

1 Point-by-point response to the reviews, a list of relevant changes.

- 2 1. Additional text for biomarker paragraph and two figures (6, flow diagram and 7, mass spectra
- 3 for reference base) to explain better the biomarker analysis procedure.
- 4 2. We inserted two sentences (incl. references) to inform better about heath management and
- 5 the consequences for species in pollen and biomarker spectra.
- 6 3. We corrected the tables of the ^{14}C and OSL dates and the depths of some spectra.
- 7 4. We removed the two microphotographs and replaced them by 3 brand new pictures (figures
- 8 3, 4 and 5).

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11 **The added value of biomarker analysis to the genesis of Plaggic Anthrosols; the identification of**

12 **stable fillings used for the production of plaggic manure.**

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20 **Abstract.**

Plaggic Anthrosols are the result of historical forms of land management in cultural landscapes on chemically poor sandy substrates. Application of plaggic manure was responsible for the development of the plaggic horizons of these agricultural soils. Pollen diagrams reflect aspects of the environmental development but the interpretation of the pollen spectra is complicated due to the mix of the aeolian pollen influx of crop species and species in the surroundings, and of pollen occurring in the used stable fillings. Pollen diagrams and radiocarbon dates of plaggic Anthrosols suggested a development period of more than a millennium. *Calluna* is present in almost all the pollen spectra, indicating the presence of heath in the landscape during the whole 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 fillings, used for the production of plaggic manure and responsible for the rise of the soil surface, cannot be identified with pollen diagrams alone. Biomarker analyses provide more evidence about the sources of stable fillings. The oldest biomarker spectra of the plaggic horizons of three typical plaggic Anthrosols examined in this study, were dominated by biomarkers of forests species as *Quercus* and *Betula* while the spectra of middle part of the plaggic horizons were dominated by biomarkers of stem tissue of crop species as *Secale* and *Avena*. Only the youngest spectra of the plaggic horizons were dominated by biomarkers of *Calluna*. This indicates that the 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 sods. We conclude that other sources of materials, containing mineral grains must have been responsible for the raise of the plaggic horizon.

42 **Key words**

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

47 **1. Introduction.**

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).

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

66 The use of ectorganic matter from forest soils in the Dutch coversand area, must have been strongly reduced in
67 the 11th-13th century, due to commercial forest clear cuttings as recorded in archived documents (Vera, 2011).
68 These deforestations resulted in a regional extension of sand drifting and the managers of the heaths had to
69 protect their valuable ecotopes against this 'historical environmental catastrophe' (Vera, 2011).
70 Heaths were already present in the Late Paleolithic landscape (Doorenbosch, 2013) and played a ceremonial
71 role in the society of our ancestors. People already had the knowledge to manage the heath as sustainable
72 grazing areas for cattle (Doorenbosch, 2013).

73 The use of heath for sheep grazing and other purposes as honey and oil production could continue until the
74 middle of the 18th century (Vera, 2011). In SE-Netherlands sustainable use of the heaths was promoted by
75 many management rules and laws (van Mourik, 1978; Veera 2011). Over the course of the 18th century, the
76 population growth resulted in an increasing food demand. In the course of the 18th century, the deep stable
77 economy was introduced and the booming demand for manure resulted in intensification of manure production.
78 Farmers started with the use of heath sods as (additional) stable filling (Spek, 2004). This caused heath
79 degradation and initiated the second extension of sand drifting. The use of sods finished at the end of the 19th
80 century after the introduction of chemical fertilizers (Spek, 2004).

81 Through the combination of OSL and ^{14}C dating, historical records and the conventional paleoecological proxy
82 of fossil pollen analysis we have a good impression of the paleoecological environment and the age of such
83 deposits. However, it remains problematic to reconstruct the combination of crop residues and various
84 materials used by farmers as stable filling to produce the stable manure, together responsible for the rise of
85 the surface of Anthrosols. This is also hindering a detailed interpretation of the agricultural practices and shifts
86 therein related to the plaggic agriculture system, and specifically the timing of the onset of the intensive heath
87 sod driven deep stable agriculture with which plaggic Anthrosols are most commonly associated. To address
88 this issue, in the present study we expanded our paleoecological toolset with an adapted application of the
89 recently developed biomarker approach (Jansen et al., 2010). This biomarker approach consists of a
90 combination of analytical chemical analysis and modelling with the VERHIB model to unravel concentration
91 patterns of higher chain length ($\text{C}_{20}\text{-C}_{36}$) *n*-alkanes of higher plant origin preserved in a soil or sedimentary
92 archive into the (groups of) species responsible for their production (Jansen et al., 2010). The approach was
93 originally developed to unravel past local vegetation composition. Upon successful application in a tropical
94 ecosystem setting (Jansen et al., 2013), its applicability in palaeopedology was explored (van Mourik & Jansen,
95 2013b). This pilot application concerned a polycyclic soil sequence in driftsand deposits. It showed that the
96 comparison of pollen and biomarker spectra allowed us to indicate the plant species responsible for carbon
97 sequestration in the humic horizons (van Mourik & Jansen, 2013b). Important conclusion was that biomarker
98 analysis showed promise not only in the reconstruction of past local vegetation composition of a specific site,
99 but also in studies where the emphasis lies not on the vegetation per se, but rather on reconstructing various
100 sources of soil organic matter input (van Mourik & Jansen, 2013b).

101 Goal of the present study was to further explore the applicability of biomarker analysis as part of a multi-proxy
102 reconstruction aimed at unraveling the sources of stable fillings used for the production of plaggic manure in
103 the context of the historic development of the plaggic agriculture ecosystem. For this, we applied biomarker
104 analysis on three previous investigated plaggic Anthrosol.

105 **Materials and methods.**

106 **2. Profile selection**

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Fig. 2

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112 The distribution area of plaggic Anthrosols in NW-Europe is indicated in fig.1. [Pape \(1972\)](#) published the first
113 map of the distribution of plaggic agriculture in NW-Europe. [Bastiaens & van Mourik \(1995\)](#) found traces of
114 intensification and extension of this area in Vlaanderen (Belgium) while [van Mourik \(1999b\)](#) also reported
115 plaggic Anthrosols in Schleswig (Germany). Beside this area with 'real' plaggic Anthrosols, [Spek \(2004, p. 724\)](#)
116 summarized information about the occurrence of soils with some evidence of application of plaggic manure in
117 the Atlantic coastal zones of Norway, Denmark, France, Galicia, Scotland and Ireland.

118 For this pilot study we selected three previously investigated plaggic Anthrosols in the Netherlands with an
119 undisturbed plaggic horizon: Valenakker, Nabbegat and Posteles (fig.2). Pollen diagrams, ^{14}C and OSL dates of
120 these profiles were available and previously published separately in various articles. Here we combined these,
121 and re-sampled the plaggic horizons of the profiles for biomarker analysis and new fossil pollen analysis to
122 allow for comparison. Vertical sampling resolution was: Valenakker 20 cm, Nabbegat and Posteles 10 cm.

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

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

132

Fig. 3

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

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141 2.1. Pollen analysis.

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

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Fig. 4

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Fig. 5

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152 Samples for pollen extraction were collected in 10 ml tubes in profile pits. For a correct matching of pollen and
153 biomarker spectra of the plaggic deposits, the same samples were treated for both pollen and biomarker
154 extraction and analysis. The pollen extractions were carried out using the tufa extraction method ([Moore et al.,
155 1991, p. 50](#)). For the identification of pollen grains, the pollen key of [Moore et al. \(1991, p. 83-166\)](#) was applied.
156 Pollen scores were based on the total pollen sum of arboreal and non-arboreal plant species. For the
157 estimation of the pollen concentrations of the various soil horizons, the exotic marker grain method was
158 applied ([Moore et al., 1991, p. 53](#)).

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161 2.2. ^{14}C and OSL dating.

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163 The determination of the age of plaggic deposits is subjected to various complications ([Spek, 2004](#)). Pollen
164 stratification is disturbed by bioturbation and ploughing. Besides, the pollen content is a mix of the regular

165 pollen influx and pollen in stable fillings, used for the production of stable manure (van Mourik et al., 2011).
166 The ages of humic horizons of buried Podzols cannot be correctly determined by ¹⁴C dating due to the complex
167 composition of soil organic carbon (van Mourik et al., 1995). During a period of active soil formation, hard
168 decomposable organic carbon can accumulate in the Ah horizon, especially in the humin fraction but also in the
169 humic acid fraction. Especially the accumulation of charcoal fragments in the organic aggregates is responsible
170 for overestimation of the ¹⁴C ages (fig.4). During the Early Holocene small amounts of charcoal fragments were
171 released after (natural) forest fires, but the amount increased drastic in the iron time due to the charcoal
172 production for the melting of iron from placic horizons and iron stone (Beukenkamp and Sevink, 2005). The age
173 of the humic acid fraction was considered the best estimate of the moment of fossilization of the Ah horizon
174 after burying by driftsand. The difference between humin and humic acids ages was interpreted as a measure
175 for the period of soil activity and humin accumulation. Later, OSL dating confirmed that radiocarbon dates, not
176 only of the humin fraction but also of the humic acids, overestimate the true ages (Bokhorts et al., 2005).
177 Conventional radiocarbon dating of humin and humic acids showed in presented diagrams, extracted from
178 plaggic deposits, was performed in the CIO (Centre for Isotope Research of the University of Groningen).
179 OSL dates provide reliable information about the moment of fossilization of plaggic material under the rising
180 furrow because the quartz grain were perfectly bleached during active ploughing (Bokhorts et al., 2005). OSL
181 dating of quartz grains, extracted from plaggic deposits, was performed in the NCL (Netherland Centre for
182 Luminesce Dating, Wageningen University).

