, where the most common shrub was Green alder (Alnus viridis), 383 with associated "megaphorbiae", while Ericaceae (*Rhododendron ferrugineum* and *Vaccinium* spp.) 384 never reached covering values higher than 5%. The dominance by Rhododendron ferrugineum in 385 the Lys proglacial area was observed in sites stabilized before 1945 by Treter et al. (2002), who 386 associated this dominance to light grazing. The ecosystem mosaic, related with microtopography, 387 was particularly visible in the 1921-1945 time span, where stable sites covered by a larch-388 Rhododendron forest were mixed with grazed open grassland patches and stony mounds; quasi-389 climax subalpine shrubs (*Rhododendron ferrugineum* in particular) colonized most of the older 390 surfaces, except where grazing pressure was higher. Stony mounds (not considered in this study) 391 were common in the area younger than 91 years, and were colonized by pioneer species (Treter et 392 al., 2002).

393 Grazed sites, devoid of ericaceous shrubs and Larch trees, were colonized by different herbaceous 394 species compared to alpine grasslands, and the oldest sites had a species assemblage close to the 395 typical subalpine acidophilous *Nardus stricta* prairie. Grazing locally inhibits shrub and tree growth 396 also on "old" surfaces (SG7), but a low-pressure grazing influence on the abundance of *Ericaceae* 397 and scarcity of *Alnus viridis* and associated species can be hypothesized. 398

### **399 4.2 Soil properties along the chronosequences**

Together with the primary vegetation succession, the speed of diagnostic horizon development and taxonomic reference group change (IUSS Working Group, 2006) was faster below timberline than at higher elevation. Under subalpine grazed grassland, in fact, a Haplic Cambisol (Dystric) formed in 130 years (SG7), while above timberline the same taxonomic level was reached in 260 years (A5). 60 years old soils below subalpine grassland were morphologically similarly developed as 190 years old ones above timberline (A4).

406 This higher soil development rate was probably caused by the higher productivity of ecosystems

407 below timberline, thanks to less harsh climatic conditions and by hypothetically weaker

- 408 cryoturbation phenomena. However, cryogenic features were not significantly more developed
- 409 above than below timberline, as demonstrated by the ubiquitous presence of thin and weak silt caps
- 410 (table 1, 3) in shallow subsurface horizons particularly in young soils (process well described by
- 411 Forman and Miller, 1994). Above timberline, silt caps were best developed in 60 years old soils,
- 412 and became less visible with increasing ages, until they disappeared in 260 years old soils. Below
- 413 timberline, silt caps were not evident under forest/shrub vegetation, while thin silt caps were visible

in shallow subsurface horizons in 60 and 90 years old soils and only in deeper C horizons of 130
years old soils under grazed grassland. Here they were not visible in soils older than 130 years. Silt
caps disappearance was probably related to the bioturbation caused by roots, which was only

417 slightly faster below than above timberline.

418 A higher ecosystem productivity below than above timberline was verified by the higher TOC 419 concentration in the fine earth of upper mineral horizons in the 130 years old soil (SG7, table 4, 6); 420 however, its trend with time was disturbed by a large variability. Above timberline, the rate of TOC concentration increase declined from 0.28  $g^{k}g^{-1}y^{-1}$  in 70 years old soils to 0.16  $g^{k}kg^{-1}y^{-1}$  on 260 421 422 years old moraines; at this point, the balance between organic carbon inputs and loss via 423 decomposition in surface horizons (steady state) was reached. Subsurface accumulation continued, 424 in quantity and depth, as shown by the increase in TOC concentration and in thickness of Bw and 425 BC-CB horizons (Table 6). Overall, under grassland the TOC concentration in surface horizons 426 tended to increase for the first few hundreds years until reaching a steady state between inputs (via 427 root and litter decay) and outputs (mineralization, erosion and leaching), according to the indicative 428 chronofunction (Figure 7a,  $R^2 = 0.284$ , p-value < 0.05 for all coefficients):

429 **430**   $TOC = -0.287 + 0.616*\ln(age)$ (3)

433 Lower TOC concentrations characterized surface mineral horizons in forest-shrub soils (Figure 7b), 434 which also showed a completely different TOC variation with time (Figure 7a). The highest TOC 435 concentration was reached in 60 years old soils. After this time, TOC concentrations started to 436 decrease, according to the descriptive chronofunction ( $R^2 = 0.201$ , p-value < 0.1):

437

 $TOC = 1.723 - 0.09 * \ln(age)$  (4)

438 The slowdown of the organic matter accumulation started quite early compared to other 439 chronosequences (e.g. Dümig et al., 2011), because of the quick establishment of the podzolization 440 process under the subalpine vegetation (He and Tang, 2008) and the development of an albic 441 horizon below the litter layer. This trend was not observed above the treeline or under subalpine 442 grazed grassland, because of limited vertical migration of soluble organic molecules and of a lower 443 organic matter production caused by the less productive vegetation. The increasing trend of TOC 444 concentration in the subsurface horizons of forest soils was faster than in the grassland ones (Figure 445 7c). Forest soils had higher TOC concentrations in subsurface horizons (Figure 7d) than grassland 446 soils, probably thanks to a vertical migration of soluble organic compounds (possibly associated to 447 a higher organic matter produced by root decay).

The C/N ratio was significantly related to the vegetation cover, with the highest values observed in
the organic horizons composed of ericaceous shrubs and Larch leaves (e.g. Boettner and Kalisz,
1990). Below the forest-shrub vegetation, the lowest C/N value was found under an understory

451 vegetation dominated by the N-fixing Alnus viridis (S8). Where the alpine prairie was enriched in 452 dwarf Ericaceae, such as Loiseleuria procumbens and Vaccinium uliginosum subsp. gaulterioides 453 (A5), the C/N ratio was higher. This ratio is characteristic of slowly decomposing organic matter, 454 rich in fulvic and low-molecular weight acids, and is associated to low pH values. Indeed the fastest 455 and strongest pH decrease was observed under subalpine forest/shrub vegetation, where values 456 below 5 characterized the surface mineral horizon already after 60 years since deglaciation. Under 457 grazed grassland and alpine prairie the pH reached comparable values after 130 and 260 years, 458 respectively. Since the leaching of organic acids derived from organic matter degradation is the 459 main acidifying factor in proglacial soils (Bernasconi et al. 2011), the slower pH decrease under 460 alpine vegetation could be attributed to the lower biomass production and the stronger disturbances 461 characterizing alpine habitats. However, a weak acidification was visible also in the youngest soils, 462 extremely poor in organic matter, thanks to incipient mineral weathering and leaching caused by 463 rain and snowmelt.

