

Palaeoecological response to Greenlandian (Early Holocene) climatic changes: Insight from an abandoned-channel sequence of the Meuse River at Autrecourt-et-Pourron (Ardennes, France)

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- 20 ABSTRACT
- 21 A Greenlandian (Early Holocene) palaeochannel of the Meuse River is described from
- 22 Autrecourt-et-Pourron in the Ardennes region of northern France. During the Younger Dryas,
- 23 fluvial deposits represent a high-energy, sinuous palaeochannel, but at the onset of the
- 24 Holocene, progressive channel abandonment resulted in the establishment of a low-energy
- 25 meandering river system. Well-dated studies using palynology, carpology, malacology and
- 26 geomorphology reveal a Greenlandian succession of changes in fluvial dynamics and
- 27 vegetation. Between 11,700 and 11,400 cal yr BP, warmer temperatures led to the development
- of a birch community (Betula sp.) within an open grassland, dominated by herbaceous cold-
- 29 adapted plants (Linum alpinum, Plantago montana, Artemisia sp., Juniperus sp.). Between
- 30 11,400 and 11,100 cal yr BP, the river and the vegetation dynamics reflected two short-term

climate events. Around 11,400 cal yr BP, the palaeochannel became separated from the active channels of the Meuse River but remained dominated by waterlogged wetlands. This cold and wet phase represents the "Preboreal Oscillation" also referred as the "11.4 ka event". Around 11,300 cal yr BP, the records suggest that fire frequency increased close to the channel, and may indicate a dry episode that reflects a sharp, short-lived rise in temperature. Between 11,100 and 10,700 cal yr BP (Late Preboreal phase), the channel received flood deposits again. *Salix* and helophytes developed while *Populus* colonised the riverbank. From 10,700 cal yr BP, at the beginning of the Boreal phase, the palaeochannel was finally abandoned and dense temperate woodland with *Corylus avellana* and *Quercus* spp. became established. Based on its high chronological resolution, Autrecourt-et-Pourron represents a key locality in north-western Europe to investigate environmental change in connection with Early Holocene short climate variations. In addition, the results highlight the valuable contribution of carpological analyses in improving the spatial resolution of reconstructions of vegetation cover.

Keywords: Palaeoenvironment; Palaeochannel; Preboreal; Multiproxy investigation; Plant macrofossils; Short climate variation

1. Introduction

Valley floor wetlands from northern Europe can provide valuable palaeoecological archives that preserve information regarding environmental changes in several biotic and abiotic proxies. Such studies have been conducted in Northern France (Antoine, 1997; Antoine et al., 2003b, 2003a; Coutard et al., 2010; Le Jeune et al., 2005; Limondin-Lozouet, 1998; Limondin-Lozouet and Antoine, 2001; Pastre et al., 2003, 1997), in Belgium (Deforce et al., 2014; Meylemans et al., 2013; van der Sloot et al., 2014), in Netherlands (Bos, 2001, 1998; Bos et al., 2018, 2007, 2006, 2005; Bos and van Geel, 2017; Huissteden et al., 1986), and in Poland

(Kołaczek et al., 2018a, 2018b). Holocene warming in northern Europe was prone to several climate fluctuations (Magny, 2019, 2004; Rasmussen et al., 2014a, 2014b). In valley floors, the beginning of the Preboreal was characterised by an intense fluvial activity, the cutting of large palaeomeanders and avulsions. However, an increase in temperature (10±4°C) and precipitation (Hoek and Bos, 2007; Kobashi et al., 2008; Magny, 2013) was interspersed with a brief cooling due to melting ice and a change in the thermohaline circulation. This cooling phase, called the "11.4 ka event", has been observed all over North-western Europe: in the GRIP and NGRIP Greenland ice cores (Björck et al., 1997; Rasmussen et al., 2014b), in French and Swiss lakes (Arbogast et al., 2006; Magny, 2004; Magny et al., 2006; Magny and Ruffaldi, 1995), in Norway (Birks et al., 2000; Birks and Ammann, 2000) and the British Isles (Lowe et al., 1999). In this framework, the French Meuse Valley (Northern France) provides the opportunity to study climate changes and landscape evolution at the beginning of the Holocene (11,700 – 10,700 cal yr BP) especially during the Preboreal. So far, the French part of the Meuse valley (Ardennes region) has had only one palaeoenvironmental investigation published, at Mairy site (Lefèvre et al., 1993). A multidisciplinary approach (morphosedimentary, palynology and malacology) was undertaken in two palaeochannels (My87 and My90), which characterised the Greenlandian dynamics of the sedimentary and biological environments (Lefèvre et al., 1993). In 2012, we had the opportunity to excavate the Autrecourt-et-Pourron site (Ardennes, France), and we detected an extremely thick sedimentological sequence spanning from the beginning of the Holocene at ca. 11,700 up to 10,200 cal yr BP. As a result, this unique high-resolution sequence is shed light on the interactions between climate, vegetation and the river. This paper discusses significant variations in fluvial and environmental dynamics, using a multidisciplinary palaeoecological approach. We aim to describe Greenlandian environmental changes and investigate their links with climate variations. The patterns observed at Autrecourtet-Pourron are compared with regional and global climate trends in order to characterise the

