1 The added value of biomarker analysis to the genesis of Plaggic Anthrosols; the identification of 2 stable fillings used for the production of plaggic manure.

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10 Abstract.

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11 12 Plaggic Anthrosols are the result of historical forms of land management in cultural landscapes on chemically 13 poor sandy substrates. Application of plaggic manure was responsible for the development of the plaggic 14 horizons of these agricultural soils. Pollen diagrams reflect aspects of the environmental development but the 15 interpretation of the pollen spectra is complicated due to the mix of the aeolian pollen influx of crop species 16 and species in the surroundings, and of pollen occurring in the used stable fillings. Pollen diagrams and 17 radiocarbon dates of plaggic Anthrosols suggested a development period of more than a millennium. Calluna is 18 present in almost all the pollen spectra, indicating the presence of heath in the landscape during the whole 19 period of soil development. Optically stimulated luminescence dating of the plaggic horizon made clear that the deposition of plaggic covers started in the 16th century and accelerated in the 18th century. The stable 20 21 fillings, used for the production of plaggic manure and responsible for the rise of the soil surface, cannot be 22 identified with pollen diagrams alone. Biomarker analyses provide more evidence about the sources of stable 23 fillings. The oldest biomarker spectra of the plaggic horizons of three typical plaggic Anthrosols examined in this 24 study, were dominated by biomarkers of forests species as Quercus and Betula while the spectra of middle part 25 of the plaggic horizons were dominated by biomarkers of stem tissue of crop species as Secale and Avena. Only 26 the youngest spectra of the plaggic horizons were dominated by biomarkers of *Calluna*. This indicates that the 27 use of heath sods as stable filling was most likely introduced very late in the development of the Anthrosols. Before the 19th century the mineral component in plaggic manure cannot be explained by the use of heath 28 29 sods. We conclude that other sources of materials, containing mineral grains must have been responsible for 30 the raise of the plaggic horizon.

32 Key words33

Plaggic Anthrosols, plaggic manure, radiocarbon/luminescence dating, palynology, biomarkers, Netherlands.

1. Introduction.

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Plaggic Anthrosols occur in cultural landscapes, developed on coversands. These chemical poor Late-glacial aeolian sand deposits dominate the surface geology of an extensive area in northwestern Europe. Plaggic Anthrosols are the characteristic soils that developed on ancient arable fields, fertilized with plaggic stable manure. Plaggic Anthrosols have a complex genesis and are valuable records of environmental and agricultural history (van Mourik et al., 2011).

44 In previous palaeopedological studies of such soil records in The Netherlands (van Mourik et al, 2011, 2012, 45 2013a), information was unlocked by application of pollen analysis, radiocarbon (¹⁴C) and Optically Stimulated 46 Luminescence (OSL) dating. Radiocarbon dates of soil organic carbon, extracted from humic horizons from 47 plaggic Anthrosols, suggested the start of sedentary agriculture between 3000 and 2000 BP but are not 48 indicative for the age of the plaggic sediments due to the complexity of soil organic carbon in plaggic sediments 49 (Mook & Streurman, 1983; van Mourik et al., 1995). It was assumed that farmers used organic sods as stable 50 filling, firstly dug on forest soils and later on heaths for the production of stable manure to fertilize the fields. 51 The mineral fraction of the sods was supposed to be responsible for the development of the plaggic horizon 52 and the raise of the land surface. OSL dating applied on guartz grains extracted from plaggic sediments 53 provides more reliable ages of the plaggic sediments. The OSL dates suggested that the rise of the plaggic horizons started in the 16th century and accelerated in the 18th century (Bokhorst et. al., 2005). This is rather 54 55 well in line with historical data, as presented by Spek, (2004, p 965).

56 The use of ectorganic matter from forest soils in the Dutch coversand area, must have been strongly reduced in

57 the 11th-13th century, due to commercial forest clear cuttings as recorded in archived documents (Vera, 2011).

58 These deforestations resulted in a regional extension of sand drifting and the managers of the heaths had to 59 protect their valuable ecotopes against this 'historical environmental catastrophe' (Vera, 2011).

Heaths were already present in the Late Paleolithic landscape (Doorenbosch, 2013) and played a ceremonial
 role in the society of our ancestors. People already had the knowledge to manage the heath as sustainable
 grazing areas for cattle (Doorenbosch, 2013).

63 The use of heath for sheep grazing and other purposes as honey and oil production could continue until the

64 middle of the 18th century (Vera, 2011). In SE-Netherlands sustainable use of the heaths was promoted by

- 65 many management rules and laws (van Mourik, 1978; Veera 2011). Over the course of the 18th century, the
- population growth resulted in an increasing food demand. In the course of the 18th century, the deep stable
 economy was introduced and the booming demand for manure resulted in intensivation of manure production.
- 68 Farmers started with the use of heath sods as (additional) stable filling (Spek, 2004). This caused heath
- 69 degradation and initiated the second extension of sand drifting. The use of sods finished at the end of the 19th
- 70 century after the introduction of chemical fertilizers (Spek, 2004).
- Through the combination of OSL and ¹⁴C dating, historical records and the conventional paleoecological proxy of fossil pollen analysis we have a good impression of the paleoecological environment and the age of such
- 73 deposits. However, it remains problematic to reconstruct the combination of crop residues and various
- 74 materials used by farmers as stable filling to produce the stable manure, together responsible for the rise of 75 the surface of Anthrosols. This is also hindering a detailed interpretation of the agricultural practices and shifts
- 75 the surface of Anthrosols. This is also hindering a detailed interpretation of the agricultural practices and shifts 76 therein related to the plaggic agriculture system, and specifically the timing of the onset of the intensive heath
- 77 sod driven deep stable agriculture with which plaggic Anthrosols are most commonly associated. To address
- this issue, in the present study we expanded our paleoecological toolset with an adapted application of the
- recently developed biomarker approach (Jansen et al., 2010). This biomarker approach consists of a combination of analytical chemical analysis and modelling with the VERHIB model to unravel concentration
- 81 patterns of higher chain length (C_{20} - C_{36}) *n*-alkanes of higher plant origin preserved in a soil or sedimentary 82 archive into the (groups of) species responsible for their production (Jansen et al., 2010). The approach was
- archive into the (groups of) species responsible for their production (Jansen et al., 2010). The approach was
 originally developed to unravel past local vegetation composition. Upon successful application in a tropical
 ecosystem setting (Jansen et al., 2013), its applicability in palaeopedology was explored (van Mourik & Jansen,
- 85 2013b). This pilot application concerned a polycyclic soil sequence in driftsand deposits. It showed that the 86 comparison of pollen and biomarker spectra allowed us to indicate the plant species responsible for carbon 87 sequestration in the humic horizons (van Mourik & Jansen, 2013b). Important conclusion was that biomarker 88 analysis showed promise not only in the reconstruction of past local vegetation composition of a specific site, 89 but also in studies where the emphasis lies not on the vegetation per se, but rather on reconstructing various
- 90 sources of soil organic matter input (van Mourik & Jansen, 2013b).
- 91 Goal of the present study was to further explore the applicability of biomarker analysis as part of a multi-proxy 92 reconstruction aimed at unraveling the sources of stable fillings used for the production of plaggic manure in 93 the context of the historic development of the plaggic agriculture ecosystem. For this, we applied biomarker 94 analysis on three previous investigated plaggic Anthrosol. 95