184 2.3. Biomarker analysis.

187 2.3.1. The application of the VERHIB model

188 A detailed description of the biomarker approach using the VERHIB method is presented in our previous
189 publications (Jansen et al., 2010; Jansen et al., 2013; Van Mourik & Jansen 2013). Briefly, the basis of the
190 method lies in the unraveling of the preserved concentration patterns of C₂₀-C₃₆ *n*-alkanes, which are exclusive
191 to the epicuticular wax layers on leaves and roots of higher plants (Kolattukudy et al., 1976). While such an
192 application in itself is not new (e.g. Pancost et al., 2002; Hughen et al., 2004) the novelty of our approach lies in
193 the application of the VERHIB model that we specifically developed to unravel the mixed *n*-alkane signal
194 encountered in soil or sedimentary archives (Jansen et al., 2010). The VERHIB model consists of a linear
195 regression model that describes how a certain input of plant derived compounds such as *n*-alkanes over time in
196 a certain archive at a certain location, results in accumulation of these compounds. An inversion of the forward
197 model is used to reconstruct the accumulation encountered with depth into its most likely vegetation origin
198 (Jansen et al., 2010). An important aspect of biomarker analysis using VERHIB is that it is an indirect
199 reconstruction. While the biomarker patterns, in the present study the *n*-alkanes, are directly measured, the
200 reconstruction into the most likely combination of vegetation biomass input responsible for the observed
201 pattern is inferred by the model. For this, several parameters must be inputted into the model (Jansen et al.,
202 2010) the most important of which is the selection of the expected plant species that have been responsible for
203 the input of biomass in the archive in question, and subsequent inclusion of their *n*-alkane signature in the
204 VERHIB reference base. In the present study, the selection of species to include was based on the (expected)
205 crop history of the sites under study, as well as the anticipated origin of the stable fillings used as manure. An
206 important matter of debate when using *n*-alkane patterns to reconstruct past vegetation input is the genotypic
207 plasticity of the *n*-alkane patterns, in particular in relation to prevailing environmental factors such as climate
208 (e.g. Shepherd and Griffiths, 2006). In a previous study focusing on vegetation of relevance for reconstructions
209 in ecosystems in North-Western Europe where plaggic agriculture occurred, we found that while genotypic
210 plasticity related to climatic factors may influence the signal, such influence does not eradicate the different
211 vegetation origins (Kirkels et al., 2013). To limit external influences as much as possible, the vegetation selected
212 for inclusion in the VERHIB reference base was sampled in close vicinity to the three study sites as much as
213 possible. The first group of selected plant species concerned the main sources of stable fillings, used for the
214 manure production: fermented litter from deciduous forest soils (*Quercus robur*, *Betula pendula*), grass sods
215 from brook valleys (*Molinia caerulea*) and heath sods (*Calluna vulgaris*).
216 The second group concerned crop species. Close to the educational Field Study Centre Orvelte (Drenthe) is a
217 traditional plaggic field where they continued with the cultivation of traditional crop species. There we sampled
218 *Fagopyrum esculentum*, *Spergula arvensis*, *Avena sativa*, *Secale cereal*, *Spergula arvensis*. The modern crop
219 species *Zea mays* corn was sampled on the Posteles.

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

223 A second parameter that must be considered in the application of VERHIB, is input of leaf and root material.
224 VERHIB considers the species specific *n*-alkane patterns in plant roots separately from the patterns in plant
225 leaves, and uses this to deal with the input of young root material at depth (Jansen et al., 2010). A first
226 selection criterion here concerns whether or not leaf and root material can be expected to have entered the
227 soil at all. For the deciduous forest soil material potentially used as stable fillings (*Quercus robur*, *Betula*
228 *pendula*), exclusively leaf derived biomass input is expected as the trees did not grow on-site. In contrast, for
229 the crop species *Zea Mays* and *Spergula Arvensis* only root material is expected to have entered the soil in
230 appreciable amounts as the leaf material is mostly removed during harvest. For the other species considered,
231 both leaf and root material must be taken into account. A selection of root and/or leaf derived *n*-alkane
232 patterns to be considered in the VERHIB reference base was made in accordance with the previous. With
233 respect to the ratio of input of leaf vs. root biomass as required by the model, no exact information is available
234 for the soil profile under study. Therefore, for those species where both leaf and root material is considered to
235 have possibly entered the soil, in line with the exploratory nature of the present study, we applied an assumed
236 leaf/root biomass input ratio of 1.0 and assumed that while input of leaf material always occurred at the top of
237 the soil profile, root input also occurred with depth. Since our pilot study in polycyclic driftsand deposits
238 showed that VERHIB was unable to filter out root input sufficiently (Van Mourik & Jansen, 2013), when
239 interpreting the occurrence of a certain species with depth in the profiles under study as modelled by VERHIB,
240 the possibility of young root input being responsible for the signal was explicitly taken into account.

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

243

Fig. 6

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2.3.2. Extraction and analysis of the biomarkers.

247 Approximately 0.1 g of each of the freeze-dried and ground vegetation and soil samples was extracted by
248 accelerated Solvent Extraction (ASE) using a Dionex 200 ASE extractor. The extraction temperature was 75°C
249 and the extraction pressure 17×10^6 Pa, employing a heating phase of 5 min and a static extraction time of 20
250 min. Dichloromethane/methanol (DCM/MeOH) (93:7 v/v) was used as the extractant (Jansen et al., 2006). The
251 extracts were subsequently fractionated into three fractions containing the *n*-alkanes, the esters and the
252 combination of alcohols and fatty acids respectively. For this, a silica column consisting of extracted cotton
253 wool and silica gel was used, followed by elution with hexane, hexane/DCM (4:1) and DCM/Methanol (9:1)
254 respectively. Separation of the *n*-alkanes took place by on-column injection of 1.0 µl of the first fraction on a 30
255 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
256 carrier gas. Temperature programming was: 50°C (hold 2 min); 40°C/min to 80°C (hold 2 min); 20°C/min to
257 130°C; 4°C/min to 350°C (hold 10 min). Subsequent MS detection in full scan mode used a mass-to-charge
258 ratio (*m/z*) of 50-650 with a cycle time of 0.65 s and followed electron impact ionization (70 eV). The *n*-alkanes
259 were identified from the total ion current (TIC) signal by their mass spectra (dominant fragment ion
260 represented by *m/z* = 57) and retention times and quantified using a deuterated internal standard (*d*₄₂-*n*-C₂₀
261 alkane (Jansen et al. 2010) as well as a conventional external *n*-alkane standard.

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Fig.7

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

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

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3.1. Profile Valenakker

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

275 The post sedimentary pollen spectra in the 2BS show percentages of tree species as *Corylus* and *Quercus* of the
276 Middle Subatlantic. The presence of Poaceae, Cyperaceae, *Rumex* and Ranunculaceae reflects a period of
277 pasture. The high scores of Cerealia in ploughed 2ABp and even the 2B indicate a form of sedentary agriculture
278 before the start of plaggic agriculture. ¹⁴C dating indicate a carbon age of the base (60 cm) of the Aan horizon
279 of ≈ 600 AD. The OSL age of the lower part of the plaggic horizon is 800-900 year younger, ≈ 1560 AD.
280

Fig.8.

Table 1.

Fig.9.

283 Micromorphological observations (fig.5ab) of the plaggic deposits show the complexity of soil organic matter.
284 There are various sources of organic carbon as plants roots, tissue of table fillings and sods. Also the
285 composition of pollen spectra is complex, a mix of the regular pollen influx of plants on the fields and in the
286 surrounding infiltrating into the soil and pollen, and pollen present in various stable fillings.
287 In previous studies the origin of stable fillings, used in plaggic agriculture, was reconstructed on the base of
288 pollen diagrams (Spek, 2004; van Mourik et al., 2012a, 2012b). The pollen spectra of the Aan horizon show very
289 low scores of arboreal trees but reasonable scores of Ericaceae and Poaceae. Ericaceae pollen may indicate the
290 use of heath sods, Poaceae pollen the use of grassland sods, the combination of sods from degrading heath and
291 the rise of the land surface by plaggic manure is caused by the mineral fraction in such sods. However, the rise
292 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
293 sods (with a mineral fraction) was introduced in the course of the 18th century when better construction
294 materials enabled the farmers to build deep stables (Vera, 2011). In fact, the sources of stable fillings cannot be
295 satisfactorily detected with pollen diagrams.
296 The biomarker spectrum of the base is dominated by *Quercus*. Despite the low percentages *Quercus* pollen it is
297 very likely that the farmers used forest litter as stable filling. The middle spectrum is dominated by markers of
298 *Avena* and *Secale*. This points to the use of straw from these crop species as stable filling. Pollen of Cerealia is
299 present in the whole diagram. In the upper spectrum biomarkers of *Calluna* are present together with *Avena*
300 and *Secale*. This points to the use of heath sods as additional stable filling during the last phase in the
301 development of the plaggic horizon.
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304 3.2. Profile Nabbegat

Fig. 10.

Table 2.

Fig. 11.