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palaeohydrological and palaeoenvironmental events related to the Preboreal. Special attention is given to the 11.4 ka event and its effects on river dynamics and the local environment.

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2. Study area

The fluvio-palustrine sequence of Autrecourt-et-Pourron (49°36'57,391" N / 5°2'53,781" E/ altitude ca. 155 m a.s.l.) is located in the Meuse Valley, about 30 km upstream from Charleville-Mézières in the Ardennes hilly region (Fig. 1). The Meuse River flows in a northwestly direction across Jurassic (Late-Lias) marly bedrock, following the Jurassic cuesta scarp of the Eastern Parisian Basin. Airborne imagery and LIDAR-DEM from the 2018 copyrighted IGN (Institut Géographique National) resources provide a comprehensive overview of the present-day alluvial landscape, which includes a low sinuosity channel (Canal de l'Est), partly-located on the north-western border of the valley floor and a ~ 2 km-wide floodplain. This floodplain was shaped by both recent meandering planforms and older irregular palaeochannels with narrowing expansion. Such palaeochannels of the PalaeoMeuse river system were incised in a ca. 4 m-thick layer composed of fluvial gravels corresponding to the Pleniglacial valley infilling of Weichselian age (Lefèvre et al., 1993). The study carried out here was undertaken at the same time archaeological excavations (2008 and 2012). Both field surveys uncovered several palaeochannels of the PalaeoMeuse River, with the last field survey in 2012 revealing a cross-section of a well-preserved palaeochannel sequence (Fig. 2). Its continuous vertical succession (Section 3-6) exposed from the bottom of Trench N°6 has allowed us to define stratigraphical units (SU) and undertake continuous sampling for palaeoecological reconstruction.

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3. Materials and methods

3.1. Radiocarbon dating

Samples were directly collected from a 160 cm thick section comprising a fluvio-palustrine deposit with organic horizons. Based on the different lithostratigraphic units and organic material availability, 10 samples of wood and terrestrial plants were dated by the AMS radiocarbon method (¹⁴C) in the Beta-Analytic (US) and Poznan-Radiocarbon (Poland) laboratories (Table 1). Oxcal 4.3 (©Christopher Bronk Ramsey 2018) was used to calibrate the AMS ¹⁴C ages based on the Northern Hemisphere terrestrial IntCal13 calibration curve (Reimer et al., 2013). Calibrated radiocarbon dates are expressed as cal yr BP, while the absolute chronology based on an age-depth model was constructed with a 2σ confidence envelope using the CLAM package (Blaauw, 2010). Linear interpolation between the dated intervals was applied taking into account the main changes in the sedimentation rate in yr/cm (Fig. 3).

3.2. Geochemical analyses

For each sediment sample, the mineral sediment grain-size range was assessed with a laser granularity analyser (Coulter LS 230). Samples were prepared by decarbonation (H_2O_2) and dissolution of the organic matter (HCL). They were first sieved using a 2 mm mesh and afterwards deflocculated with a few millilitres of sodium hexametaphosphate ($Na_6O_{18}P_6$). After homogenisation, the samples were introduced gradually using an eyedropper, and the concentration values were checked. The different granulometric fractions of the sample were measured by laser sensor. The analyser measured, with an accuracy of <1%, the relative proportions of clay (0.04-2 μ m), silt (2-50 μ m) and sand (50-2000 μ m).

Using 100 mg samples of dry sediment, analyses of Total Organic Carbon (TOC) were carried out with a CNS analyser. The TOC content was measured by thermocatalytic oxidation (Elementar, Vario TOC cube analyser).