464

### 465 4.3 Evidences of podzolization processes

Both Feo and Alo in the initial soil (P1) were higher in the top C1 horizon than in the underlying
C2. This distribution could be attributed to an initial mineral weathering caused by the organic
matter accumulation produced by the few plants which colonize these sites, and by oxidative
reactions involving fresh, reactive particles. The already detectable Fe and Al oxi-hydroxides in the
fresh material of the C2 horizons in the 6 years old soils could be attributed to subglacial
weathering processes, a well-known source of already weathered material to initial soil formation
processes (Anderson et al., 2000 and Dümig et al., 2011).

473 This depth trend was soon reversed: an early redistribution of pedogenic Fe oxi-hydroxides to

474 subsurface horizons was observed under subalpine larch forest with Rhododendron understory.

475 Higher values of Feo and Fed in subsurface BC horizons compared to the overlying A ones were

476 already observed in the ca. 60 years old soils. Despite the weaker Alo redistribution with depth in

477 young soils, the ISratio tended to increase with time, according to the (qualitative) chronofunction

478 ( $R^2 = 0.801$ , p-value < 0.01 for the age coefficient, Figure 8a):

479 ISratio = 
$$0.676^{*}$$
age<sup>0.185</sup> (5).

480 Thus, around 350 years are needed in order to meet the ISratio requirements for the diagnostic

481 spodic horizon according to Soil Survey Staff (2010) and IUSS Working Group (2006).

482 Observing the IS and Fed depth trends (Figure 5 and Figure 6), the cheluviation seems to have

483 actively translocated Fe and Al oxi-hydroxides into subsurface BC and CB horizons. The IS, in

484 subsurface horizons of soils under subalpine forest-shrubs (*Rhododendron* and *Larch*), increased

(6)

- 485 according to the chronofunction (Figure 8b):
- 486 IS= $0.045 + 0.003 * age^{0.8}$

487 therefore, the time needed was ca. 530 years.

488 Based on the few data available, all the regressions between time since deglaciation and the spodic 489 properties had similar results: the time required to obtain a "chemical" Podzol, according to the Soil 490 Survey Staff (2010) and IUSS Working Group (2006) is around 300-600 years, which is a much 491 shorter period than the 3000 years previously calculated on the Alps (Egli et al. 2001). 492 However, these findings apply only to sites located below the timberline and covered by larch forest 493 with ericaceous understory. Below subalpine grazed grassland and below alpine prairie, Fe and Al 494 oxyhydroxides were more abundant in surface A horizons compared to subsurface BC or Bw 495 horizons. An exception was the 60 years old grassland soil (SG3), influenced by nearby larch trees, 496 and A5, thanks to abundant ericaceous dwarf shrubs (Vaccinium uliginosum subsp. gaulterioides

497 and *Loiseleuria procumbens*).

498 Similarly to other chronosequences (e.g., Burt and Alexander 1996, Egli et al. 2001), the slower 499 release and mobility of Al from the parent material, compared with Fe, could be related to the 500 presence of easily weatherable Fe-rich minerals included in the small amounts of mafic rocks in the 501 parent material. The sialic minerals weathered more slowly in the considered soils, as demonstrated 502 by the Alo/Feo ratio increase with time, both in surface (not shown) and in subsurface horizons 503 (Table 7). Conversely, in more acidic parent materials, the onset of the podzolization process tended 504 to remove Al faster than Fe from the young and weakly developed E horizons (Mokma et al., 2004). 505 The E horizon formation in alpine soils has sometimes been interpreted as a stagnogley feature, 506 caused by seasonal waterlogging and alternation of reductive and oxidative conditions, common in 507 high altitude soils thanks to the abundant snow cover and high water input during the snowmelt 508 (e.g. Gensac, 1990). However, the E horizon formation under subalpine vegetation in the Lys 509 proglacial area cannot be ascribed specifically to seasonal waterlogging, as stronger seasonal 510 reductive conditions should be met above timberline, thanks to deeper winter and spring snow 511 covers. Here, E and CBs/Bs horizons were never found, and Fe/Al redistribution to subsurface 512 horizons normally was not observed. Moreover the coarse texture of the soil material should 513 facilitate drainage, decreasing the waterlogging potential in the topsoil.

514 The Feo/Fed ratio, called "activity ratio" and considered as a measure of the crystallinity of

515 pedogenic Fe oxides (Burt and Alexander, 1996), did not show any significant time trends but had

516 significantly lower values under subalpine forest-shrub vegetation than under grassland (Figure 9),

517 particularly in subsurface horizons. Alternation of reductive and oxidative conditions are known to

favor the crystallization of pedogenic Fe-oxi-hydroxides; however, the lower values found under subalpine vegetation, particularly in subsurface BC and Bs horizons, are not explainable according to surface seasonal redox conditions, for the same reason explained above. Nor the higher crystallinity found in soils under forest vegetation, compared to the soils developed under even more humid climates on the Alps (Dümig et al., 2011), could be easily explainable. Low Feo/Fed values, however, are typical of many well developed podzolic soils in the Valle d'Aosta Region (e.g. D'Amico et al., 2008).

525

526 On the Alps, Podzols are known to develop from silica-rich glacial till in late stages of pedogenesis 527 (3300 and 10000 years), as reported by Egli et al. (2001). Faster rates of E horizon formation have 528 often been measured in other boreal or mountain areas (Sauer et al., 2007). Faster Podzol 529 development in Alaska (Alexander and Burt, 1996, Ugolini, 1966), Norway (Mellor, 1986) or China 530 (He and Tang, 2008) compared to the European Alps was attributed to the maritime climate (Egli et 531 al., 2006, Dümig et al., 2011), which accelerates plant growth, soil organic matter accumulation 532 and, consequently, the rate of soil development, thanks to longer growing season and much higher 533 winter temperature. However, the establishment of mixed coniferous forests caused an immediate 534 onset of the podzolization process also in continental climates (ca. 225 years, Lichter, 1998). An 535 even faster redistribution of Fe oxi-hydroxides was observed on sandy parent materials under boreal 536 Scots pine in north-western Russia (Abakumov et al., 2010), where higher Feo contents in BCs than 537 in AE horizons were measured in 10 years old soils, and E horizons appeared in 20 years; in fact, 538 sandy parent materials are known to increase the speed of the podzolization process thanks to the 539 already weathered grain surfaces (Schaetzl and Anderson, 2005) and higher water mobility that 540 promotes vertical fluxes of elements.