308 Profile Nabbegat is a haplic Arenosol (with Mormoder humus form), overlying a plaggic Anthrosol, overlying a
309 ploughed umbric Podzol. The pollen diagram (fig.6.) and the absolute dates (table 2) reflect a soil development
310 of ≈ 3000 year.
311 The post sedimentary pollen spectra of the 3ABp reflect the start of agriculture (increase of Cerealia) on a
312 former heath (decrease of *Ericaceae*) in a surrounding with coppice hedges (*Quercus*, *Corylus*). Based on
313 radiocarbon dates, the agricultural activities started before ≈ 1000 BC, the OSL dates point to deposition of
314 plaggic material after ≈1500 AD.
315 The radiocarbon ages indicate that the farmers used organic matter with very few mineral 'contamination' for
316 a long time. The OSL ages indicate that the rise of the plaggic horizon started ≈ 1500 AD due to mineral grains
317 as part of the manure. The plaggic horizon developed between 1500 and 1800 AD. Around 1800 AD, short after
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319 the introduction of the deep stable economy (Vera, 2011), the plaggic Anthrosol was overblown by driftsand.
320 Apparently, the use of heath sods resulted in heath degradation, sand drifting and acceleration of the rise of
321 the plaggic horizon (van Mourik et al., 2012a). The sand drifting stabilized under planted *Quercus* trees; the
322 roots of these trees reached the buried Anthrosol and may have contributed the scores of biomarkers in the
323 upperpart of the buried plaggic horizon. The composition of the pollen spectra of the plaggic horizon is rather
324 uniform, dominated by Ericaceae and Cerealia.

325 Fig.7. shows the results of biomarker analysis. Biomarkers of *Quercus* were present in all the spectra, dominant
326 in the lower spectra, regular in the other spectra. This points to the use of forest litter as stable filling during
327 the development of the lower part of the plaggic horizon. The main crop species during this time was *Spergula*.
328 The middle part is dominated by markers of *Avena* and *Secale*, indicating the use of straw. Only in the upper
329 spectrum *Calluna* was found, indicating the use of heath sods during the last phase of the development of the
330 plaggic horizon.

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333 3.3. Profile Posteles

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Fig.12.

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Table 3.

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Fig.13.

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

340 The pollen content of the buried ploughed Podzol (2Ap, 2B) is post-sedimentary infiltrated in Late-Glacial
341 coversand by bioturbation and agriculture. Characteristic is the sharp decrease of pollen concentrations with
342 depth, shown by the pollen density curve. The spectra of the 2B horizon already reflect evidence of agriculture
343 (Cerealia) in a deforested landscape (low percentages of *Alnus*, *Quercus*, *Fagus*). The spectra of the 2Ap horizon
344 show increasing percentages of Cerealia.

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

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

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

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

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361

362 4. Discussion

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364 Pollen diagrams of plaggic Anthrosols provide valuable paleoecological information to reconstruct the soil
365 dynamics during the plaggic agriculture. However, interpretation of pollen diagrams is complicated. Pollen
366 grains, extracted from plaggic deposits, may originate from two sources (van Mourik et al., 2011). The first
367 source concerns the regional pollen influx from flowering species and local flowering crop species. Pollen grains

368 precipitate on the soil surface and may infiltrate into the Anthrosols by ploughing and bioturbation. This pollen
369 influx will be mixed with the pollen content of materials, used as stable filling to produce manure.
370 Pollen will be preserved in plaggic deposits in the anaerobic and acid micro environment of humic aggregates,
371 produced by worms and micro arthropods (van Mourik, 1999b, 2001). In general it is not possible to make a
372 clear separation between pollen grains originating from the regular pollen influx or from materials as sods.
373 Therefore, the identification of the various sources of stall fillings cannot be based on pollen analysis alone.
374 Additional information, acquired by biomarker analysis proved very useful for this purpose.
375 In the pollen diagrams *Fagopyrum* is found in almost all the spectra of the plaggic deposits and in Valenakker
376 and Nabbegat even in the top spectra of the buried ploughed Podzol, probably as result of pollen infiltration.
377 *Fagopyrum* as crop species on sandy soils was introduced after 1350 AD (Leenders, 1996). Based on this
378 palynological time marker, plaggic deposition started around 1350 AD.
379 The radiocarbon ages of plaggic deposits are much older. This is caused by (1) older organic carbon, present in
380 the applied stable fillings (as forest litter) for the manure production and (2) accumulation of hardly
381 decomposable organic carbon during active soil formation. Consequently the radiocarbon dates overestimate
382 the ages of the plaggic sediments, but approach the age of the introduction of agricultural soil management
383 (van Mourik et al., 1995, 2011, 2012a, 2012b). Manuring of infertile soils came already in use in the Bronze Age
384 and also the Celtic fields are an example of a prehistorical agricultural system based on manure management
385 (Spek, 2004).
386 The mineral component of stable manure, applied on the fields, was responsible for the thickening of the
387 plaggic horizon. Ploughing of the furrow will bleach the OSL signal of the mineral grains until the moment that
388 the grains are no longer part of the active soil furrow. For that reason, OSL dating of the plaggic horizon provide
389 reliable ages of the plaggic deposits (Bockhorst et al., 2005). The OSL dates of the profiles Valenakker,
390 Nabbegat and Posteles indicate a start of the thickening \approx 1550 AD.
391 It was not possible to determine the sources of stable fillings palynologically. Possible stable fillings were forest
392 litter, sods from moist grass lands and heath sods. But in almost all spectra of the pollen diagrams Ericaceae,
393 Poaceae and arboreal pollen occur. Biomarkers extracted from plaggic deposits, originate from two sources.
394 The first source concerns biomarkers from decomposed roots of crop species, the second source of organic
395 material as straw and sods, used as stable filling for manure production.
396 In the three diagrams we find *Quercus* as dominant marker in the lowest part of the Aan-horizon, indicating the
397 use of forest litter. In Nabbegat, *Quercus* markers can also originate from roots of the planted *Quercus* forest
398 after the stabilization of the sand drifting. This is not the case on Valenakker and Posteles. The middle part of
399 the Aan-horizon is dominated by markers of *Avena* and *Secale*, indicating the use of straw as stable filling.
400 Only in the top of the Aan-horizon markers of *Calluna* are present, indicating the use of heath sods as stable
401 filling. Based on the results of the biomarker analysis we can conclude that heath sods were used as stable
402 filling only in the 18th and 19th century. This fits with the observations about the use of heaths in historical
403 archives Vera (2011).
404 So the question rises about heath management before the introduction of the deep stable economy. Some
405 researchers point to careful heath management before the 19th century. In interviews with farmers, born
406 before 1950, Burny (1999) collected essential information about historical heaths management in the Belgian
407 Kempen. A historical study of land use in the Campina also indicated carefully maintenance and sustainable use
408 of valuable common fields (de Keyzer, 2014). Before the 19th century, heath sods were never dug on the dry
409 *Calluna* heath, only on the moist *Erica* heath. These organic sods were not used as stable filling but as fuel for
410 the furnace. Burning of *Calluna* heaths was the most important management action to rejuvenate the heath.
411 Juvenile heath is food for cows. Sods digging was a bad action due to the resistance and incoherence of these
412 dry sods and also the long recovery period. Mowing of older *Calluna* shrubs took place. Twigs were used for
413 roofs, burning and brooms. (Burny, 1999). Because of the very low nutrient contribution to the manure of
414 mowed *Calluna*, the farmers preferred the use of twigs of broom (*Genista*). When in the course of the 18th the
415 authority relationships changed and the population growth and the demand for food increased, farmers
416 started to intensify their production (Vera, 2011). They needed more manure and started with the deep stable
417 economy and the use of *Calluna* heath sods.
418 An important factor may be the presence of pollen and biomarkers content in sheep droppings. According to
419 Simpon et al., (1999) biomarkers survive the congestion process and stay in the manure. But what do sheep
420 consume? Grazing sheep are very selective in collecting food (Oom et al., 2008; Smits & Noordijk, 2013). They
421 prefer grasses (*Molinia*, *Festuca* and *Corynephorous*). Only in years that there is insufficient grass available at
422 the end of the summer, they eat shoots of *Calluna*, at that time nourishing with high concentrations Ca, Mg and
423 but no P. Pollen extractions from sheep droppings showed that only in droppings, collected during the summer
424 season *Calluna* pollen is present. During the flowering season of *Calluna*, the animals consume pollen,

425 precipitated on the grasses. That explains the presence of *Calluna* pollen and the absence of *Calluna*
426 biomarkers in the lower parts of the plaggic horizons.

427 If it is true that *Calluna* heath sods were dug only in the 18th and 19th century, how can we explain the mineral
428 component in the plaggic manure, responsible of the rise of the land surface before that time?

429 According to [Smits and Noordijk \(2013\)](#) there are several sources of minerals. Firstly, a small amount of mineral
430 grains will be incorporated in the manure during emptying out the manure of the stable. Secondly, farmers had
431 the knowledge that the addition of sand could improve the fertility of the soil. Not the leached and acid sand
432 from heath sods but not leached sand, dug on sheep walks and in blown out depressions in nearby drift sand
433 landscapes.

434

435

436 5. Conclusions

437

438 • The vertical zoning of biomarkers and pollen in plaggic horizons are different. Palynologically, the plaggic
439 horizon is a homogenous, the biomarker diagrams show differentiation.

440 • We can identify various stable fillings used, based on the vertical distribution of biomarkers.