3.3. Palaeozoological analysis

Within the context of an exploratory analysis, 11 samples were studied to assess the malacological preservation potential. Shells were sieved and sorted by the carpologist. Only the fraction between 0.5 and 2 mm (the most abundant in malacological remains) was studied, which was a bias compared to the methodological standards of malacology (Limondin-Lozouet, 2002). Another bias was that the volume of sieved sediments (between 2 and 4 litres) was insufficient to reach the minimum number of 200 shells required for a malacological analysis (Evans, 1972). Despite these biases, shells were determined in order to estimate the major changes in the composition of the malacofauna within the sequence. This qualitative approach was intended both to estimate the ratio between freshwater and terrestrial species, which is important for assessing the palaeochannel dynamics, and to determine whether species used as biostratigraphic markers on a northern French scale were identifiable at Autrecourt-et-Pourron.

In addition, as part of the exploratory analysis, microfaunal, micromammal and fish remains found in 29 carpological samples were handpicked and counted. Although no identification was undertaken, it seemed relevant to present the total abundances of microfaunal and ichthyofaunal remains as these might also reflect environmental changes.

3.4. Palaeobotanical analysis

Palynological samples were extracted from cores with a sampling interval of 5 cm, and 34 sediment samples of 1 cm³ volume were prepared following standard pollen analytical procedures (Faegri and Iversen, 1989), with chemical treatments (NaOH, HCl, HF, ZnCl₂) followed by acetolysis. Pollen preservation was sufficient and between 250 and 800 pollen grains were counted in each slide. Pollen types were identified using determination keys (e.g. Beug, 2004), pollen reference guides (Moore, 1991; Reille, 1992-1998) and comparison with the reference collection of the Chrono-environnement Laboratory (Besançon, France). The identified species were sorted according to their morphological type (trees, herbaceous) and the

herbaceous taxa were categorised as ecological groups: heliophilous (*Artemisia*, Chenopodiaceae, *Thalictrum*, *Helianthemum*, *Potentilla*, *Plantago*, *Centaurea*, etc.), mesohygrophilous (*Filipendula*, *Equisetum*, *Sanguisorba officinalis*, *Polemonium*, *Lythrum*) and aquatic (*Typha/Sparganium*, *Potamogeton*, *Myriophyllum*).

The 29 carpological samples with a thickness of 5 centimetres and a volume of 10 to 20 litres were extracted continuously on a stratigraphic column 85 cm wide. After sampling, sediments were water-sieved with several meshes (2, 1, 0.5 and 0.25 mm). Sieved macrofossils were fully sorted under a stereomicroscope with a magnification of 2x to 60x. Extracted elements were seeds, bryophytes, buds, vegetative parts (internodes, leaf abscission scars), twigs larger than 2 mm in diameter, whole leaves, needles, and charcoal over 1 mm in diameter. Plant macrofossils were identified using standard specialised literature (Beijerinck, 1976; Berggren, 1969; Cappers et al., 2006; Tomlinson, 1985) and on the basis of the reference collections from the Chrono-environnement Laboratory and GéoArchÉon company. The remains were counted according to the principle that a whole macrofossil equalled one specimen. The fragments were also counted and then converted into an estimated whole (minimum number of individuals) using the «judgmental method» (Pradat, 2015). All data were recorded in the AGAM database (Schaal, 2019a). To measure functional diversity, the ecological requirements (Ellenberg, 1992; Julve, 1998) of species were averaged and weighted by the abundance of fossil remains. As a result, a hierarchical cluster analysis was constructed to categorise groups of samples.

For plotting all the stratigraphic diagrams, the C2 program (©2014 Steve Juggins) was used and we have defined the local pollen assemblage (LPAZ) from the botanical data.