96 Materials and methods.

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2. Profile selection

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	Fig. 1
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	Fig. 2 a,b,c
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- 102 The distribution area of plaggic Anthrososl in NW-Europe is indicated in fig.1. Pape (1972) published the first 103 map of the distribution of plaggic agriculture in NW-Europe. Bastiaens & van Mourik (1995) found traces of 104 intensivation and extension of this area in Vlaanderen (Belgium) while van Mourik (1999b) also reported 105 plaggic Anthrosols in Schleswig (Germany). Beside this area with 'real' plaggic Anthrosols, Spek (2004, p. 724) 106 summarized information about the occurrence of soils with some evidence of application of plaggic manure in 107 the Atlantic coastal zones of Norway, Denmark, France, Galicia, Scotland and Ireland.
- For this pilot study we selected three previously investigated plaggic Anthrosols in the Netherlands with an undisturbed plaggic horizon: Valenakker, Nabbegat and Posteles (fig.2). Pollen diagrams, ¹⁴C and OSL dates of these profiles were available and previously published separately in various articles. Here we combined these, and re-sampled the plaggic horizons of the profiles for biomarker analysis and new fossil pollen analysis to allow for comparison. Vertical sampling resolution was: Valenakker 20 cm, Nabbegat and Posteles 10 cm.

113 Valenakker (van Mourik et al. 2012) is situated southwest of the city Weert (middle Limburg) on the sport fields 114 of a former college. As a result, during the 20th century the soil has never been ploughed or subjected to land 115 consolidation. This profile has never been affected by roots of *Zea mays*, introduced in The Netherlands in the 116 middle of the 20th century (van Mourik & Horsten, 1995).

117 Nabbegat (van Mourik et al. 2013a) is situated on the Maashorst (eastern North-Brabant). The plaggic deposits 118 were buried by drift sand around 1800 AD. Consequently, the plaggic deposits have perfectly been protected 119 against damage by land consolidation or pollution afterwards (van Mourik et al. 2013a). The site is now 120 vegetated by oak and birch trees. Roots of these trees may have caused input of organic matter by 121 decomposed roots in the upper part of the plaggic horizon (fig.3).

Fig. 3

Posteles (van Mourik et.al, 2011) is situated in Twente (eastern Overijssel). The landowner informed us that during the last three generations this land was never subjected to deep ploughing or land consolidation but since 1960 Zea mays was regular sowed. In contrast to Valenakker and Nabbegat we can expect biomarkers of this deep rooting cultivated plant.

2.1. Pollen analysis.

Pollen diagrams of plaggic Anthrosols provide paleoecological information about plant species, present on site and in the region during the formation of the plaggic horizon. Previous research showed that pollen grains, infiltrated in soils and incorporated in plaggic deposits, are well preserved in the anaerobic and acid microenvironment of excremental aggregates (van Mourik, 1999a, 2001) (fig 4,5).

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138	Fig. 4
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140	Fig. 5
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Samples for pollen extraction were collected in 10 ml tubes in profile pits. For a correct matching of pollen and biomarker spectra of the plaggic deposits, the same samples were treated for both pollen and biomarker extraction and analysis. The pollen extractions were carried out using the tufa extraction method (Moore et al., 1991, p. 50). For the identification of pollen grains, the pollen key of Moore et al. (1991, p. 83-166) was applied. Pollen scores were based on the total pollen sum of arboreal and non-arboreal plant species. For the estimation of the pollen concentrations of the various soil horizons, the exotic marker grain method was applied (Moore et al., 1991, p. 53).

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2.2. ¹⁴C and OSL dating.

152 153 The determination of the age of plaggic deposits is subjected to various complications (Spek, 2004). Pollen 154 stratification is disturbed by bioturbation and ploughing. Besides, the pollen content is a mix of the regular 155 pollen influx and pollen in stable fillings, used for the production of stable manure (van Mourik et al., 2011). The ages of humic horizons of buried Podzols cannot be correctly determined by ¹⁴C dating due to the complex 156 composition of soil organic carbon (van Mourik et al., 1995). During a period of active soil formation, hard 157 158 decomposable organic carbon can accumulate in the Ah horizon, especially in the humin fraction but also in the 159 humic acid fraction. Especially the accumulation of charcoal fragments in the organic aggregates is responsible 160 for overestimation of the ¹⁴C ages (fig.4). During the Early Holocene small amounts of charcoal fragments were 161 released after (natural) forest fires, but the amount increased drastic in the iron time due to the charcoal 162 production for the melting of iron from placic horizons and iron stone (Beukenkamp and Sevink, 2005). The age 163 of the humic acid fraction was considered the best estimate of the moment of fossilization of the Ah horizon 164 after burying by driftsand. The difference between humin and humic acids ages was interpreted as a measure 165 for the period of soil activity and humin accumulation. Later, OSL dating confirmed that radiocarbon dates, not 166 only of the humin fraction but also of the humic acids, overestimate the true ages (Bokhorts et al., 2005).

167 Conventional radiocarbon dating of humin and humic acids showed in presented diagrams, extracted from168 plaggic deposits, was performed in the CIO (Centre for Isotope Research of the University of Groningen).

OSL dates provide reliable information about the moment of fossilization of plaggic material under the rising
 furrow because the quartz grain were perfectly bleached during active ploughing (Bokhorts et al., 2005). OSL
 dating of quartz grains, extracted from plaggic deposits, was performed in the NCL (Netherland Centre for
 Luminesce Dating, Wageningen University).