441 • The biomarker spectra of the base layer of the plaggic horizon are dominated by biomarkers of deciduous
442 trees litter (dominated by *Quercus*), indicating the use of organic matter from the forest floor.

443 • The biomarker spectra of the middle part of the plaggic deposits are dominated by crop species (*Avena*,
444 *Secale*), indicating the use of straw from these species as stable filling during a relatively long time.

445 • Only the top spectra of the plaggic horizons are dominated by *Calluna*, indicating that heath sods were
446 used as stable filling only during the last phase in the development of the plaggic horizon.

447 • Profile Posteles shows the impact of the contribution of biomarkers of roots of *Zea mays*, introduced
448 around 1950 AD, suppressing the other species.

449 • The negligible percentages of *Calluna* in biomarker spectra of plaggic deposits with exception of the top,
450 suggest an overestimating of the use of heath sods in the traditional interpretation of the genesis of
451 plaggic horizons, the dominance of crop species in biomarker spectra of plaggic deposits suggests
452 underestimating of the use of straw as source material for the production of organic stable manure to
453 fertilize ancient arable fields.

454

455

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458 agriculture. We are grateful to Annemarie Philip (IBED, University of Amsterdam) for the preparation of the
459 pollen slides, Hans van der Plicht (CIO, University Groningen) for production of the radiocarbon dates and Jakob
460 Wallinga (NCL, Wageningen University) for the realization of the OSL dates. The digital illustration were
461 produced by Jan van Arkel (IBED, University of Amsterdam).

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463

464 References

465 Bakker, J., Moscol, M., Hooghiemstra, H., 2008. Holocene environmental change at the upper forest line in
466 northern Ecuador. *The Holocene* 18, 877-893.

467

468 Bastiaens, J., van Mourik, J.M., 1995. Bodemsporen van beddenbouw in het zuidelijk deel van het
469 pluggenlandbouw areaal. *Historisch Geografisch Tijdschrift* 1995, 81-90.

470

471 [Beukenkamp, P.C. and Sevink, J., \(2005\). Natuur en landschap. In: De Hoge Veluwe, natuur en kunst. Stichting
472 nationaal park De Hoge Veluwe en Wanders Uitgeverij, Zwolle. P. 40-96.](#)

473

474 Bokhorst, M.P., Duller, G.A.T., van Mourik, J.M., 2005. Optically Stimulated Luminescence Dating of a fimic
475 anthrosol in the Southern Netherlands. *Journal of Archaeological Science* 2005, 547-553.

476

477 Burny, J., 1999. Bijdrage tot de historische ecologie van de Limburgse Kempen (1910-1950). *Natuurhistorisch
478 Genootschap in Limburg XLII(1)*, Maastricht.

479
480 De Keyzer, M., 2014. The common denominator; the survival of the commons in the late medieval Campine
481 area. University Antwerpen, department of history, Belgium.
482
483 Doorenbosch, M., 2013. Ancestral heaths; reconstructing the barrow landscape in the central and southern
484 Netherlands. Sidestone Press, Leiden.
485
486 Everitt, B.S., Landau, S., Leese, M. and Stahl, D., 2011. Cluster Analysis, 5th Edition, John Wiley & Sons, Ltd,
487 Chichester, UK
488
489 Gocke, M., Kuzyakov, Y., Wiesenberg, G.L.B., 2010. Rhizoliths in loess - evidence for post-sedimentary
490 incorporation of root-derived organic matter in terrestrial sediments as assessed from molecular proxies.
491 Organic Geochemistry 41, 1198-1206.
492
493 Huguen, K.A., Eglinton, T.I., Xu, L., Makou, M., 2004. Abrupt tropical vegetation response to rapid climate
494 changes. Science 304, 1955-1959.
495
496 Jansen, B., Nierop, K.G.J., Hageman, J.A., Cleef, A.M., and Verstraten, J.M., 2006. The straight-chain lipid
497 biomarker composition of plant species responsible for the dominant biomass production along two altitudinal
498 transects in the Ecuadorian Andes. Organic Geochemistry 37, 1514-1536.
499
500 Jansen, B. Haussmann, N.S., Tonneijck, F.H., De Voogt, W.P. and Verstraten, J.M., 2008. Characteristic straight-
501 chain lipid ratios as a quick method to assess past forest - páramo transistons in the Ecuadorian Andes,
502 Palaeogeography, Palaeoclimatology, Palaeoecology, 262: 129-139.
503
504 Jansen, B., Van Loon, E.E., Hooghiemstra, H., and Verstraten, J.M., 2010. Improved reconstruction of palaeo-
505 environments through unravelling of preserved vegetation biomarker patterns. Palaeogeography,
506 Palaeoclimatology, Palaeoecology 285, 119-130.
507
508 Kirkels, F.M.S.A, Jansen, B., and Kalbitz, K., 2013. Consistency of plant-specific n-alkane patterns in plaggen
509 ecosystems: A review, The Holocene, 23: 1355-1368
510
511 Kolattukudy, P.E., Croteau, R., and Buckner, J.S., 1976. Biochemistry of plant waxes. In: Kolattukudy, P. E. (Ed.),
512 Chemistry and biochemistry of natural waxes. Elsevier, Amsterdam.
513
514 Leenders KHAW, 1996. De boekweitcultuur in historisch perspectief. Geografisch Tijdschrift, 21-3, 213-227.
515
516 ISRIC-FAO, 2006. World Reference Base for Soil Recourses 2006. World soil resources reports 103.
517
518 Mook, W.G., Streurman, H.J. 1983. Physical and chemical aspects of radiocarbon dating. First Symposium on
519 14C and Archaeology, Groningen, PACT 8: 31-55.
520
521 Moore, P.D., Webb, J.A., Collinson, M.E., 1991. Pollen analyses. Blackwell Scientific Publications.
522
523 Oom, S.P., Sibbald, A.M., Hester, A.J., Miller, D.R., Legg, C.J., 2008. Impacts of sheep grazing a complex
524 vegetation mosaic: Relating behavior to vegetation change. Agriculture, Ecosystems and Environment 124
525 (2008) 219–228.
526
527 Pape, J.C., 1972. Oude bouwlandgronden in Nederland. Boor en Spade 18, pp. 85-115
528
529 Rieley, G., Collier, R.J., Jones, D.M., Eglinton, G., 1991. The biogeochemistry of Ellesmere Lake, U.K. -I: source
530 correlation of leaf wax inputs to the sedimentary lipid record. Organic Geochemistry 17, 901-912.
531
532 Shepherd, T., Griffiths, D.W., 2006. The effects of stress on plant cuticular waxes. New Phytologist 171, 469-
533 499.
534

535 Simpson, I.A., van Bergen, P.F., Perret, V., Elhmmali, M.M., David, J., Roberts, D.J., Richard, R.P., 1999. Lipid
536 biomarkers of manuring practice in relict anthropogenic soils. *The Holocene* 9,2 (1999) 223–229.
537
538 Smits, J., Noordijk, J., 2013. Heidebeheer, moderne methoden in een eeuwenoud landschap. KNNV uitgeverij,
539 september 2013.
540
541 Spek, T., 2004. Het Drentse esdorpenlandschap, een historisch geografische studie. *Matrijs*, Utrecht, Volume 2,
542 part VI,725-967.
543
544 Tareq, S.M., Tanoue, E., Tsuji, H., Tanaka, N., Ohta, K., 2005. Hydrocarbon and elemental carbon signatures in a
545 tropical wetland: Biogeochemical evidence of forest fire and vegetation changes. *Chemosphere* 59, 1655-1665.
546
547 van Mourik, J.M., 1987. Het stuifzand van Heeswijk-Dinther. *Geografisch Tijdschrift* 21-4, 327-337.
548
549 van Mourik, J.M., 1999a. The use of micromorphology in soil pollen analysis. *Catena* 35, 239-257.
550
551 van Mourik, J.M., 1999b. Spuren von Plaggenlandbau im Gebiet der Schleswiger Landenge. *Offa* 47, 1990, 169-
552 176.
553
554 van Mourik, J.M., 2001. Pollen and spores, preservation in ecological settings. In: Briggs, E.G., Crowther, P.R.
555 (eds). *Palaeobiology II*. Blackwell Science, 315-318.
556
557 Van Mourik, J.M. and Horsten, F., 1995. De paleogeografie van de Valenakker. *Weerter Jaarboek* 1996, pp.
558 105-118.
559
560 van Mourik, J.M., Wartenbergh, P.E., Mook W.J., Streurman, H.J., 1995. Radiocarbon dating of palaeosols in
561 eolian sands. *Mededelingen Rijks Geologische Dienst* 52, 425-439.
562
563 van Mourik, J.M., Slotboom, R.T., Wallinga, J., 2011. Chronology of plaggic deposits; palynology, radiocarbon
564 and optically stimulated luminescence dating of the Posteles (NE-Netherlands). *Catena* 84, 54-60.
565
566 van Mourik, J.M., Seijmonsbergen, A.C., Jansen, B., 2012. Geochronology of Soils and Landforms in Cultural
567 Landscapes on Aeolian Sandy Substrates, Based on Radiocarbon and Optically Stimulated Luminescence Dating
568 (Weert, SE-Netherlands). *InTech (2012) Radiometric Dating* 75-114.
569
570 van Mourik, J.M., Seijmonsbergen, A.C., Slotboom, R.T., Wallinga, J., 2013a. The impact of human land use on
571 soils and landforms in cultural landscapes on aeolian sandy substrates (Maashorst, SE Netherlands). *Quaternary*
572 *International* 265 (2012) 74-89.
573
574 van Mourik, J.M., Jansen, B., 2013b. The added value of biomarker analysis in palaeopedology; reconstruction
575 of the vegetation during stable periods in a polycyclic driftsand sequence in SE-Netherlands. *Quaternary*
576 *International* 306 (2013) 14-23
577
578 Vera, H., 2011. 'dat men het goed van de ongeboornen niet mag verkoopen'; Gemene gronden in de Meierij
579 van Den Bosch tussen hertog en hertgang 1000-2000. Uitgeverij BOXpress, Oisterwijk, Netherlands (with
580 English summary).
581
582 Zech, M., Zech, R., Morras, H., Moretti, L., Glaser, B., Zech, W., 2009. Late Quaternary environmental changes in
583 Misiones, subtropical NE Argentina, deduced from multi-proxy geochemical analyses in a palaeosol-sediment
584 sequence. *Quaternary International* 196, 121-136.