Thus, on a global scale, the "fast" podzolization process characterizing subalpine soils developed on 541 542 the gneissic materials of our study area is in the "normal" ranges, but much faster than in other 543 alpine chronosequences. Similarly developed soils have been observed on the LIA morainic arcs of 544 the Miage glacier, Val Veny, in the Mont Blanc Massif and in the Verra Grande forefield, Monte 545 Rosa Group (D'Amico 2011, unpublished data). These proglacial areas in Valle d'Aosta are 546 characterized by temperature and precipitation regimes similar to other proglacial areas, in 547 particular in the northern parts of the Alps, where the podzolization processes start later and seem 548 slower than in our study area. Thus, climatic differences cannot be the cause of such a slower 549 pedogenesis on the northern side of the Alps, while the early establishment of ericaceous shrubs 550 below the treeline appeared the main driver of the early appearance of the bleached E horizon in 90 551 years old soils.

552 Plant communities on Swiss glacier forefields were dominated by alder (Alnus viridis), normally 553 associated with N-fixing bacterial communities (Egli et al., 2001, Burga et al., 2011), and by 554 herbaceous plant species, while Rhododendron ferrugineum appeared in later stages of succession 555 (Dümig et al., 2011). This vegetation creates a litter which has a weak complexing capacity, while 556 the ericaceous shrubs are able to begin a quick podzolization process in previously non-podzolic 557 soils (Bernier and Gillet 2012, 1993, Boettcher and Kalisz, 1990). This is caused by the slow 558 decomposition rates of the litter of ericaceous shrubs, due to their high amount of lignin, cellulose 559 and other recalcitrant substances, such as phenolic compounds, which reduce the soil biological 560 activity (Pornon and Doche, 1995). The litter of coniferous trees and Ericaceae produces large 561 quantities of low molecular weight and fulvic acids, which cause intense mineral weathering 562 (Schaetzl and Anderson, 2005). As humus forms are considered a good indicator of forest 563 ecosystem functioning (Michalet et al., 2001), the fast development of Mor humus forms under 564 subalpine typical Rhododendron-larch vegetation confirms the slow mineralization of the soil 565 organic matter, typically associated with the onset of podzolization. 566 The strong vegetation effect on pedogenesis could be enhanced by mycorrhizal fungi associated 567 with different plant species. In fact, ectomycorrhizal and ericoid (associated to Ericaceae) fungi are

568 known to increase the weathering rate in surface mineral horizons, particularly under coniferous or 569 ericaceous species, where they form mat-like structure at the boundary between the organic layer 570 and the upper mineral horizon, and extending down to the E horizons (Koele et al., 2011). These 571 fungi are able to dissolve mineral grains, extracting and chelating metals and nutrients via the 572 release of phenolic compounds, low weight organic acids, oxalate, citrate and malate (Landeweert 573 et al., 2001), which have a stronger acidifying and weathering capacity than humic molecules 574 (Ochs, 1996). Ericoid mycorrhizal fungi, in particular, produce siderophores molecules, able to efficiently extract and bind Fe and other metals from primary metals (Hoffland et al., 2004). 575 576 Substances produced and released by ectomycorrhizal fungi, thus, increase the podzolization rate 577 under coniferous trees (Lundström et al., 2000, van Breemen et al., 2000), and even more under 578 Ericaceae, thanks to siderophore substances. Hence, also this process may contribute to the faster 579 podzolization rates found in this study.

580

### 581 **5. Conclusions**

582

In this study we characterize the main pedogenetic processes occurring in recently deglaciated areasunder different vegetation covers. We furthermore provide evidence of fast rates of podzolization

- under a subalpine larch-Rhododendron forest, previously undocumented for the Alps: bleached E
- 586 horizons are visible in 60 years old soils, and diagnostic albic horizons appear in 120 years. In this
- 587 time span the cheluviation of organo-metal compounds in the underlying BC horizons was not
- 588 sufficient to create diagnostic spodic features, but 300-500 years seemed enough for the formation
- 589 of a diagnostic spodic Bs horizon and, consequently, of a "real" Podzol.
- 590 We suggest this relatively fast rate of podzolization be due to the specific plant community
- 591 succession rather than to the climatic conditions in the study area, characterized by cold
- 592 temperatures and significant but not exceptional precipitation amounts. The appearance of a larch-
- 593 Rhododendron forest could significantly influence the soil organic matter characteristics, driving
- the soil development and the onset of soil forming processes in this proglacial area.
- 595 596

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- 605

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815 816	<b>Figure captions</b> Fig. 1: the Lys proglacial area in the Aosta Valley Region, North-western Italy. The sampling sites
817	and the main phases of glacial retreat since 1821 are also shown.
818 819	Fig. 2: 1961-1990 average daily temperature, monthly rainfall or Snow Water Equivalent (SWE,
820	white columns) and snow on the ground (dark columns) in the Gressoney d'Eyola weather station,
821	only few km far from the studied proglacial area.
822 823	Fig. 3: conceptual diagram of vegetation development in the Lys proglacial area; the plant
824	successions observed below timberline are shown on the left, above timberline (i.e. alpine
825	successions) on the right.
826 827	Fig. 4: soil chronosequences under alpine (a), subalpine grassland (b) and typical subalpine forest-
828	heath (c).
829 830	Fig. 5: comparison between C/N ratio in surface mineral horizons (a); surface pH values (b);
831	cumulative thickness of O horizons (OL+OF+OH, c); ratio between Feo (d) and Fed (e) in
832	subsurface CB, BC, Bs and Bw horizons and in surface A, AE and E ones; ISratio (Spodicity Index
833	ratio between subsurface and surface horizons, $IS_{(B-BC)}/IS_{(E-AE)}$ , f) in grassland and forest-shrub soils
834	(p-value $< 0.05$ ). In a), the forest soils low-values outlier is caused by the presence of N-fixing
835	Alnus viridis (S8)
836 837	Fig. 6: Fed depth trends in the LIA sites analyzed in the Lys proglacial area. Different depth trends
838	are visible for the soils under different primary successions.
839 840	Fig. 7: (a), chronofunctions of TOC concentration in surface A, AE and E horizons under grassland
841	(empty circles, continuous line) and forest-shrub vegetation (dashed line, filled squares); (b),
842	surface TOC concentration (%) in subalpine forest-shrub soils and grassland; (c), TOC
843	concentration increase with time in grassland (filled squares, continuous line) and in forest soils
844	(dashed line, empty circles); (d), almost significantly different (p-value = 0.08) TOC concentration
845	in forest-shrub compared to grassland subsoil (CB, BC, BCs, Bs and Bw horizons).
846 847	Fig. 8: (a) descriptive chronofunctions of the ISratio in forest (straight line) and grassland soils
848	(dashed line); . (b) chronofunction of IS increase in subsurface forest soil horizons.
849	
850	Fig. 9: activity ratio in surface and subsurface horizons of grassland and forest soils (p-value <
851	0.05).