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4. Results

4.1. Lithostratigraphy

The detailed stratigraphic study of the section 3 (Trench N° 6), described in Table 2, showed an alluvial sequence including an abandoned channel-fill, which was observed to be over 2.5 m thick and composed of 22 stratigraphic units (SU). Those units have been grouped into main associations of facies corresponding to Lateral-accretion Deposits (Element LA), Sandy Bedforms (Element SB), Abandoned Channel Fills (Element FF(CH)) and Floodplain Fines (Element FF) sensu Miall (1996). From a sedimentary point of view, the sequence corresponds to contrasting depositional settings related to the fluvial history of a palaeochannel in the Meuse Valley. It is proposed that the basal units (i.e. SU 301, SU 339, SU 341,) were formed in a fluvial bar domain composed of lateral accretion structures (LA) and sand bars (SB). Indeed, we suggest a south-western lateral adjustment of a low to moderate sinuosity stream within a multi-channeled river (i.e. the Palaeo-Meuse River system). According to the age-depth model, we consider that these Late-Glacial fluvial macroforms (LA and SB) were the result of a transitional river system at the expense of a previous braided-plain, as observed over the Weichselian Pleniglacial period (Antoine et al., 2003a, 2003b; Coutard et al., 2010; Limondin-Lozouet and Antoine, 2001; Pastre et al., 2003). Field observations of the SU 326a/b, SU 327, SU 328 and SU 329 support the progressive burial of the preexisting lateral bars by sedimentation of suspended-load deposits with varying amounts of sand (overbank to backswamp deposits). We interpret this fine-grained sediment deposition as the result of channel plugging (CH-FF) controlled by a diminishing flow regime. At the supra-regional scale, the large valleys from the Parisian Basin showed a reduction in the number of channels with generalised clogging and subsequent peat infilling from the Lateglacial (Pastre et al., 2000). At Autrecourt, the floodplain reworking process of the PalaeoMeuse River was characterised by the slow abandonment of its secondary channels followed by vegetated swamp development (CH-FF), with local peat bogs, as observed in SU 330 to SU 332. The upper part of the alluvial sequence was dominated by vertical accretion of floodplain fines (FF), with the occurrence of

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an embryonic humic soil (SU 334), interbedded with clayey silts units (SU 333 to SU 335). The overlying SU 336 is part of 1.5 m-thick subsurface alluvial clay deposits which exhibited reworking evidence, including protohistoric (308-184 BC dendrochronological date on *Quercus* sp.) remains (which were not sampled, and thus not described in this paper). Finally, the current soil (ca. 0.5 m) covered alluvial deposits.

4.2. Microfaunal data

In sedimentary units SU 326, SU 327 and SU 328, an average of 5 fish remains/litre (scales and vertebrae) were found (Fig. 4). In SU 329, fish remains were scarce. In SU 330, few fish remains were recorded while an average of 6/litre of micro-mammal remains were found. No fish remains were observed in SU 331, SU 332 and SU 333.

Malacological samples provided assemblages that reflected two environmental phases. First, in SU 326, terrestrial molluses were dominant (Table 3). Almost half of the specimens belonged to a single species: *Trochulus hispidus*. Besides this mesophile species, the openground species *Pupilla muscorum* and the marshland species *Oxyloma elegans* were the most common. In addition, two terrestrial species indicative of cold environments, were listed during this first environmental phase: *Pupilla alpicola* and *Columella columella*, which are mainly restricted today to boreo-alpine areas (Welter-Schultes, 2012). The list of freshwater species, with *Pisidium subtruncatum* as the main taxon, suggested permanent water in the palaeochannel (Welter-Schultes, 2012). In SU 327 and SU 328, terrestrial molluses virtually disappeared. The species composition of freshwater molluse assemblages changed with *Valvata macrostoma*, characteristic of periodic pools in floodplain contexts (Horsák et al., 2013), which was observed as the main taxon. The upper sedimentary units were devoid of any malacological remains, probably because of the acidity of peat deposits, which prevented the development of molluses and/or the preservation of their shells.

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4.3. Palaeobotanical data

In SU 326, SU 327 and SU 328 (local pollen assemblage zone LPAZ AUT.1), NAP values (> 60%) are high, which suggest an open landscape, dominated by Poaceae and heliophilic taxa (Artemisia, Thalictrum, Helianthemum and Chenopodiaceae). Pinus (20% of records) and *Betula* (not exceeding 10%, with a peak in layer SU 327) represent woodland (Fig. 5). Taxa from wet meadows are also well represented with Cyperaceae (30%) and Filipendula (20 %). The carpological diagram (Fig. 6) specifies the composition of the herbaceous assemblage with some taxa representative of cold climates such as Linum alpinum, which is found today at high altitudes (Alps Mountain). Carpological results also give accurate information about ecology, particularly for mesohygrophilic and aquatic plants. The plant macrofossil assemblage is composed of some algae (Nitella sp.), submerged plants (Myriophyllum spicatum) and floating leaf plants (Potamogeton sp.). Hygrophytes and seasonal amphibious vegetation complement the aquatic spectrum (Carex rostrata, Rorippa palustris and Phragmites australis). Riparian and wet meadow plants also occur (Carex elata, Filipendula ulmaria, Geum rivale, Viola cf. palustris, Stellaria palustris and Peuceudanum officinale). Some weed species (Oxybasis rubra, Urtica dioica and Ranunculus repens) and Cenococcum geophilum sclerotia have been found. In the carpological diagram, trees are represented by macrofossils of *Betula* sp. and *Salix* sp. (seeds and buds).