585 **Tables (including table captions)**

586

Table 1. ¹⁴C and OSL dates of the plaggic deposits of Valenakker.

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

587

Table 2. ¹⁴C and OSL dates of the plaggic deposits of Nabbeget.

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	–
2An	130	1182 ± 139 BC	811 ± 101 BC	1676 ± 14 AD
3ABp	140	–	1299 ± 78 BC	–
3ABp	150	–	1385 ± 72 BC	–

588

Table 3. ¹⁴C 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

589

590

591 **Figure Captions**

592

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

595

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

598

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

603

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

606

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

610

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

612

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

615

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

617

618 Fig. 9. Biomarker diagram Valenakker.

619

620 Fig. 10. Pollen diagram Nabbegat. Log D = pollen density in log k.grain/ml.

621

622 Fig. 11. Biomarker diagram Nabbegat.

623

624 Fig. 12. Pollen diagram Posteles; Pollen density in k.grain/ml.

625

626 Fig. 13. Biomarker diagram Posteles.

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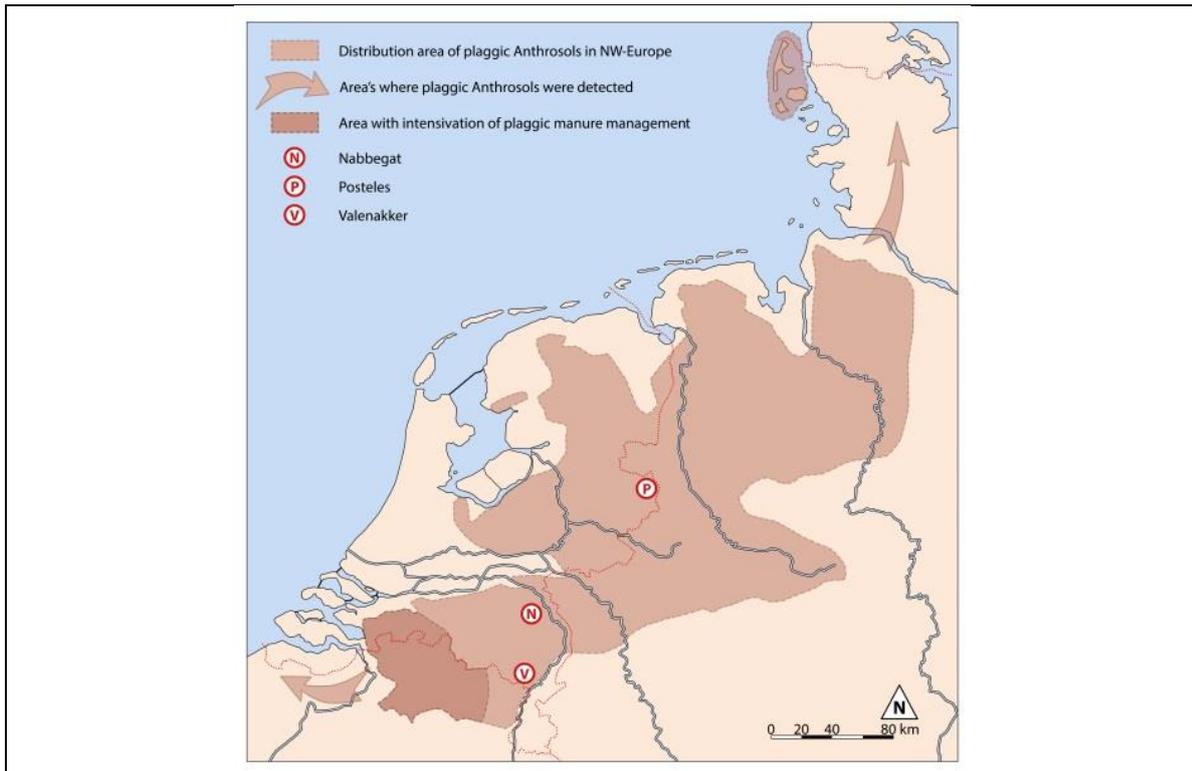


Fig. 1. The location of sampled profiles Valenakker, Nabbegat and Posteles in the distribution area of plaggic agriculture.

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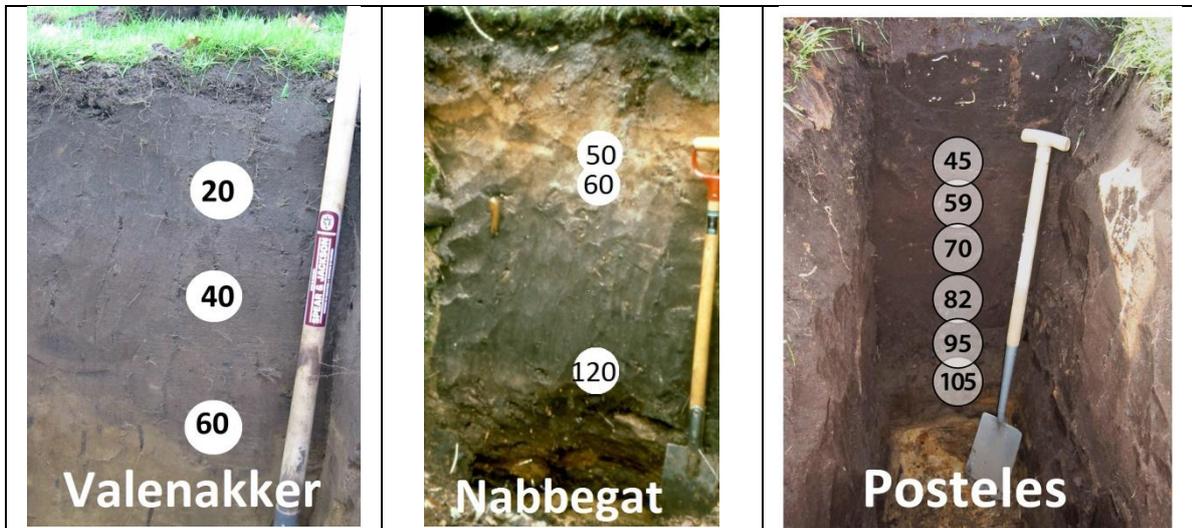


Fig. 2. The plaggic Anthrosols Valenakker, Nabbegat and Posteles. The location of the OSL samples are indicated in the white circles (depth in cm); the locations of the profiles are indicated in fig. 1.

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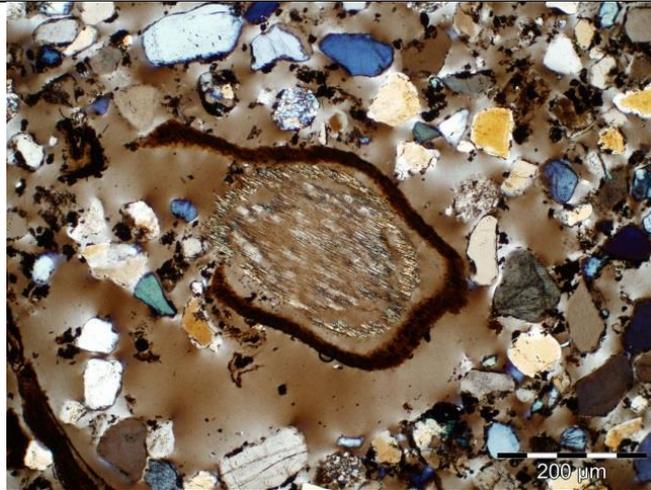


Fig. 3. Cross-section of a tree root in a thin section of the 2 An of Nabbegat (70-80cm). Characteristic is the double fringing of the root tissue. Such roots were only found in the upper part of the 2An of Nabbegat. Roots of crop species were not found in thin sections of the three profiles. The decompose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

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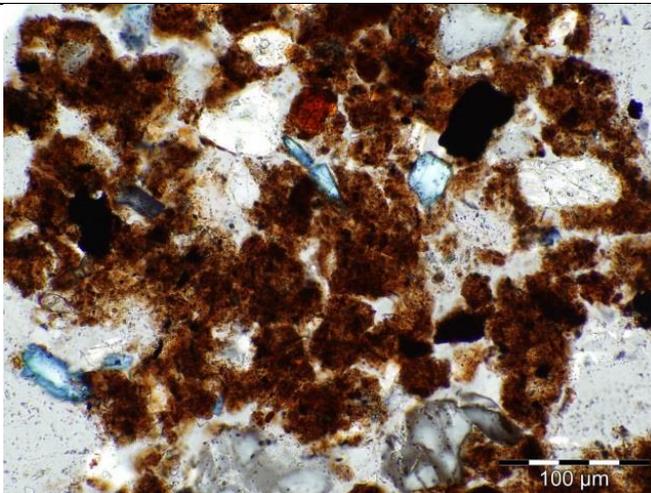