	Phytoclimatic belt	Year of moraine deposition	Altitude	Aspect	Slope	Land cover, total plant cover (%)
P1	Alpine/subalpine	2000	2385	/	5°	Pioneer veg. (5)
A2	Alpine	1945	2420	140°	15°	Grassland (25)
A3	Alpine	1945	2415	180°	5°	Grassland (80)
A4	Alpine	1820	2460	140°	20°	Grassland (100)
A5	Alpine	1755	2480	280°	30°	Grassland / dwarf shrubs (100)
A6	Alpine	ca. 11500 BP	2510	220°	30°	Grassland (100)
SG2	Subalpine	1945	2180	200°	5°	Grazed grassland (100)
S3	Subalpine	1945	2180	220°	5°	Larch-rhododendron Forest (100)
S4	Subalpine	1920	2155	340°	2°	Larch-rhododendron Forest (100)
SG5	Subalpine	1920	2155	160°	2°	Grazed grassland (80)
S6	Subalpine	1880	2105	180°	2°	Larch-rhododendron Forest (100)
SG7	Subalpine	1880	2138	160°	2°	Grazed grassland (100)
S8	Subalpine	1821	2005	20°	30°	Larch-rhododendron Forest (100)
<b>S</b> 9	Subalpine	1821	2005	0°	30°	Larch-rhododendron Forest (100)
S10	Subalpine	ca. 11500 BP	1995	90°	20°	Larch-rhododendron Forest (100)

Table 1: Main environmental properties of the study sites; phytoclimatic level (alpine/subalpine), years of deposition of the parent material, altitude, aspect, slope steepness, land cover type and total plant cover (%) are shown.

Table 2: Alpine primary succession; increasing age of the site is from left to right; the % cover of each species (based on visual estimation) is shown in each site. The differentiation between early successional, mid successional, late successional and ubiquitous species is based on Pignatti (1992) and on the Landolt indices (Landolt, 1977): on acidic parent materials, basophilous species should be restricted on weakly developed, initial soils, not yet acidified and desaturated.

	P1	A2	A3	A4	A5	A6
Early successional						
Cerastium uniflorum	1	3	0	0	0	0
Epilobium fleischeri	1	5	2	0	0	0
Linaria alpina	1	1				
Oxyria digyna	1	0	0	0	0	0
Saxifraga oppositifolia	0	0	1	0	0	0
Trisetum distichophyllum	1	3	0	0	0	0
Mid-successional						
Achillea nana	0	5	0	5	0	0
Agrostis schraderiana	1	0	0	0	0	0
Anthyllis vulneraria subsp valesiaca	0	2	2	0	0	0
Aster alpinus	0	1	1	0	0	0
Campanula cochleariifolia	0	0	0	1	0	0
Carex norvegica	0	0	1	0	0	0
Carex ornithopodioides	0	0	1	0	0	0
Erigeron alpinus	0	1	1	0	0	0
Gentiana nivalis	0	0	1	0	0	0
Luzula spicata	0	0	2	3	0	0
Myosotis alpestris	0	0	10	10	0	0
Saxifraga exarata	1	0	2	0	0	0
Saxifraga paniculata	0	0	2	0	0	0
Silene excapa	1	0	5	1	0	0
Silene rupestris	0	0	0	1	0	0
Trifolium pallescens	0	30	10	5	1	0
Late successional						
Antennaria dioica	0	0	0	0	1	0
Anthoxanthum alpinum	0	0	1	0	0	0
Bellis perennis	0	0	0	0	1	0
Botrychium lunaria	0	0	0	1	0	0
Carex curvula	0	0	20	5	5	0
Carex sempervirens	0	0	0	0	0	5
Coeloglossum viride	0	0	0	0	1	0
Euphrasia rohoskoviana	0	0	3	0	0	0
Festuca halleri	0	1	3	5	2	2