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2.3. Biomarker analysis.

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2.3.1. The application of the VERHIB model

178 A detailed description of the biomarker approach using the VERHIB method is presented in our previous 179 publications (Jansen et al., 2010; Jansen et al., 2013; Van Mourik & Jansen 2013). Briefly, the basis of the 180 method lies in the unraveling of the preserved concentration patterns of C_{20} - C_{36} *n*-alkanes, which are exclusive 181 to the epicuticular wax layers on leaves and roots of higher plants (Kolattukudy et al., 1976). While such an 182 application in itself is not new (e.g. Pancost et al., 2002; Hughen et al., 2004) the novelty of our approach lies in 183 the application of the VERHIB model that we specifically developed to unravel the mixed n-alkane signal 184 encountered in soil or sedimentary archives (Jansen et al., 2010). The VERHIB model consists of a linear 185 regression model that describes how a certain input of plant derived compounds such as *n*-alkanes over time in 186 a certain archive at a certain location, results in accumulation of these compounds. An inversion of the forward 187 model is used to reconstruct the accumulation encountered with depth into its most likely vegetation origin 188 (Jansen et al., 2010). An important aspect of biomarker analysis using VERHIB is that it is an indirect 189 reconstruction. While the biomarker patterns, in the present study the n-alkanes, are directly measured, the 190 reconstruction into the most likely combination of vegetation biomass input responsible for the observed 191 pattern is inferred by the model. For this, several parameters must be inputted into the model (Jansen et al., 192 2010) the most important of which is the selection of the expected plant species that have been responsible for 193 the input of biomass in the archive in question, and subsequent inclusion of their n-alkane signature in the 194 VERHIB reference base. In the present study, the selection of species to include was based on the (expected) 195 crop history of the sites under study, as well as the anticipated origin of the stable fillings used as manure. An 196 important matter of debate when using *n*-alkane patterns to reconstruct past vegetation input is the genotypic 197 plasticity of the *n*-alkane patterns, in particular in relation to prevailing environmental factors such as climate 198 (e.g. Shepherd and Griffiths, 2006). In a previous study focusing on vegetation of relevance for reconstructions 199 in ecosystems in North-Western Europe where plaggic agriculture occurred, we found that while genotypic 200 plasticity related to climatic factors may influence the signal, such influence does not eradicate the different 201 vegetation origins (Kirkels et al., 2013). To limit external influences as much as possible, the vegetation selected 202 for inclusion in the VERHIB reference base was sampled in close vicinity to the three study sites as much as 203 possible. The first group of selected plant species concerned the main sources of stable fillings, used for the 204 manure production: fermented litter from deciduous forest soils (Quercus robur, Betula pendula), grass sods 205 from brook valleys (Molinia caerulea) and heath sods (Calluna vulgaris).

The second group concerned crop species. Close to the educational Field Study Centre Orvelte (Drenthe) is a traditional plaggic field where they continued with the cultivation of traditional crop species. There we sampled *Fagopyrum esculentum, Spergula arvensis, Avena sativa, Secale cereal, Spergula arvensis.* The modern crop species *Zea mays* corn was sampled on the Posteles.

The concentration patterns of the *n*-alkanes with carbon numbers 20-36 in the selected vegetation samples and in the soil samples were subsequently used as input for the VERHIB model (see 2.3.2 for a description of

212 the extraction and analysis of the biomarkers).

213 A second parameter that must be considered in the application of VERHIB, is input of leaf and root material. 214 VERHIB considers the species specific *n*-alkane patterns in plant roots separately from the patterns in plant 215 leaves, and uses this to deal with the input of young root material at depth (Jansen et al., 2010). A first 216 selection criterion here concerns whether or not leaf and root material can be expected to have entered the 217 soil at all. For the deciduous forest soil material potentially used as stable fillings (Quercus robur, Betula 218 pendula), exclusively leaf derived biomass input is expected as the trees did not grow on-site. In contrast, for 219 the crop species Zea Mays and Spergula Arvensis only root material is expected to have entered the soil in 220 appreciable amounts as the leaf material is mostly removed during harvest. For the other species considered, 221 both leaf and root material must be taken into account. A selection of root and/or leaf derived n-alkane 222 patterns to be considered in the VERHIB reference base was made in accordance with the previous. With 223 respect to the ratio of input of leaf vs. root biomass as required by the model, no exact information is available 224 for the soil profile under study. Therefore, for those species where both leaf and root material is considered to 225 have possibly entered the soil, in line with the exploratory nature of the present study, we applied an assumed

leaf/root biomass input ratio of 1.0 and assumed that while input of leaf material always occurred at the top of the soil profile, root input also occurred with depth. Since our pilot study in polycyclic driftsand deposits showed that VERHIB was unable to filter out root input sufficiently (Van Mourik & Jansen, 2013), when interpreting the occurrence of a certain species with depth in the profiles under study as modelled by VERHIB, the possibility of young root input being responsible for the signal was explicitly taken into account.

Figure 6 shows a flow diagram that illustrates the functioning of the VERHIB modelling as well as the selectionof parameters and reference base species as described above.

Fig. 6

2.3.2. Extraction and analysis of the biomarkers.

237 Approximately 0.1 g of each of the freeze-dried and ground vegetation and soil samples was extracted by 238 accelerated Solvent Extraction (ASE) using a Dionex 200 ASE extractor. The extraction temperature was 75°C 239 and the extraction pressure 17 × 106 Pa, employing a heating phase of 5 min and a static extraction time of 20 240 min. Dichloromethane/methanol (DCM/MeOH) (93:7 v/v) was used as the extractant (Jansen et al., 2006). The 241 extracts were subsequently fractionated into three fractions containing the *n*-alkanes, the esters and the 242 combination of alcohols and fatty acids respectively. For this, a silica column consisting of extracted cotton 243 wool and silica gel was used, followed by elution with hexane, hexane/DCM (4:1) and DCM/Methanol (9:1) 244 respectively. Separation of the *n*-alkanes took place by on-column injection of 1.0 μ l of the first fraction on a 30 245 m Rtx-5Sil MS column (Restek) with an internal diameter of 0.25 mm and film thickness of 0.1 µm, using He as a 246 carrier gas. Temperature programming was: 50°C (hold 2 min); 40°C/min to 80°C (hold 2 min); 20°C/min to 247 130°C; 4°C/min to 350°C (hold 10 min). Subsequent MS detection in full scan mode used a mass-to-charge 248 ratio (m/z) of 50-650 with a cycle time of 0.65 s and followed electron impact ionization (70 eV). The n-alkanes 249 were identified from the total ion current (TIC) signal by their mass spectra (dominant fragment ion 250 represented by m/z = 57) and retention times and quantified using a deuterated internal standard $(d_{42}-n-C_{20})$ 251 alkane (Jansen et al. 2010) as well as a conventional external *n*-alkane standard.