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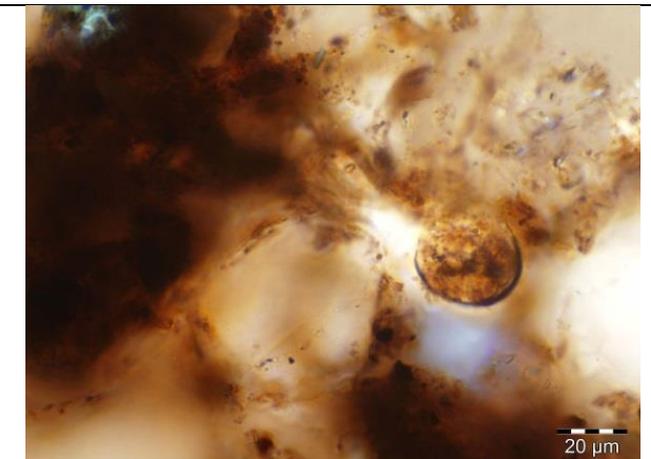


Fig. 5. Pollen grains, visible in an aged aggregate of the same thin sections. Pollen grains are in thin sections observable as not double fringing and empty spheroidal objects. The palynological characteristics as sculpture and aperture are not visible without the chemical treatments during pollen extraction.

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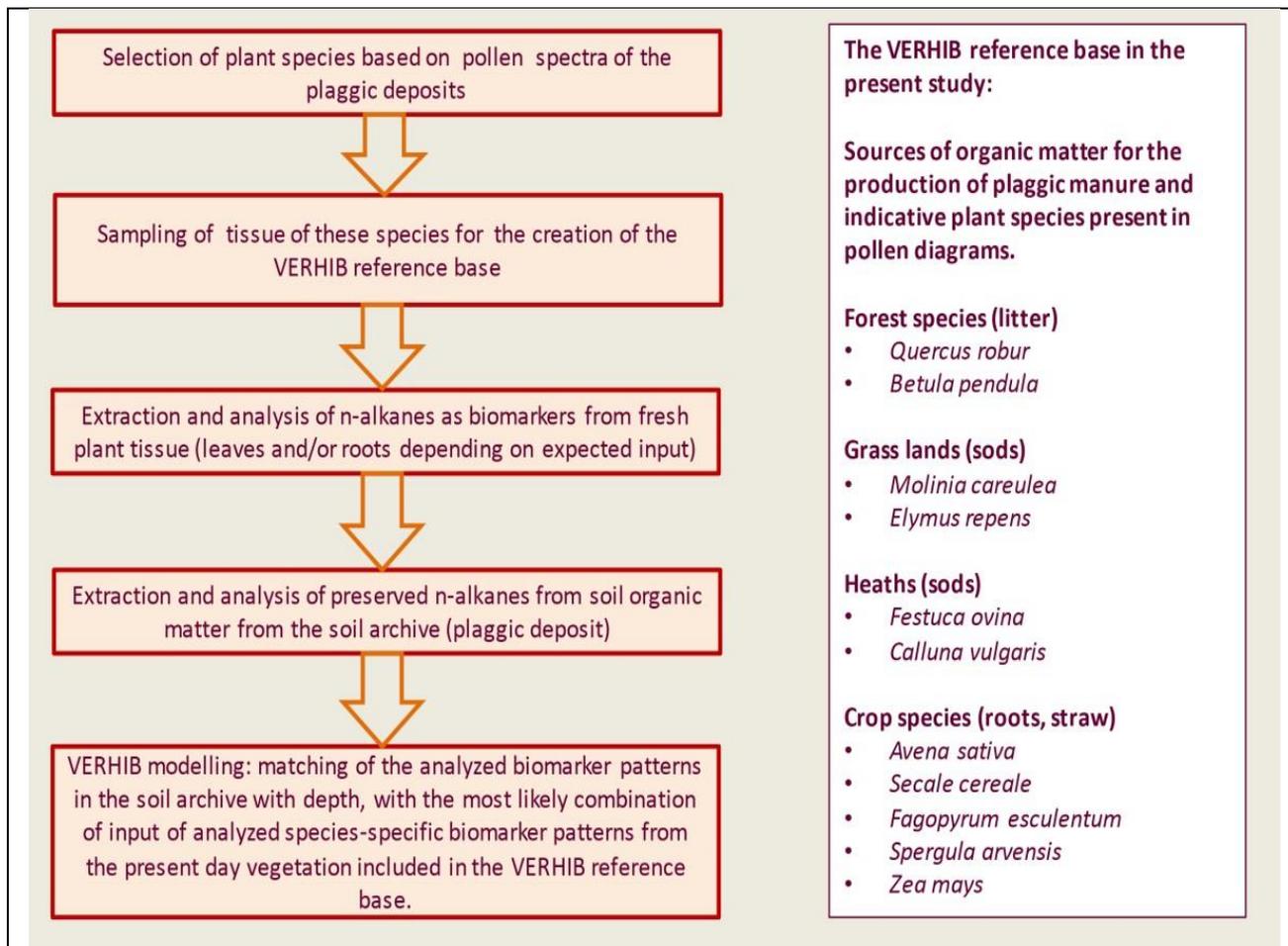


Fig. 6. Flow diagram of the methodology of biomarker analysis.

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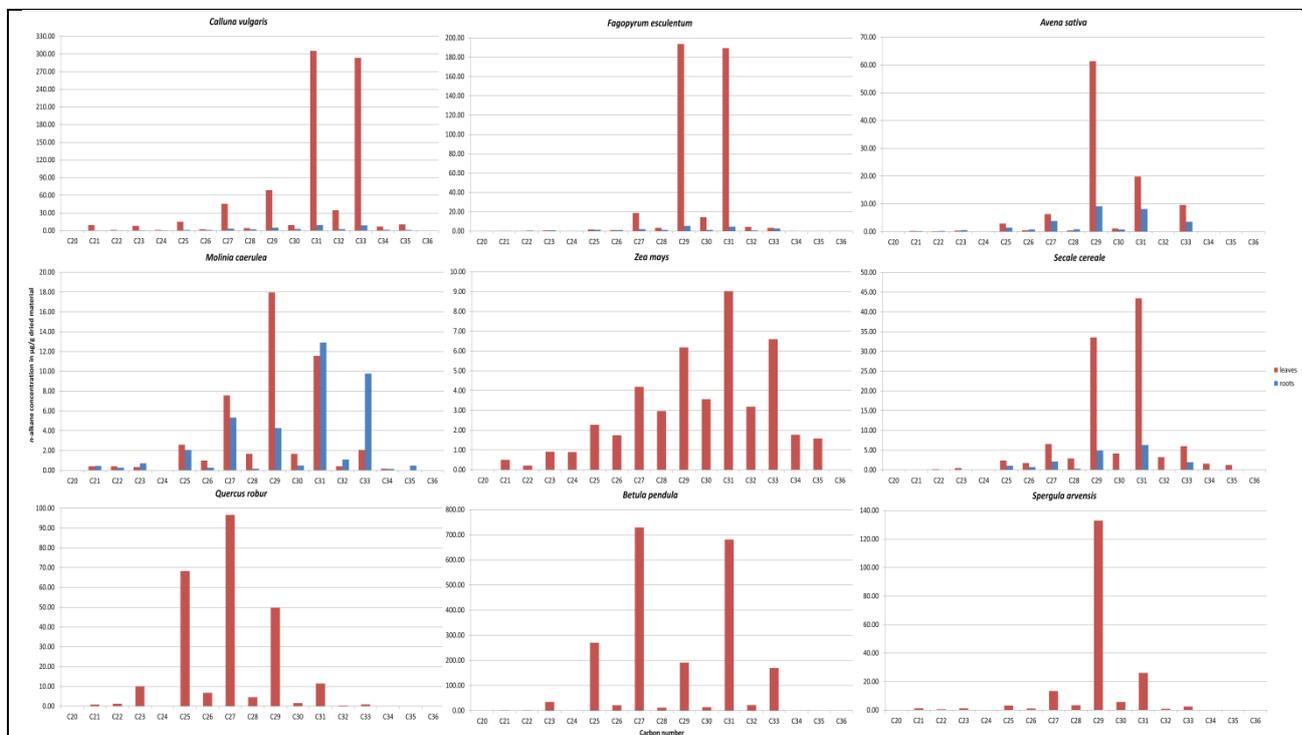