Festuca varia	0	1	10	20	0	50
Galium anysophyllon	0	0	0	5	0	0
Gentiana acaulis subsp. Koch	0	0	0	1	0	1
Geum montanum	0	0	0	5	0	10
Hieracium piloselloides	0	0	0	0	1	0
Homogyne alpina	0	0	0	0	1	0
Hupertia selago	0	0	0	1	1	0
Juncus trifidus	0	0	10	0	0	0
Juniperus communis	0	0	5	0	0	0
Loiseleuria procumbens	0	0	0	0	40	0
Lotus corniculatus subsp alpinus	0	0	5	3	0	0
Luzula alpinopilosa	0	0	0	0	2	1
Luzula lutea	0	0	0	0	1	0
Minuartia recurva	0	5	1	0	0	0
Nardus stricta	0	0	0	0	0	5
Pedicularis kerneri	0	0	0	1	1	0
Pedicularis verticillata	0	0	0	1	0	0
Poa nemoralis	0	0	2	0	0	0
Pulsatilla alpina	0	0	0	1	0	1
Rhinanthus alectorolophus	0	0	1	0	0	0
Soldanella verna	0	0	0	0	0	10
Trichophorum caespitosum	1	0	0	0	3	15
Trifolium alpinum	0	0	5	1	5	10
Vaccinium uliginosum subsp gaulterioider	0	0	20	0	60	0
Vaccinium uliginosum subsp gaulterioider Valeriana celtica	0 0	0 0	20 0	0 0	60 2	0 0
Vaccinium uliginosum subsp gaulterioider Valeriana celtica Veronica aphylla	0 0 0	0 0 0	20 0 1	0 0 0	60 2 0	0 0 2
Vaccinium uliginosum subsp gaulterioider Valeriana celtica Veronica aphylla	0 0 0	0 0 0	20 0 1	0 0 0	60 2 0	0 0 2
Vaccinium uliginosum subsp gaulterioider Valeriana celtica Veronica aphylla Ubiquitous	0 0 0	0 0 0	20 0 1	0 0 0	60 2 0	0 0 2
Vaccinium uliginosum subsp gaulterioider Valeriana celtica Veronica aphylla Ubiquitous Agrostis rupestris	0 0 0	0 0 0	20 0 1 5	0 0 0	60 2 0	0 0 2 2
Vaccinium uliginosum subsp gaulterioider Valeriana celtica Veronica aphylla Ubiquitous Agrostis rupestris Bartsia alpina	0 0 0 0	0 0 0 0	20 0 1 5 1	0 0 0 0	60 2 0 1 2	0 0 2 2 2 0
Vaccinium uliginosum subsp gaulterioider Valeriana celtica Veronica aphylla Ubiquitous Agrostis rupestris Bartsia alpina Cerastium cerastioides	0 0 0 0 0 0	0 0 0 0 0 0	20 0 1 5 1 1	0 0 0 0 0 0	60 2 0 1 2 0	0 0 2 2 2 0 10
Vaccinium uliginosum subsp gaulterioider Valeriana celtica Veronica aphylla Ubiquitous Agrostis rupestris Bartsia alpina Cerastium cerastioides Cirsium spinosissimum	0 0 0 0 0 0 0	0 0 0 0 0 0 0	20 0 1 5 1 1 1	0 0 0 0 0 0 1	60 2 0 1 2 0 0	0 0 2 2 0 10 1
Vaccinium uliginosum subsp gaulterioider Valeriana celtica Veronica aphylla Ubiquitous Agrostis rupestris Bartsia alpina Cerastium cerastioides Cirsium spinosissimum Festuca rubra	0 0 0 0 0 0 0 0	0 0 0 0 0 0 0 1	20 0 1 5 1 1 1 0	0 0 0 0 0 1 0	60 2 0 1 2 0 0 0 0	0 2 2 0 10 1 0
Vaccinium uliginosum subsp gaulterioider Valeriana celtica Veronica aphylla Ubiquitous Agrostis rupestris Bartsia alpina Cerastium cerastioides Cirsium spinosissimum Festuca rubra Festuca violacea	0 0 0 0 0 0 0 0 0	0 0 0 0 0 0 1 0	20 0 1 5 1 1 1 0 5	0 0 0 0 0 0 1 0 0	60 2 0 1 2 0 0 0 0 0	0 0 2 2 0 10 1 0 0
Vaccinium uliginosum subsp gaulterioider Valeriana celtica Veronica aphylla Ubiquitous Agrostis rupestris Bartsia alpina Cerastium cerastioides Cirsium spinosissimum Festuca rubra Festuca violacea Leontodon helveticus	0 0 0 0 0 0 0 0 0 0 0	0 0 0 0 0 0 1 0 0	20 0 1 5 1 1 1 0 5 1	0 0 0 0 0 1 0 0 2	60 2 0 1 2 0 0 0 0 0 0 0	0 0 2 2 0 10 1 0 0 0
Vaccinium uliginosum subsp gaulterioider Valeriana celtica Veronica aphylla Ubiquitous Agrostis rupestris Bartsia alpina Cerastium cerastioides Cirsium spinosissimum Festuca rubra Festuca violacea Leontodon helveticus Leucanthemopsis alpina	0 0 0 0 0 0 0 0 0 0 0 0	0 0 0 0 0 0 0 1 0 0 0	20 0 1 5 1 1 1 0 5 1 0	0 0 0 0 0 0 1 0 0 2 5	60 2 0 1 2 0 0 0 0 0 0 0 0 0	0 0 2 2 0 10 1 0 0 0 0 0
Vaccinium uliginosum subsp gaulterioider Valeriana celtica Veronica aphylla Ubiquitous Agrostis rupestris Bartsia alpina Cerastium cerastioides Cirsium spinosissimum Festuca rubra Festuca rubra Eestuca violacea Leontodon helveticus Leucanthemopsis alpina Minuartia sedoides	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	0 0 0 0 0 0 1 0 0 0 0 0	20 0 1 5 1 1 1 0 5 1 0 2	0 0 0 0 0 0 1 0 0 2 5 0	60 2 0 1 2 0 0 0 0 0 0 0 0 0 0	0 0 2 0 10 1 0 0 0 0 0
Vaccinium uliginosum subsp gaulterioider Valeriana celtica Veronica aphylla Ubiquitous Agrostis rupestris Bartsia alpina Cerastium cerastioides Cirsium spinosissimum Festuca rubra Festuca rubra Eestuca violacea Leontodon helveticus Leucanthemopsis alpina Minuartia sedoides Poa alpina	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	0 0 0 0 0 0 0 1 0 0 0 0 0 0 2	20 0 1 5 1 1 1 0 5 1 0 2 5	0 0 0 0 0 0 1 0 0 2 5 0 5	60 2 0 1 2 0 0 0 0 0 0 0 0 0 0 0 2	0 0 2 0 10 1 0 0 0 0 0 0 0 0
Vaccinium uliginosum subsp gaulterioider Valeriana celtica Veronica aphylla Ubiquitous Agrostis rupestris Bartsia alpina Cerastium cerastioides Cirsium spinosissimum Festuca rubra Festuca rubra Festuca violacea Leontodon helveticus Leucanthemopsis alpina Minuartia sedoides Poa alpina Polygonum viviparum	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	0 0 0 0 0 0 0 1 0 0 0 0 0 2 0	20 0 1 5 1 1 1 0 5 1 0 2 5 2	0 0 0 0 0 0 1 0 0 2 5 0 5 0	60 2 0 1 2 0 0 0 0 0 0 0 0 0 0 0 0 0 2 5	0 0 2 0 10 1 0 0 0 0 0 0 0 0 0
Vaccinium uliginosum subsp gaulterioider Valeriana celtica Veronica aphylla Ubiquitous Agrostis rupestris Bartsia alpina Cerastium cerastioides Cirsium spinosissimum Festuca rubra Festuca rubra Festuca violacea Leontodon helveticus Leucanthemopsis alpina Minuartia sedoides Poa alpina Polygonum viviparum Potentilla frigida	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	0 0 0 0 0 0 0 1 0 0 0 0 0 2 0 0	20 0 1 5 1 1 0 5 1 0 2 5 2 0	0 0 0 0 0 0 1 0 0 2 5 0 5 0 5 0 3	60 2 0 1 2 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	0 0 2 0 10 1 0 0 0 0 0 0 0 0 0 0 0 0
Vaccinium uliginosum subsp gaulterioider Valeriana celtica Veronica aphylla Ubiquitous Agrostis rupestris Bartsia alpina Cerastium cerastioides Cirsium spinosissimum Festuca rubra Festuca rubra Festuca violacea Leontodon helveticus Leucanthemopsis alpina Minuartia sedoides Poa alpina Polygonum viviparum Potentilla frigida Primula hirsuta	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	0 0 0 0 0 0 0 1 0 0 0 0 2 0 0 0 0	20 0 1 5 1 1 1 0 5 1 0 2 5 2 0 0	0 0 0 0 0 0 1 0 0 2 5 0 5 0 5 0 3 0	60 2 0 1 2 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	0 0 2 0 10 1 0 0 0 0 0 0 0 0 0 0 0 0 0 0
Vaccinium uliginosum subsp gaulterioider Valeriana celtica Veronica aphylla Ubiquitous Agrostis rupestris Bartsia alpina Cerastium cerastioides Cirsium spinosissimum Festuca rubra Festuca rubra Festuca violacea Leontodon helveticus Leucanthemopsis alpina Minuartia sedoides Poa alpina Polygonum viviparum Potentilla frigida Primula hirsuta Rhododendron ferrugineum	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	20 0 1 5 1 1 1 0 5 1 0 2 5 2 0 0 0 0	0 0 0 0 0 0 1 0 0 2 5 0 2 5 0 5 0 3 0 0 0	60 2 0 1 2 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	0 0 2 0 10 1 0 0 0 0 0 0 0 0 0 0 0 0 0 0
Vaccinium uliginosum subsp gaulterioider Valeriana celtica Veronica aphylla Ubiquitous Agrostis rupestris Bartsia alpina Cerastium cerastioides Cirsium spinosissimum Festuca rubra Festuca rubra Festuca violacea Leontodon helveticus Leucanthemopsis alpina Minuartia sedoides Poa alpina Polygonum viviparum Potentilla frigida Primula hirsuta Rhododendron ferrugineum Salix helvetica	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	0 0 0 0 0 0 0 1 0 0 0 0 0 0 0 0 0 0 0 0	20 0 1 5 1 1 1 0 5 1 0 2 5 2 0 0 0 0 5	0 0 0 0 0 0 1 0 0 2 5 0 2 5 0 5 0 5 0 3 0 0 0 0 0	60 2 0 1 2 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	0 0 2 0 10 1 0 0 0 0 0 0 0 0 0 0 0 0 0 0
Vaccinium uliginosum subsp gaulterioider Valeriana celtica Veronica aphylla Ubiquitous Agrostis rupestris Bartsia alpina Cerastium cerastioides Cirsium spinosissimum Festuca rubra Festuca rubra Festuca violacea Leontodon helveticus Leucanthemopsis alpina Minuartia sedoides Poa alpina Polygonum viviparum Potentilla frigida Primula hirsuta Rhododendron ferrugineum Salix helvetica Saxifraga moschata	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	20 0 1 5 1 1 1 0 5 1 0 2 5 2 0 0 0 0 5 1	0 0 0 0 0 0 1 0 0 2 5 0 5 0 5 0 5 0 5 0 3 0 0 0 0 0 0 0	60 2 0 1 2 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	0 0 2 0 10 1 0 0 0 0 0 0 0 0 0 0 0 0 0 0