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Fig.7

Figure 7 presents the n-alkane biomarker distribution in the leaves and/or roots of the species, inserted in the reference base. The results show the odd-over-even chain-length predominance typical of higher plants (Kolattukudy et al., 1976). The observed variation in patterns and concentrations is in line with the variation found in other species in previous work (e.g. Jansen et al., 2006).

3. The vertical distribution of biomarkers and pollen in the analysed profiles.

3.1. Profile Valenakker

263 Profile Valenakker is a plaggic Anthrosol (Aan), overlying a ploughed umbric Podzol (2ABp, 2Bs). The pollen 264 diagram (fig.3) and the absolute dates (table 1) reflects a soil development of \approx 1400 year.

The post sedimentary pollen spectra in the 2BS show percentages of tree species as *Corylus* and *Quercus* of the Middle Subatlantic. The presence of Poaceae, Cyperaceae, *Rumex* and Ranunculaceae reflects a period of pasture. The high scores of Cerealia in ploughed 2ABp and even the 2B indicate a form of sedentary agriculture before the start of plaggic agriculture. ¹⁴C dating indicate a carbon age of the base (60 cm) of the Aan horizon of \approx 600 AD. The OSL age of the lower part of the plaggic horizon is 800-900 year younger, \approx 1560 AD.

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- Fig.8. 271 Table 1. 272 Fig.9. 273

274 Micromorphological observations (fig.5ab) of the plaggic deposits show the complexity of soil organic matter. 275 There are various sources of organic carbon as plants roots, tissue of table fillings and sods. Also the composition of pollen spectra is complex, a mix of the regular pollen influx of plants on the fields and in thesurrounding infiltrating into the soil and pollen, and pollen present in various stable fillings.

278 In previous studies the origin of stable fillings, used in plaggic agriculture, was reconstructed on the base of

pollen diagrams (Spek, 2004; van Mourik et al., 2012a, 2012b). The pollen spectra of the Aan horizon show very

280 low scores of arboreal trees but reasonable scores of Ericaceae and Poaceae. Ericaceae pollen may indicate the 281 use of heath sods, Poaceae pollen the use of grassland sods, the combination of sods from degrading heath and

the rise of the land surface by plaggic manure is caused by the mineral fraction in such sods. However, the rise

of the plaggic horizon of ≈ 60 cm cannot be explained by the use of heath sods if it is true that the use of heath

sods (with a mineral fraction) was introduced in the course of the 18th century when better construction

285 materials enabled the farmers to build deep stables (Vera, 2011). In fact, the sources of stable fillings cannot be 286 satisfactorily detected with pollen diagrams.

The biomarker spectrum of the base is dominated by *Quercus*. Despite the low percentages *Quercus* pollen it is very likely that the farmers used forest litter as stable filling. The middle spectrum is dominated by markers of *Avena* and *Secale*. This points to the use of straw from these crop species as stable filling. Pollen of Cerealia is present in the whole diagram. In the upper spectrum biomarkers of *Calluna* are present together with *Avena* and *Secale*. This points to the use of heath sods as additional stable filling during the last phase in the development of the plaggic horizon.

3.2. Profile Nabbegat

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	Fig. 10.
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	Table 2.
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	Fig. 11.
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Profile Nabbegat is a haplic Arenosol (with Mormoder humus form), overlying a plaggic Anthrosol, overlying a
 ploughed umbric Podzol. The pollen diagram (fig.6.) and the absolute dates (table 2) reflect a soil development
 of ≈ 3000 year.

The post sedimentary pollen spectra of the 3ABp reflect the start of agriculture (increase of Cerealia) on a former heath (decrease of *Ericaceae*) in a surrounding with coppice hedges (*Quercus, Corylus*). Based on radiocarbon dates, the agricultural activities started before \approx 1000 BC, the OSL dates point to deposition of plaggic material after \approx 1500 AD.

The radiocarbon ages indicate that the farmers used organic matter with very few mineral 'contamination' for a long time. The OSL ages indicate that the rise of the plaggic horizon started \approx 1500 AD due to mineral grains as part of the manure. The plaggic horizon developed between 1500 and 1800 AD. Around 1800 AD, short after the introduction of the deep stable economy (Vera, 2011), the plaggic Anthrosol was overblown by driftsand. Apparently, the use of heath sods resulted in heath degradation, sand drifting and acceleration of the rise of the plaggic horizon (van Mourik et al., 2012a). The sand drifting stabilized under planted *Quercus* trees; the roots of these trees reached the buried Anthrosol and may have contributed the scores of biomarkers in the

upperpart of the buried plaggic horizon. The composition of the pollen spectra of the plaggic horizon is ratheruniform, dominated by Ericaceae and Cerealia.

Fig.7. shows the results of biomarker analysis. Biomarkers of *Quercus* were present in all the spectra, dominant in the lower spectra, regular in the other spectra. This points to the use of forest litter as stable filling during

the development of the lower part of the plaggic horizon. The main crop species during this time was *Spergula*.

318 The middle part is dominated by markers of *Avena* and *Secale*, indicating the use of straw. Only in the upper

319 spectrum *Calluna* was found, indicating the use of heath sods during the last phase of the development of the

320 plaggic horizon.

323 **3.3. Profile Posteles**

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	Fig.12.
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	Table 3.
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328 Profile Posteles is a plaggic Anthrosol, overlying a ploughed umbric Podzol. The pollen diagram (fig.8) and the 329 absolute dates (table 3) reflect a soil development of at least 1200 year.