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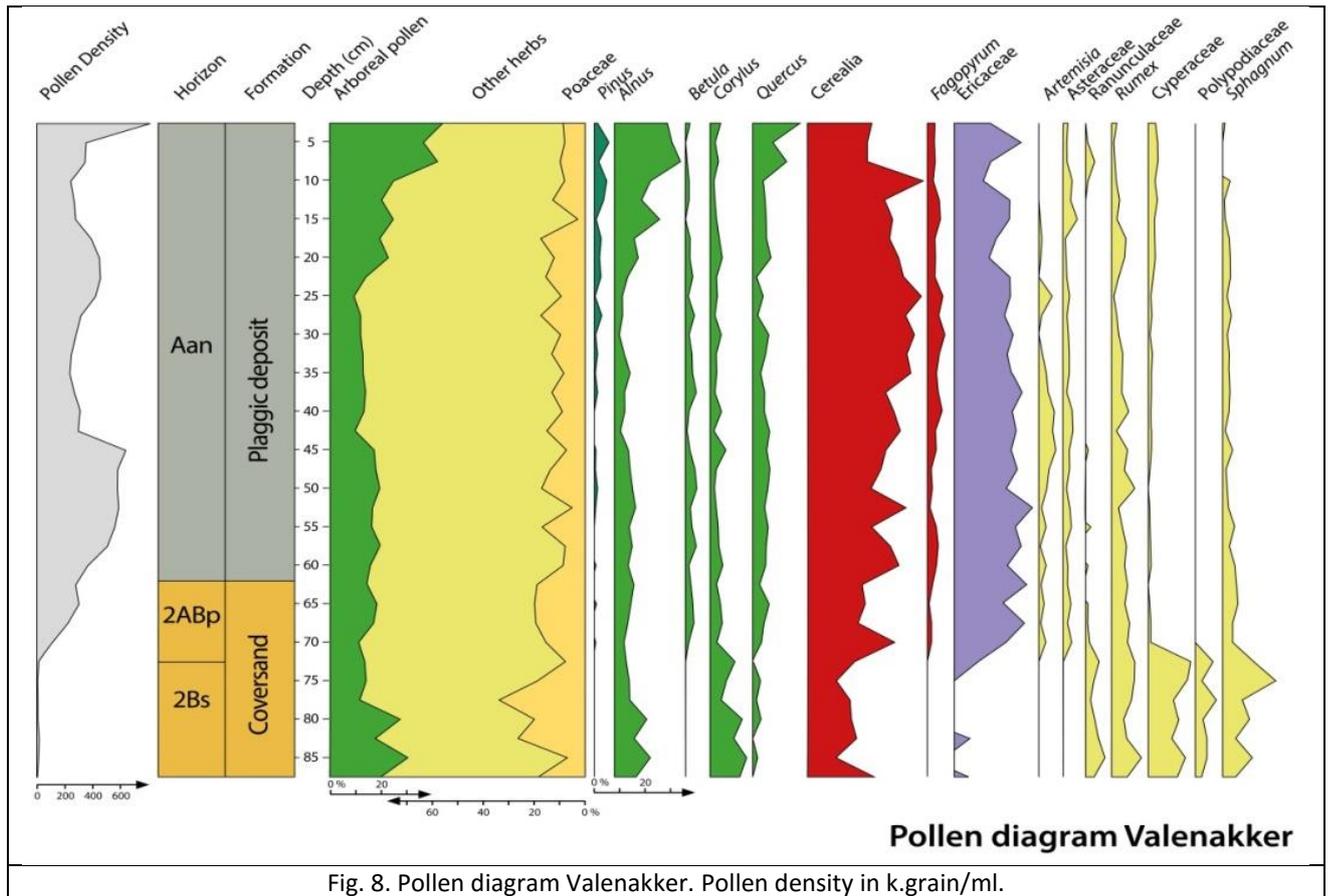


Fig. 8. Pollen diagram Valenakker. Pollen density in k.grain/ml.

637

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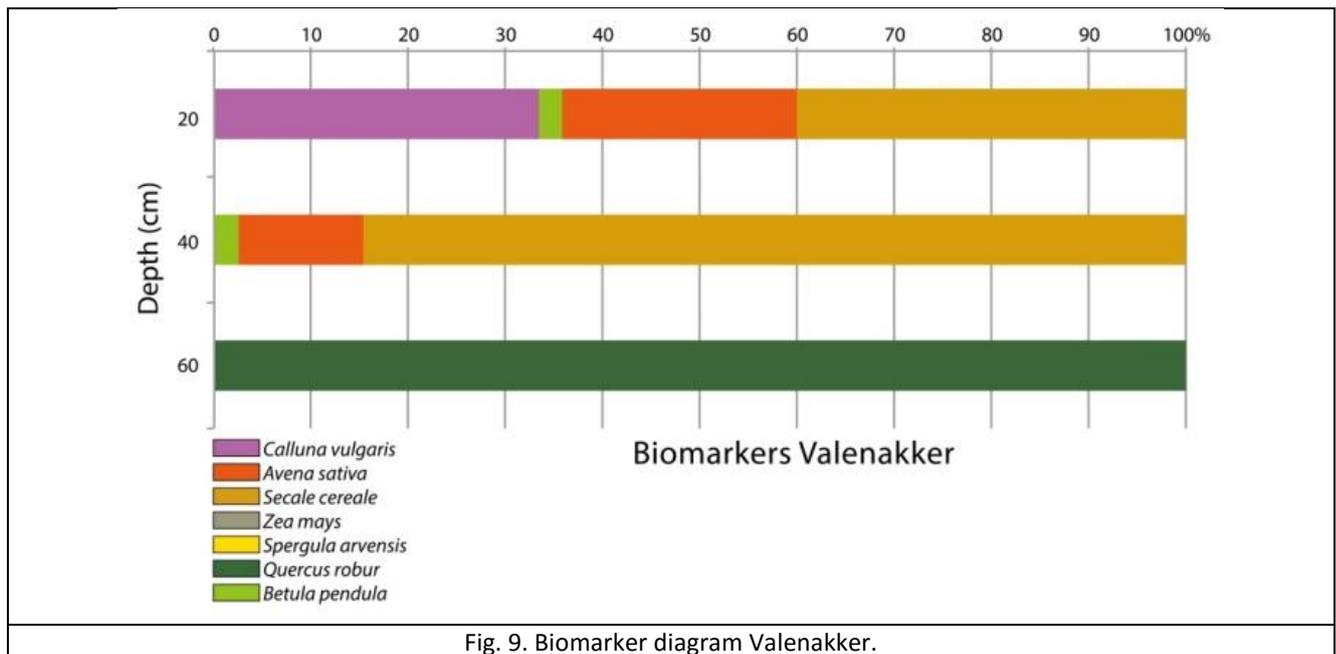


Fig. 9. Biomarker diagram Valenakker.

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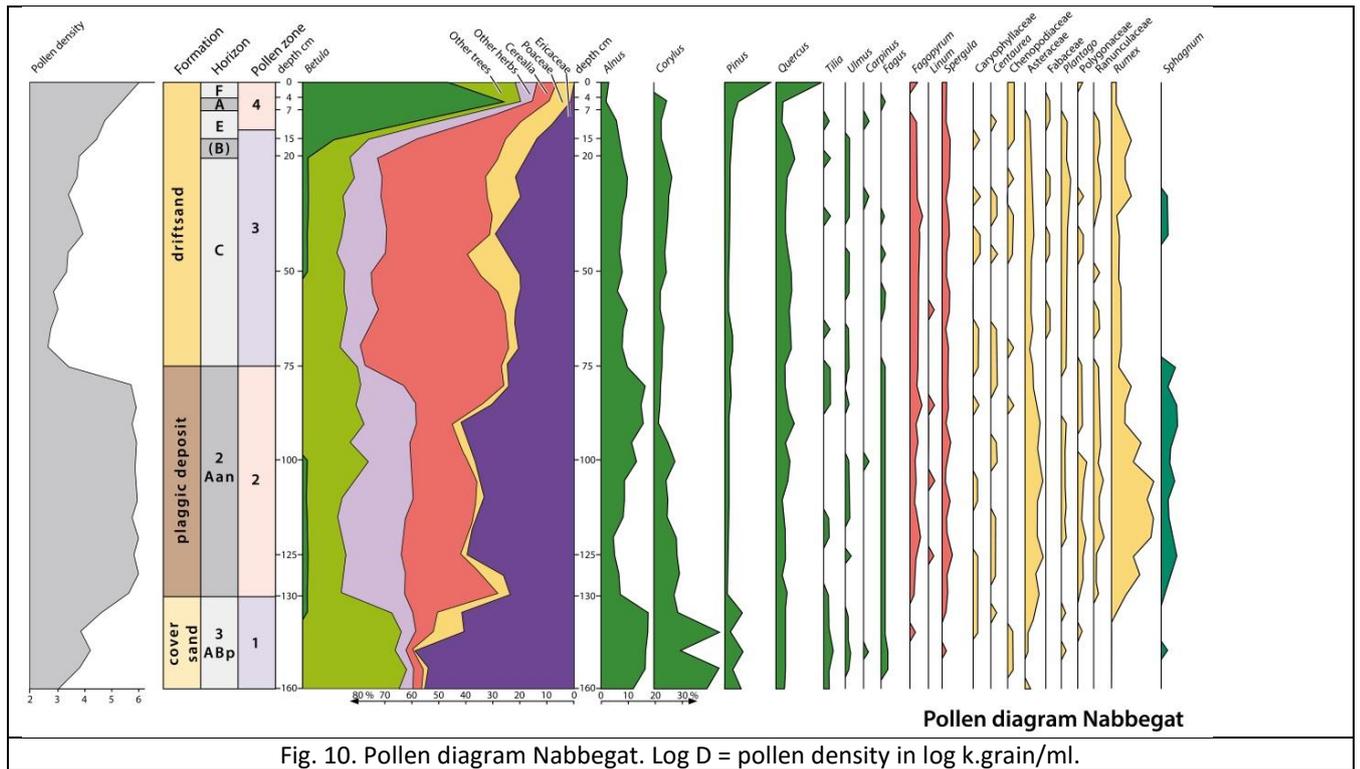


Fig. 10. Pollen diagram Nabbegat. Log D = pollen density in log k.grain/ml.