Sempervivum montanum	0	3	3	2	0	0
Veronica bellidioides	0	0	1	0	0	0
Veronica fruticans	0	10	0	0	0	0

Table 3: Subalpine primary succession; increasing age of the site is from left to right. the % cover of each species (based on visual estimation) is shown in each site. The differentiation between early-mid successional species, grassland and ubiquitous species and subalpine forest (climax) species was based on Burga et al. 2010 and on Pignatti (1992).

	P1	SG2	\$3	S4	SG5	S6	SG7	S8	<b>S</b> 9	S10
Early-mid successional										
Achillea nana	0	1	2	0	1	0	0	0	0	0
Cerastium uniflorum	1	0	0	0	0	0	0	0	0	0
Epilobium fleischeri	1	1	1	0	0	0	0	0	0	0
Linaria alpina	1	0	0	0	0	0	0	0	0	0
Oxyria digyna	1	0	0	0	0	0	0	0	0	0
Poa nemoralis	0	10	0	0	2	0	0	0	0	0
Rumex scutatus	0	0	1	0	0	0	0	0	0	0
Salix appendiculata	0	20	5	0	10	0	0	0	0	0
Salix helvetica	0	5	0	0	10	0	0	0	0	0
Saxifraga exarata	1	0	0	0	0	0	0	0	0	0
Silene excapa	1	0	0	0	0	0	0	0	0	0
Trisetum distichophyllum	1	0	1	0	0	0	0	0	0	0
Grassland and ubiquitous species (selection)										
Agrostis schraderiana	1	5	5	0	0	0	10	1	0	0
Anthoxanthum alpinum	0	1	0	0	1	0	0	0	10	0
Anthyllis vulneraria subsp valesiaca	0	0	2	0	0	0	0	0	5	0
Astragalus penduliflorus	0	0	0	0	0	0	0	0	5	0
Calluna vulgaris	0	0	0	0	0	0	0	0	5	0
Campanula barbata	0	0	0	0	0	0	0	0	1	0
Carduus defloratus	0	0	0	0	0	0	0	0	1	0
Carex sempervirens	0	0	2	0	0	0	0	0	2	0
Carlina acaulis	0	0	1	0	0	0	0	0	0	0
Cerastium arvense	0	3	0	0	0	0	0	0	1	0

Cirsium spinosissimum	0	0	0	0	0	0	0	0	1	0
Erigeron alpinus	0	0	0	0	0	0	0	0	1	0
Festuca varia	0	20	15	0	10	0	0	0	30	0
Festuca violacea	0	5	0	0	0	0	0	0	15	0
Galium album	0	10	0	0	0	0	0	0	0	0
Galium anisophyllon	0	5	0	0	0	0	0	0	0	0
Geum montanum	0	1	0	0	0	0	0	1	5	0
Helianthemum nummularium subsp. grandiflorum	0	0	0	0	0	0	0	0	1	0
Hieracium pilosella	0	0	0	0	0	0	0	0	5	0
Hieracium prenanthes	0	0	0	0	0	0	0	1	0	0
Knautia arvensis	0	0	0	0	0	0	0	0	1	0
Leucanthemum vulgaris	0	0	0	0	0	0	0	0	1	0
Lotus corniculatus aggr.	0	0	5	0	1	0	0	1	5	0
Nardus stricta	0	0	0	0	5	20	0	0	20	0
Nigritella nigra	0	0	0	0	0	0	0	0	1	0
Phyteuma orbicularis	0	1	0	0	0	0	0	0	0	0
Plantago alpina	0	0	0	0	0	0	0	0	1	0
Poa supina	0	2	0	0	0	0	0	0	0	0
Polygonum viviparum	0	1	0	0	0	0	0	1	0	0
Rhinanthus alectorolophus	0	0	5	0	0	0	0	0	5	0
Schoenus nigricans	0	2	0	0	0	0	0	0	0	0
Sempervivum montanum	0	0	2	0	0	0	0	0	0	0
Solidago virgaurea subsp minor	0	0	0	0	0	0	0	1	0	0
Thymus serpyllum	0	0	0	0	0	0	0	0	5	0
Trichophorum caespitosum	1	5	0	0	0	0	0	0	3	0
Trifolium pratense aggr.	0	1	10	0	0	0	0	0	3	0
Trollius europaeus	0	0	0	0	0	0	0	1	0	0
Subalnine forest-climax (selection)										
Alnus viridis	0	30	0	0	0	0	30	1	0	10
	0	20	0	0	0	0	20	-	0	10