330 The pollen content of the buried ploughed Podzol (2Ap, 2B) is post-sedimentary infiltrated in Late-Glacial 331 coversand by bioturbation and agriculture. Characteristic is the sharp decrease of pollen concentrations with

332 depth, shown by the pollen density curve. The spectra of the 2B horizon already reflect evidence of agriculture 333 (Cerealia) in a deforested landscape (low percentages of Alnus, Quercus, Fagus). The spectra of the 2Ap horizon 334 show increasing percentages of Cerealia.

335 The radiocarbon age of the base of the plaggic deposits (95 cm) is \approx 850 AD, The OSL age \approx 1500 AD. The OSL 336 age of the 2Ap (105 cm) is 2035 \pm 450 BC, \approx 3500 year older than sample 95. In this part of the profile we see 337 the effect of bioturbation on the age of the coversand. Grains from the base of the Aan were transported to 338 the 2Ap and reversed, which explains the large standard deviation of the OSL of sample 105 cm.

339 The actual Ap horizon (the active plough horizon) is palynologically characterized by peak percentages of Cerealia, a 340 slight extension of Pinus (planted on the abandoned heath after 1900 AD) and the appearance of Zea mays 341 (introduced in Dutch agriculture after 1950 AD). Pollen of Cerealia, Ericaceae and Poaceae were found in all the 342 spectra of the Aan.

343 The lowest spectrum (80) is dominated by the crop species Spergula and the score of Quercus indicates the use 344 of forest litter during the development of this part of the Aan.

345 The spectra 10, 20, 40, 60 are dominated by biomarkers from roots of Zea mays. This crop species was 346 introduced around 1950 AD, but the markers of the decomposed Zea roots seem to suppress all the others 347 (this was not the case in the profiles Valenakker and Nabbegat). Spectrum 50 is dominated by Avena and 348 Secale, spectrum 30 by Zea and Secale and spectrum 0 by Zea and Calluna. Again the use of heath sods seems 349 restricted to the youngest part of the plaggic horizon.

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352 4. Discussion

354 Pollen diagrams of plaggic Anthrosols provide valuable paleoecological information to reconstruct the soil 355 dynamics during the plaggic agriculture. However, interpretation of pollen diagrams is complicated. Pollen 356 grains, extracted from plaggic deposits, may originate from two sources (van Mourik et al., 2011). The first 357 source concerns the regional pollen influx from flowering species and local flowering crop species. Pollen grains 358 precipitate on the soil surface and may infiltrate into the Anthrosols by ploughing and bioturbation. This pollen 359 influx will be mixed with the pollen content of materials, used as stable filling to produce manure.

360 Pollen will be preserved in plaggic deposits in the anaerobic and acid micro environment of humic aggregates, 361 produced by worms and micro arthropods (van Mourik, 1999b, 2001). In general it is not possible to make a 362 clear separation between pollen grains originating from the regular pollen influx or from materials as sods. 363 Therefore, the identification of the various sources of stall fillings cannot be based on pollen analysis alone. 364 Additional information, acquired by biomarker analysis proved very useful for this purpose.

365 In the pollen diagrams Fagopyrum is found in almost all the spectra of the plaggic deposits and in Valenakker 366 and Nabbegat even in the top spectra of the buried ploughed Podzol, probably as result of pollen infiltration. 367 Fagopyrum as crop species on sandy soils was introduced after 1350 AD (Leenders, 1996). Based on this 368 palynological time marker, plaggic deposition started around 1350 AD.

369 The radiocarbon ages of plaggic deposits are much older. This is caused by (1) older organic carbon, present in 370 the applied stable fillings (as forest litter) for the manure production and (2) accumulation of hardly 371 decomposable organic carbon during active soil formation. Consequently the radiocarbon dates overestimate the ages of the plaggic sediments, but approach the age of the introduction of agricultural soil management 372

373 (van Mourik et al., 1995, 2011, 2012a, 2012b). Manuring of infertile soils came already in use in the Bronze Age

- and also the Celtic fields are an example of a prehistorical agricultural system based on manure management
 (Spek, 2004).
- The mineral component of stable manure, applied on the fields, was responsible for the thickening of the plaggic horizon. Ploughing of the furrow will bleach the OSL signal of the mineral grains until the moment that the grains are no longer part of the active soil furrow. For that reason, OSL dating of the plaggic horizon provide
- 379 reliable ages of the plaggic deposits (Bockhorst et al., 2005). The OSL dates of the profiles Valenakker,
 380 Nabbegat and Posteles indicate a start of the thickening ≈ 1550 AD.
- 381 It was not possible to determine the sources of stable fillings palynologically. Possible stable fillings were forest 382 litter, sods from moist grass lands and heats sods. But in almost all spectra of the pollen diagrams Ericaceae,
- litter, sods from moist grass lands and heats sods. But in almost all spectra of the pollen diagrams Ericaceae,
 Poaceae and arboreal pollen occur. Biomarkers extracted from plaggic deposits, originate from two sources.
- The first source concerns biomarkers from decomposed roots of crop species, the second source of organic material as straw and sods, used as stable filling for manure production.
- In the three diagrams we find *Quercus* as dominant marker in the lowest part of the Aan-horizon, indicating the use of forest litter. In Nabbegat, Quercus markers can also originate from roots of the planted *Quercus* forest after the stabilization of the sand drifting. This is not the case on Valenakker and Posteles. The middle part of the Aan-horizon is dominated by markers of *Avena* and *Secale*, indicating the use of straw as stable filling.
- 390 Only in the top of the Aan-horizon markers of Calluna are present, indicating the use of heath sods as stable
- filling. Based on the results of the biomarker analysis we can conclude that heaths sods were used as stable
- filling only in the 18th and 19th century. This fits with the observations about the use of heaths in historical
 archives Vera (2011).
- 393 archives Vera (2011).394 So the question rises about heath management before the introduction of the deep stable economy. Some
 - researchers point to careful heath management before the 19th century. In interviews with farmers, born 395 396 before 1950, Burny (1999) collected essential information about historical heaths management in the Belgian Kempen. A historical study of land use in the Campina also indicated carefully maintenance and sustainable use 397 of valuable common fields (de Keyzer, 2014). Before the 19th century, heath sods were never dug on the dry 398 399 Calluna heath, only on the moist Erica heath. These organic sods were not used as stable filling but as fuel for 400 the furnace. Burning of Calluna heaths was the most important management action to rejuvenate the heath. 401 Juvenile heath is food for cows. Sods digging was a bad action due to the resistance and incoherence of these 402 dry sods and also the long recovery period. Mowing of older Calluna shrubs took place. Twigs were used for 403 roofs, burning and brooms. (Burny, 1999). Because of the very low nutrient contribution to the manure of
 - mowed *Calluna*, the farmers preferred the use of twigs of broom (*Genista*). When in the course of the 18th the
 authority relationships changed and the population growth and the demand for food increased, farmers
 started to intensify their production (Vera, 2011). They needed more manure and started with the deep stable
 economy and the use of *Calluna* heath sods.
 - 408 An important factor may be the presence of pollen and biomarkers content in sheep droppings. According to 409 Simpon et al., (1999) biomarkers survive the congestion process and stay in the manure. But what do sheep 410 consume? Grazing sheep are very selective in collecting food (Oom et al., 2008; Smits & Noordijk, 2013). They 411 prefer grasses (Molinia, Festuca and Corynephorous). Only in years that there is insufficient grass available at 412 the end of the summer, they eat shoots of Calluna, at that time nourishing with high concentrations Ca, Mg and 413 but no P. Pollen extractions from sheep droppings showed that only in droppings, collected during the summer 414 season Calluna pollen is present. During the flowering season of Calluna, the animals consume pollen, 415 precipitated on the grasses. That explains the presence of Calluna pollen and the absence of Calluna 416 biomarkers in the lower parts of the plaggic horizons.
 - 417 If it is true that *Calluna* heath sods were dug only in the 18th and 19th century, how can we explain the mineral
 418 component in the plaggic manure, responsible of the rise of the land surface before that time?
 - According to Smits and Noordijk (2013) there are several sources of minerals. Firstly, a small amount of mineral grains will be incorporated in the manure during emptying out the manure of the stable. Secondly, farmers had the knowledge that the addition of sand could improve the fertility of the soil. Not the leached and acid sand from heath sods but not leached sand, dug on sheep walks and in blown out depressions in nearby drift sand landscapes.
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5. Conclusions