640

Table 2. ¹⁴C 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
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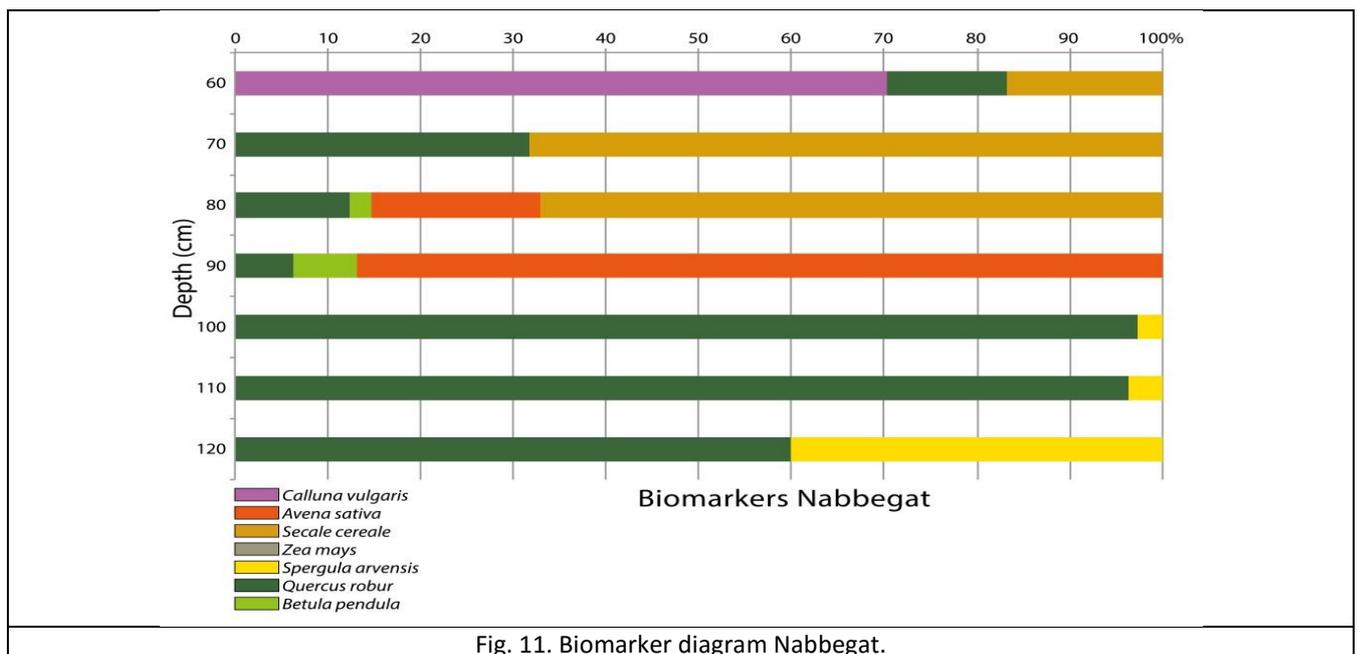
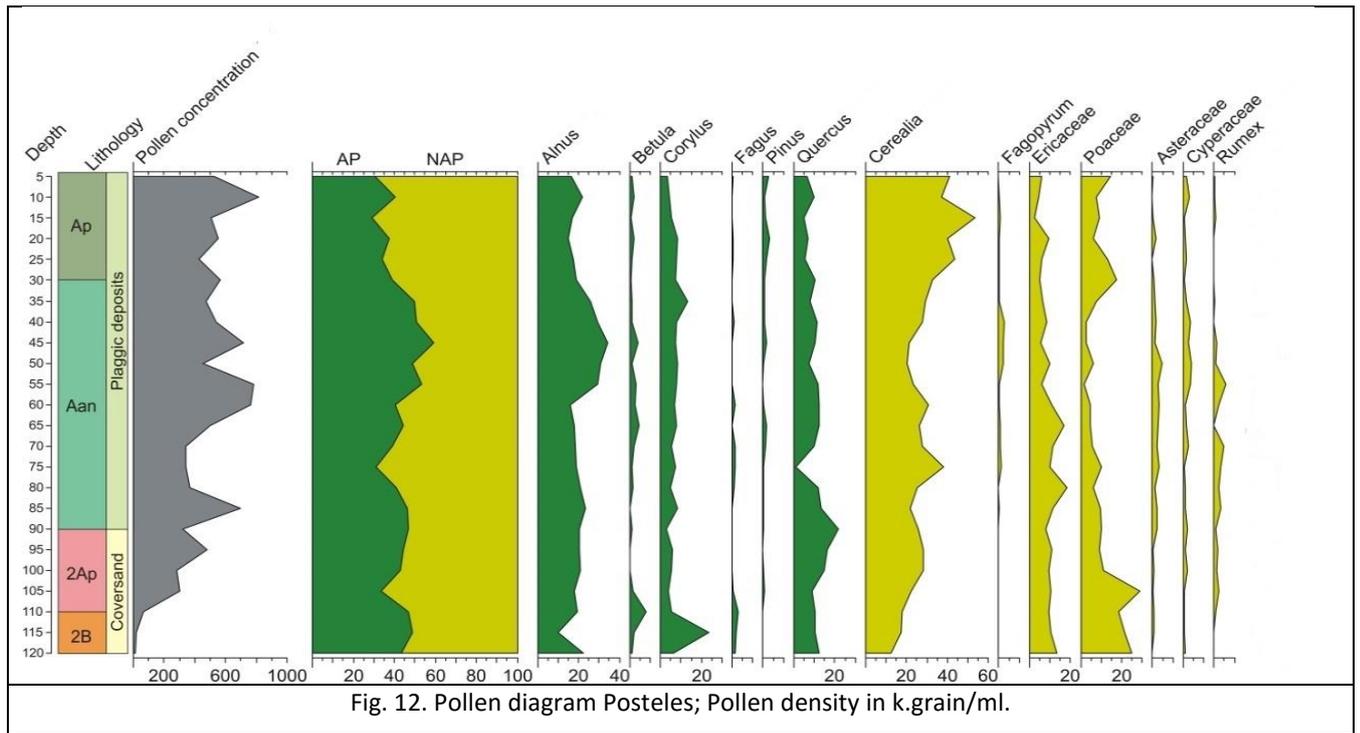


Fig. 11. Biomarker diagram Nabbegat.



642

Table 3. ¹⁴C 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
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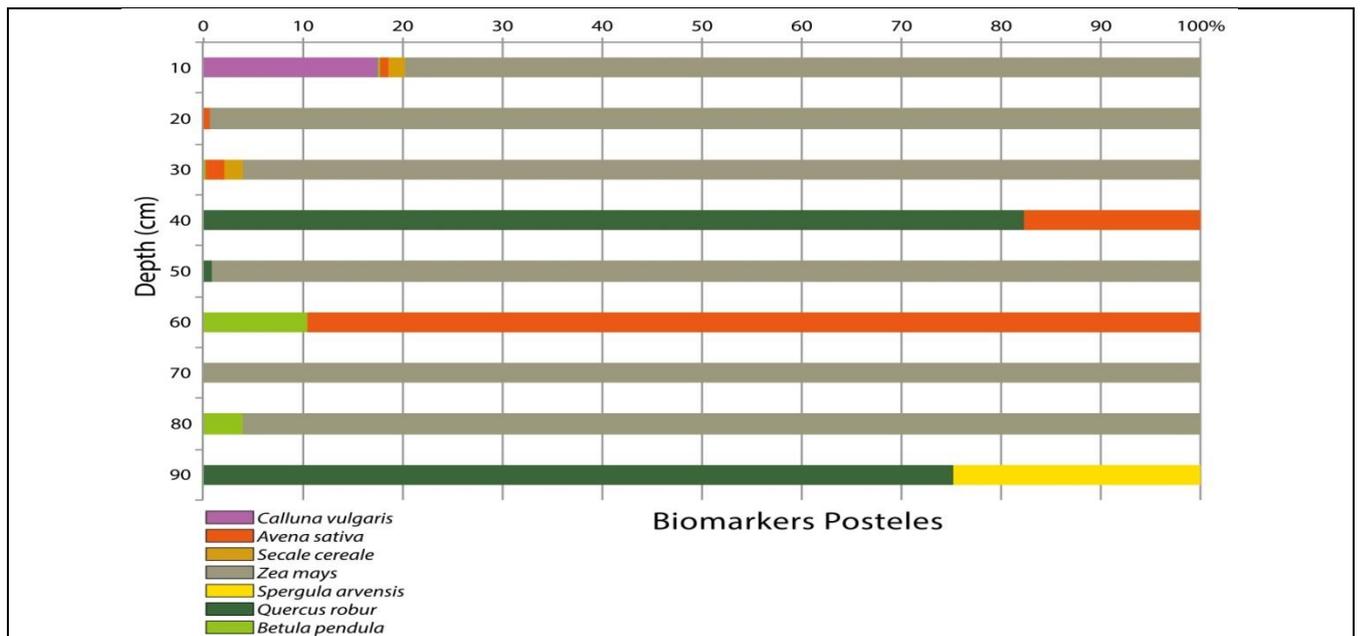


Fig. 13. Biomarker diagram Posteles.