Astrantia minor	0	0	0	0	0	0	0	2	0	0
Avenella flexuosa	0	15	0	0	0	10	50	5	0	5
Calamagrostis villosa	0	5	0	0	3	0	0	0	0	10
Dactylorhiza maculata subsp fuchsii	0	0	0	0	0	0	0	1	0	0
Hieracium murorum aggr.	0	0	0	0	0	0	0	3	0	0
Homogyne alpina	0	0	0	0	0	0	0	2	0	5
Juniperus communis	0	0	0	0	0	10	0	0	0	5
Larix decidua	0	40	30	0	60	50	80	80	0	60
Orthilia secunda	0	3	0	0	0	0	0	0	0	0
Peucedanum ostruthium	0	0	0	0	0	0	0	5	0	0
Phyteuma betonicifolium	0	1	0	0	0	0	0	0	1	0
Rhododendron ferrugineum	0	0	30	0	80	60	50	60	0	70
Vaccinium myrtillus	0	0	0	0	0	40	50	30	0	30
Vaccinium uliginosum subsp. gaulterioides	0	0	20	0	0	0	0	0	0	20
Vaccinium vitis-idaea	0	0	0	0	2	20	0	0	0	20
Viola biflora	0	0	0	0	0	0	0	1	0	1

Tab. 4: Macromorphological properties of the soils in the alpine chronosequence. Structure: GR = granular; PL = platy; PS = subangular polyhedral; MA = massive; RS = rock structure; AB = absent; M = matted (O horizons). The number indicates the size class of the soil structure: 1 = very fine, 2 = fine, 3 = medium, 4 = coarse; Consistence: LO loose; FR friable; FI firm; VFI very firm; The second number symbolizes the strength of the aggregates: 1 = very weak, 2 = weak, 3 = moderate, 4 = strong, 5 = very strong. Silt caps and their thickness are shown as well (1 = up to 1 mm thick, visible on few rock fragments, and 2 = up to 2 mm thick and visible on many rock fragments).

Profile	Year of deposition	Horizon	Depth (cm)	Munsell Color (moist)	Sand (%)	Clay (%)	Structure (or cementation)	Consistence (moist)	Rock fragments (volume, %)	Siltcaps
P1	2004	C1	0-8	5Y5.5/3	69.1	2.1	PL3/AB	LO	60	
		C2	8-47+	5Y6/1	80.3	0.8	PL3/AB	LO	60	1
A2	c. 1950	AC	0-7	2.5Y4/3	82.3	1.1	AB	LO	50	
		C1	7-16	2.5Y4/4	83.4	1.1	AB	LO	60	1
		C2	16-35+	2.5Y5/3	81.5	1.0	AB	LO	60	2
A3	c. 1950	OL	0-0.5							
		А	0-8	10YR4/3	81.2	1.8	GR2	LO1	50	
		AC	8-25	5Y4/2	81.8	1.5	AB	LO	70	
		С	25-48+	5Y4/1	82.9	1.5	AB	LO	70	2
A4	1821	OF	0-3							
		А	3-14	10YR3/2	78.5	2.1	М	LO2	40	
		AB	14-17	10YR4/4.5	79.2	2.0	SP1	LO2	60	
		BC	17-37	10YR4/3	81.2	1.6	AB	LO	50	
		С	37-45+	2.5Y5/4	80.6	1.4	AB	LO	50	1
A5	c. 1755	OL	0-1							
		OF	1-3	7.5YR2/1						
		A1	3-10	7.5YR2.5/1	79.5	2.6	M, PL3	FR3	20	
		A2	10-22	7.5YR3/2	81.3	1.9	SP2	FR2	20	
		Bw	22-40	10YR5/4	84.6	1.1	SP2	LO1	30	
		С	40-50+	2.5Y5/4	94.2	0.5	AB	LO	30	
A6	c. 11000 BP	OL-OF	0-6							
		А	6-12/16	7.5YR2.5/2	69.2	5.2	GR/GM2	LO2	5	
		BA	12/16-48	7.5YR3/3	73.0	4.7	GR2	LO2	40	
		Bw	48-72	8YR3/6	84.1	3.5	GR2	FR2	30	
		IIBC	72-106+	10YR3/6	82.1	1.7	GR1	FR1	10	

	Horizon	pН	С	Feo	Alo	Fed	Feo/Fed
			%	g/kg	g/kg	g/kg	
P1	C1	6.6	0.04	0.87	0.21	1.87	0.47
	C2	6.8	0.00	0.80	0.18	1.50	0.53
A2	AC	5.9	1.71	1.00	0.22	1.80	0.56
	C1	5.2	0.51	1.20	0.19	1.90	0.63
	C2	5.6	0.17	0.80	0.18	1.40	0.57
A3	А	6.1	1.46	1.35	0.44	3.15	0.44
	AC	6.4	0.33	1.23	0.26	2.60	0.46
	С	6.5	0.07	1.73	0.21	2.15	0.77
A4	0	5.1	28.00				
	А	4.7	1.90	1.80	0.70	3.40	0.53
	AB	5	0.93	1.60	0.70	3.50	0.46
	BC	5	0.50	1.30	0.50	2.80	0.46
	С	5.1	0.31	1.30	0.40	2.80	0.46
A5	A1	4.5	4.14	1.80	0.90	4.00	0.45
	A2	4.9	2.53	3.00	1.60	8.20	0.37
	Bw	5.4	0.55	1.70	1.10	8.90	0.19
A6	OL	4.6	23.24				
	А	4.6	4.83	7.25	2.46	16.20	0.45
	BA	5.4	2.19	7.70	5.68	14.80	0.52
	Bw	5.6	1.45	4.65	4.11	17.10	0.27
	BC	5.7	0.77	4.28	3.58	16.40	0.26

Tab. 5: Chemical properties of the alpine soil chronosequences.

Tab. 6: Macromorphological properties of the soils in the subalpine chronosequence. Structure: GR = granular; PL = platy; PS = subangular polyhedral; MA = massive; RS = rock structure; AB = absent; M = matted (O horizons). The number indicates the size class of the soil structure: 1 = very fine, 2 = fine, 3 = medium, 4 = coarse; Consistence: LO loose; FR friable; FI firm; VFI very firm; The second number symbolizes the strength of the aggregates: 1 = very weak, 2 = weak, 3 = moderate, 4 = strong, 5 = very strong. Silt caps and their thickness are shown as well (1 = up to 1 mm thick, visible on few rock fragments, and 2 = up to 2 mm thick and visible on many rock fragments, 3 = up to 2 mm thick and visible on most rock fragments).