- The vertical zoning of biomarkers and pollen in plaggic horizons are different. Palynologically, the plaggic
 horizon is a homogenous, the biomarker diagrams show differentiation.
- We can identify various stable fillings used, based on the vertical distribution of biomarkers.

- The biomarker spectra of the base layer of the plaggic horizon are dominated by biomarkers of deciduous
 trees litter (dominated by *Quercus*), indicating the use of organic matter from the forest floor.
- The biomarker spectra of the middle part of the plaggic deposits are dominated by crop species (*Avena*, *Secale*), indicating the use of straw from these species as stable filling during a relatively long time.
- Only the top spectra of the plaggic horizons are dominated *by Calluna,* indicating that heath sods were
 used as stable filling only during the last phase in the development of the plaggic horizon.
- Profile Posteles shows the impact of the contribution of biomarkers of roots of *Zea mays*, introduced around 1950 AD, suppressing the other species.
- The negligible percentages of *Calluna* in biomarker spectra of plaggic deposits with exception of the top, suggest an overestimating of the use of heath sods in the traditional interpretation of the genesis of plaggic horizons, the dominance of crop species in biomarker spectra of plaggic deposits suggests underestimating of the use of straw as source material for the production of organic stable manure to fertilize ancient arable fields.
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576 Tables (including table captions)

Table 1. ¹⁴ C	Table 1. ¹⁴ C and OSL dates of the plaggic deposits of Valenakker.					
Horizon	Depth (cm)	Calendric ¹⁴ C ages	Calendric ¹⁴ C ages	Calendric OSL ages		
		humin	humic acids			
Aan	20	-	-	1775 ± 20 AD		
Aan	40	771 ± 92 AD	1049 ± 78 AD	1635 ± 30 AD		
Aan	60	595 ± 61 AD	698 ± 54 AD	1565 ± 30 AD		

Table 2. ¹⁴ C and OSL dates of the plaggic deposits of Nabbegat.					
Horizon	Depth (cm)	Calendric ¹⁴ C ages	Calendric ¹⁴ C ages	Calendric OSL ages	
		humin	humic acids		
С	70	-	-	1803 ± 12 AD	
2An	80	428 ± 107 AD	626 ± 45 AD	1770 ± 11 AD	
2An	105	37 ± 133 BC	3 ± 101 AD	-	
2An	130	1182 ± 139 BC	811 ± 101 BC	1676 ± 14AD	
ЗАВр	140	-	1299 ± 78 BC	-	
ЗАВр	150	_	1385 ±72 BC	_	

Table 3. ¹⁴ C and OSL dates of the plaggic deposits of Posteles.					
Horizon	Depth cm	Calendric ¹⁴ C ages	Calendric ¹⁴ C ages	Calendric OSL ages	
		humin	Humic acids		
Aan	45	-	-	1758 ± 14 AD	
Aan	59	-	-	1711 ± 20 AD	
Aan	70	1132 ± 68 AD	1172 ± 51 AD	1651 ± 31 AD	
Aan	82	-	-	1626 ± 20 AD	
Aan	95	884 ± 82 AD	861 ± 85 AD	1517 ± 31 AD	
2ABp	105	_	_	2035 ± 450 BC	

581 Figure Captions

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Fig. 1. The location of sampled profiles Valenakker, Nabbegat and Posteles in the distribution area of plaggic
agriculture.

Fig. 2. The plaggic Anthrosols Valenakker, Nabbegat and Posteles. The location of the OSL samples are indicatedin the white circles (depth in cm); the locations of the profiles are indicated in fig. 1.

Fig. 3. Cross-section of a (living) tree root in the thin section of the 2 Aan of Nabbegat (70-80cm). Characteristic
is the double fringing of the root tissue. Such roots were only found in the upper part of the 2Aan of Nabbegat.
Roots of crop species were not found in the thin sections of the three profiles; they decompose rather fast
compared with tree roots.

Fig. 4. Distribution pattern of organic aggregates in a thin section of the Aan of Valenakker (40-50 cm). In the fabric of the aggregates are charcoal particles visible

Fig. 5. Pollen grains, visible in a welded aggregate of the same thin sections. Pollen grains in thin sections are
 observable as not double fringing, empty spheroidal objects. The palynological characteristics as sculpture and
 aperture are not visible without the chemical treatments during pollen extraction.

601 Fig. 6. Flow diagram of the methodology of biomarker analysis.

Fig. 7. The *n*-alkane biomarker distribution in leaves and/or roots of species sampled, for the reference base of this pilot study.