Profile	Year of deposition	Hor.	Depth (cm)	Munsell Color (moist)	Sand (%)	Clay (%)	Structure (or cementation)	Consistence (moist)	Rock fragments (volume, %)	Silt caps
P1	2004	C1	0-8	5Y5.5/3	69.1	2.1	PL3/AB	LO	60	
		C2	8-47+	5Y6/1	80.3	0.8	PL3/AB	LO	60	1
SG2	c. 1950	OL	0-1							
		OH	1-3	7.5YR2/2						
		А	3-6.5	10YR4/2	81.3	1.1	GR1	LO	40	
		CB	6.5-18	10YR5/4	82.3	1.1	GR1	LO	50	
		С	18-40+	2.5Y5/3	86.4	0.8	М	М	50	3
<b>S</b> 3	c. 1950	OL	0-1							
		OF	1-3	7.5YR2/2						
		OH	3-6	7.5YR2/2			М	3		
		А	6-12	2.5Y4/1.5	78.6	1.9	GR1	LO	30	
		CB	12-25+	1Y5/4	81.3	1.1	AB	LO	80	
S4	1921	OL	0-1							
		OF	1-2	5YR2/2						
		OH	2-6	5YR2/2						
		Е	6-9	5YR5/1	78.3	2.1	GR1	LO	40	
		CBs/C	9-37+	10YR5/4, 10YR6/3	81.7	1.6	AB	LO	60	
SG5	1921	OF	0-4	10YR2/1			GR1	LO		
		А	4-17	10YR4/3	80.5	1.7	GR1	LO	60	
		С	17-30+	10YR5/3	81.7	0.8	AB	LO	80	1
S6	c. 1880	OL	0-1							
		OF-OH	1-5	5YR2.5/2			Μ	3		
		Е	5-7.5	10YR6/2	68.7	2.2	GR1	Μ	50	
		BCs	7.5-21	10YR4.5/4	74.5	1.1	PS2	FR1	70	
		С	21-27+	10YR5/3	75.6	0.9	AB	LO	70	
SG7	c. 1880	OH	0-1							
		А	1-3	7.5YR2/1	76.5	3.1	М	3	20	
		Bw	3-20/30	10YR4/5	78.3	2.2	PS2	FR1	70	
		СВ	20/30- 30/38	10YR5/3	80.2	1.1	PS1	LO	50	1
		С	30/38-45	2.5Y5/3	81.6	1.0	AB	Lo	60	3
<b>S</b> 8	1821	OL	0-2							

		OF	2-4	7.5YR2.5/2			M1	LO	
		Е	4-7	10YR5.5/2, 10YR3/1	67.5	2.1	GR1	LO1	50
		BCs	7-21	10YR5/4	81.9	1.0	GR1	LO1	60
		С	21-42	10YR5/3	82.6	0.8	AB (PS1)	LO1	80
S9	1821	OL	0-1						
		OF	1-5	5YR2/2					
		OH	5-10/7	5YR2/2					
		AE	10/7-12	10YR4/3	68.2	1.9	GR0	М	30
		Е	12-18	10YR5/2	66.5	2.1	GR1	М	40
		BCs	18-30	10YR4/4	79.0	2.0	PS2	FR1	40
		С	30-40+	10YR5/3	84.2	0.9	AB	LO	50
S10	c. 11000 BP	OL-OF	0-3						
		OH	3-21/12	7.5YR2/1			GR1	LO	
		AE/OH	21/12- 20/30	7.5YR2/2	59.2	8.6	AB	MA3	0
		Eh	20/30- 34/46	10YR5/3	64.4	7.6	AB/PL2	LO	60
		EBh	34/46-54	7.5YR4/4	66.3	7.5	PL2	LO	60
		Bhs	54-68	5YR4/4	77.8	6.1	PS2	LO2	60
		Bsm1	68-87	5YR4/4	79.2	5.9	Cemented	М	80
		Bsm2	87-120	6YR5/8	81.3	5.1	Cemented	М	80
		CBm	120-135	10YR5/8	83.4	4.9	Strongly cemented	М	80
		C(m)	135-150+	2.5Y5/6	79.6	4.8	Partly cemented	M2/LO	80

	Horizon	pH	С	Feo	Alo	Fed	Feo/Fed	Alo/Feo
			%	%	%	%		
P1	C1	6.4	0.04	0.09	0.20	0.19	0.47	0.22
	C2	6.6	0	0.08	0.18	0.15	0.53	0.25
SG2	OH	5.9	26.58					
	А	5.8	1.5	0.11	0.27	0.19	0.58	0.27
	CB	5.8	0.44	0.14	0.36	0.25	0.56	0.29
	С	5.9	0.15	0.10	0.31	0.21	0.48	0.30
<b>S</b> 3	OL/OF	5.6	26.89					
	А	5	1.92	0.07	0.19	0.18	0.39	0.29
	CB	5.2	0.68	0.11	0.31	0.23	0.48	0.27
S4	OL/OF	5.1	41.2					
	OH	4.9	18.44					
	Е	5.1	1.24	0.07	0.18	0.32	0.22	0.29
	CBs	5.2	0.34	0.14	0.22	0.42	0.33	0.14
SG5	OF	5.8	14.32					
	А	5.5	0.55	0.17	0.44	0.38	0.45	0.24
	С	5.8	0.19	0.12	0.51	0.16	0.75	0.50
<b>S</b> 6	OF-OH	4.4	35.24					
	Е	4.7	1.12	0.07	0.49	0.16	0.44	0.71
	BCs	5.1	0.859	0.17	0.58	0.40	0.43	0.35
	С	5.3	0.51	0.15	0.71	0.35	0.43	0.47
SG7	OH	5.6	11.78					
	А	5.6	8.18	0.15	0.61	0.28	0.54	0.40
	Bw	5	0.74	0.15	0.43	0.27	0.56	0.27
	CB	5.7	0.33	0.13	0.34	0.24	0.54	0.23
	С	5.4	0.14	0.09	0.22	0.18	0.5	0.22
<b>S</b> 8	OL-OF	4.6	16.32					
	Е	5	1.19	0.11	0.47	0.17	0.65	0.45
	BCs	5.1	0.75	0.17	0.80	0.67	0.26	0.47
	С	5.2	0.31	0.15	0.48	0.29	0.52	0.33
<b>S</b> 9	OH/OF	5	18.72					
	Е	4.5	0.8	0.11	0.38	0.24	0.46	0.36
	Bs	4.7	0.76	0.20	0.74	0.41	0.49	0.35
S10	OL-OF	3.6	45.21					
	OH	3.5	38.53					
	AE/OH	3.6	5.4	0.12	0.15	0.38	0.32	1.17
	Е	4.1	1.02	0.09	0.09	0.23	0.41	1.00
	EBh	4.2	1.1	0.45	0.13	0.77	0.58	0.29
	Bs	4.9	2.63	1.13	0.77	4.06	0.28	0.68
	Bsm1	5.3	2.31	0.76	0.82	4.02	0.19	1.08
	Bsm2	5.4	0.86	0.29	0.73	1.56	0.19	2.52
	CBm	5.4	0.3	0.11	053	0.81	0.14	4.82
	C(m)	6	0.11	0.11	0.34	0.44	0.25	3.00

Tab. 7: Chemical properties of the subalpine soils along the Lys forefield chronosequences.

