- 606 Fig. 8. Pollen diagram Valenakker. Pollen density in k.grain/ml.
- 608 Fig. 9. Biomarker diagram Valenakker.
- Fig. 10. Pollen diagram Nabbegat. Log D = pollen density in log k.grain/ml.
- 612 Fig. 11. Biomarker diagram Nabbegat.
- 614 Fig. 12. Pollen diagram Posteles; Pollen density in k.grain/ml.
- 616 Fig. 13. Biomarker diagram Posteles.

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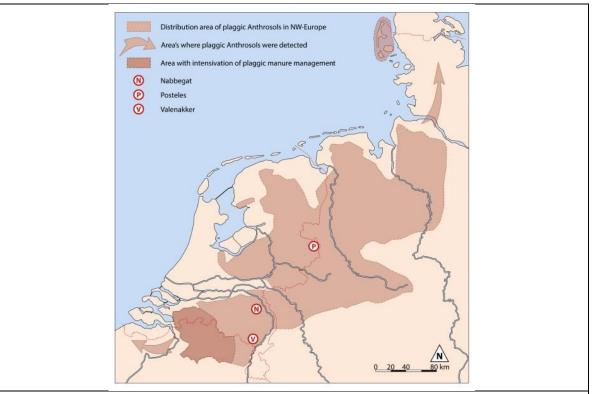


Fig. 1. The location of sampled profiles Valenakker, Nabbegat and Posteles in the distribution area of plaggic agriculture.



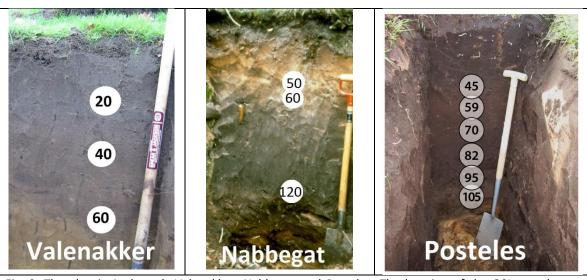


Fig. 2. The plaggic Anthrosols Valenakker, Nabbegat and Posteles. The location of the OSL samples are indicated in the white circles (depth in cm); the locations of the profiles are indicated in fig. 1.



Fig. 3. Cross-section of a tree root in a thin section of the 2 Aan of Nabbegat (70-80cm). Characteristic is the double fringing of the root tissue. Such roots were only found in the upper part of the 2An of Nabbegat. Roots of crop species were not found in thin sections of the three profiles. The decompose rate of such roots is fast.

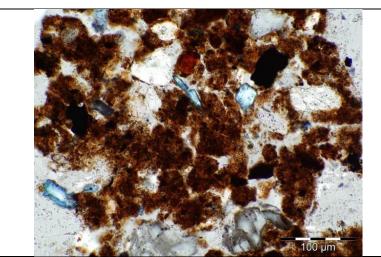


Fig. 4. Distribution pattern of organic aggregates in a thin section of the Aan of Valenakker (40-50 cm). In the intern fabric of the aggregates are charcoal particles visible.

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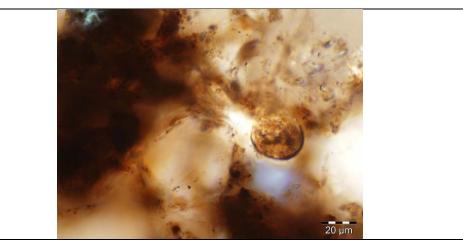
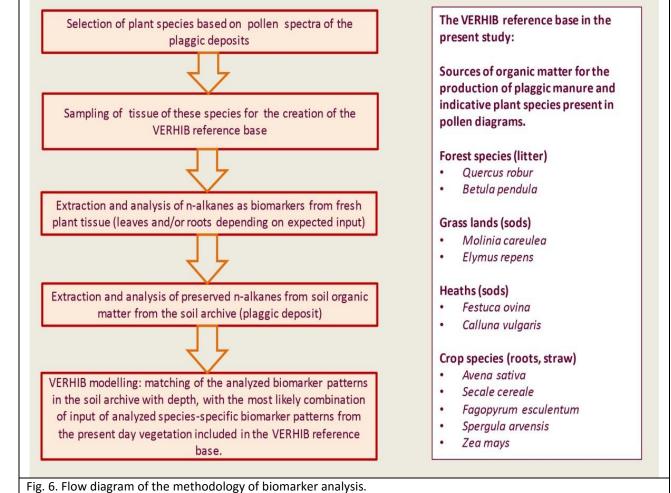
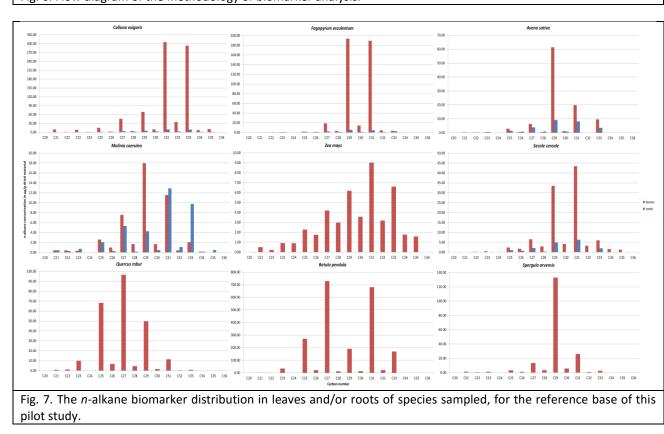


Fig. 5. Pollen grains, visible in an aged aggregate of the same thin sections. Pollen grains are in thin sections observable as not double fringing and empty spheroidal objects. The palynological characteristics as sculpture and aperture are not visible without the chemical treatments during pollen extraction.







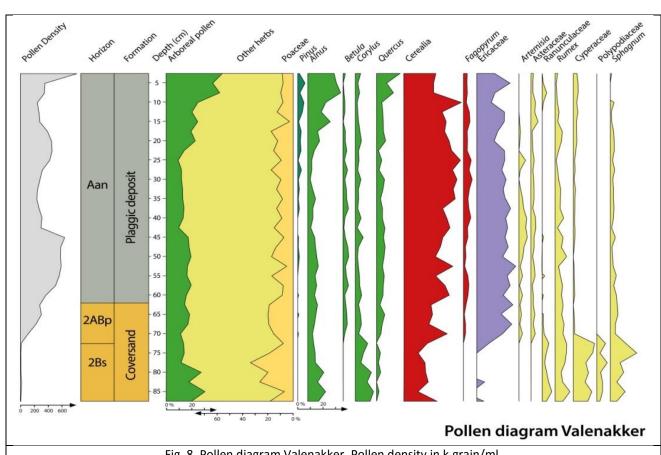
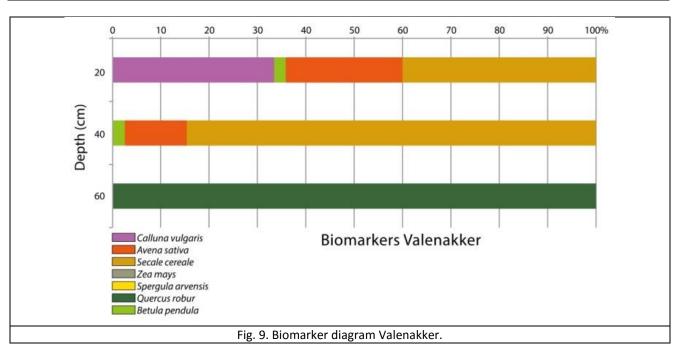


Fig. 8. Pollen diagram Valenakker. Pollen density in k.grain/ml.

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Horizon	Depth (cm)	Calendric ¹⁴ C ages humin	Calendric ¹⁴ C ages humic acids	Calendric OSL ages		
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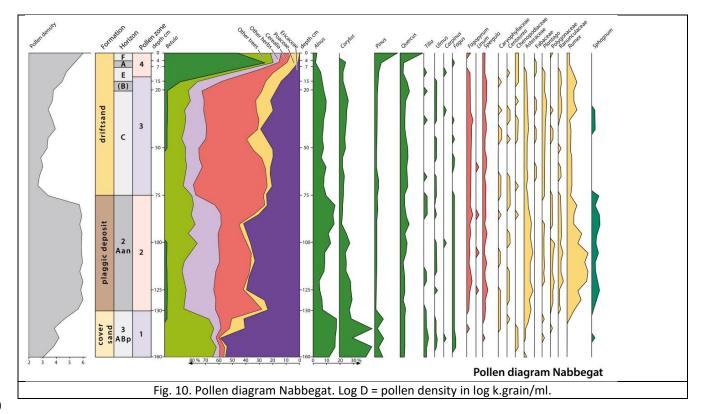
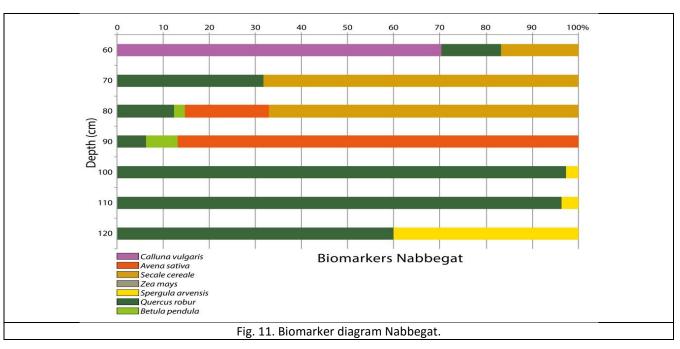




Table 2. ¹⁴ C and OSL dates of the plaggic deposits of Nabbegat.					
Horizon	Depth (cm)	Calendric ¹⁴ C ages humin	Calendric ¹⁴ C ages humic acids	Calendric OSL ages	
C	70	-	_	1803 ± 12 AD	
2An	80	428 ± 107 AD	626 ± 45 AD	1770 ± 11 AD	
2An	105	37 ± 133 BC	3 ± 101 AD	-	
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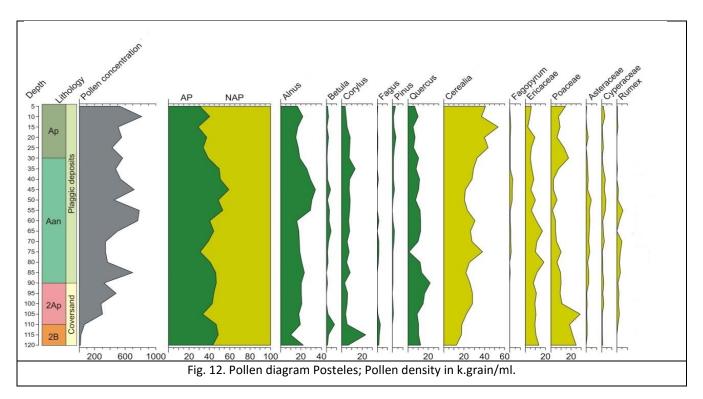
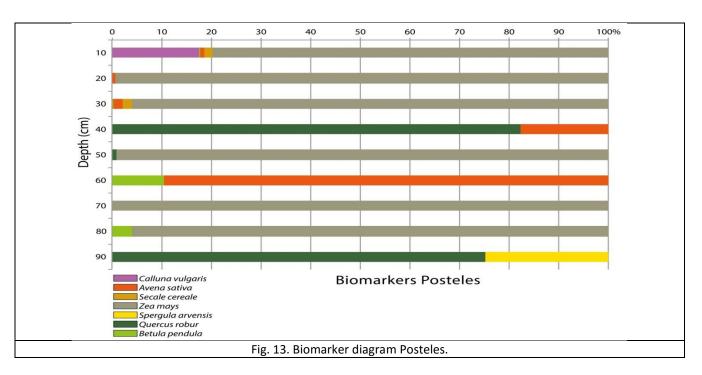


Table 3. ¹⁴ C and OSL dates of the plaggic deposits of Posteles.						
Horizon	Depth cm	Calendric ¹⁴ C ages humin	Calendric ¹⁴ C ages Humic acids	Calendric OSL ages		
Aan	45	-	-	1758 ± 14 AD		
Aan	59	-	_	1711 ± 20 AD		
Aan	70	1132 ± 68 AD	1172 ± 51 AD	1651 ± 31 AD		
Aan	82	-	_	1626 ± 20 AD		
Aan	95	884 ± 82 AD	861 ± 85 AD	1517 ± 31 AD		
2ABp	105	-	-	2035 ± 450 BC		



634 635 636 Figures 1 – 13 (including figure captions).