In SU 329 (LPAZ AUT.2), the percentage of *Pinus* pollen increases slightly, while *Betula* pollen and macrofossils decreased (Figs. 5 and 6). At the same time, oogonia from *Chara* algae sharply increase, as well as the seed densities of *Potamogeton, Carex rostrata, Rorippa palustris* and *Veronica longifolia*. The riparian vegetation is imperceptible and taxa from meadow environments decline.

In SU 330 (LPAZ AUT.3), numerous wood charcoal fragments (between 100 and 200 > 1mm per litre of sediment) were recovered. Carbonised seeds of wetland plants, such as *Carex*, were also found with an average of 5 seeds per litre. Macrofossils of aquatic plants almost disappeared whereas seasonal amphibious and hygrophilous plant remains such as *Sparganium erectum, Carex elata, Filipendula ulmaria, Ranunculus repens* and *Thalictrum flavum* increased. The first riparian forest indicators are *Stachys sylvatica, Prunus spinosa* and *Rhamnus cathartica*, while at the top of this pollen zone the end of the *Juniperus* phase is recorded.

In SU 331 (LPAZ AUT.4), charcoal and carbonised seeds were still recovered but in far less abundances (Fig. 4). *Pinus* pollen reaches its maximum percentages between 40 and 60% (Fig. 5). The main components of the aquatic environment change with *Myriophyllum spicatum* and *Ranunculus aquatilis* agg. being recorded as the dominant species. Helophytic vegetation decreases despite the presence of *Sparganium erectum* seeds. Riverside vegetation with *Salix, Solanum dulcamara* and *Filipendula ulmaria* expand, while taxa indicative of alluvial forest develop (*Prunus spinosa, Rhamnus cathartica* and *Origanum vulgare*).

In SU 332 (LPAZ AUT.5), high percentages of *Betula and Salix* pollen and macrofossils characterize this phase, while Cyperaceae and *F. ulmaria* gradually decline (pollen and seeds). High abundances of *Hippuris vulgaris* and *R. aquatilis* agg. correlate with the aquatic environment of the channel. The riparian vegetation shows a diverse spectrum including *Lycopus europaeus* and *Valeriana officinale*.

SU 333 (LPAZ AUT.6) is marked by a *Pinus* decline (13%) and the development of *Corylus*, comprising 45% of the pollen assemblage, together with the appearance of mesothermophilic taxa, such as *Quercus* (2.5%) and *Ulmus* (2%). Forest herbs such as *Origanum vulgare, Stachys sylvatica* and *Valerianella dentata* are also important. This last phase is highlighted by a strong decrease in Cyperaceae (3%) and mesohygrophilous taxa (e.g.

Filipendula and Equisetum). With the presence of Ranunculus aquatilis and Hippuris vulgaris in both palynological and carpological diagrams, increasing drier conditions can be reconstructed within the channel. The occurrence of Lycopus europaeus and Eupatorium cannabinum indicate a wet meadow environment in its vicinity.

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5. Discussion

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5.1. Palaeoenvironmental succession at Autrecourt-et-Pourron

The phase AUT.0 was defined from the current landforms and the lithostratigraphy since no samples were taken within the gravels and sands of the excavation floor (SU 301, SU 339 and SU 341). The depositional settings observed at the base of the section 3-6 are characterised by lateral bar deposits and sandy bedforms recorded along the course of a narrow channel (ca. 5 to 10 m-width). Thus, it appears that this palaeochannel probably occurred in a relatively stable multi-channeled floodplain as a wandering gravel-bed river controlled by both high hydrodynamic conditions and lateral accretion processes. It is assumed that this transitional river pattern, including several channels with such geometry, incised the underlying pleniglacial gravels. An age of between 12,377 and 11,626 cal yr BP is supported by two radiocarbon measurements on waterlogged wooden twigs (Table 1). This time-span covers the Younger Dryas period, reported in the GRIP isotope record as the last cold phase before the postglacial warming (Ammann et al., 2007; Rasmussen et al., 2014b). A succession of silt, clay and peat dated between c. 11,700 and c. 10,200 cal yr BP, then slowly filled the channel with the low sedimentation rate being of an oxbow lake type (Fig. 3). The phases established from the agedepth model were very accurate in comparison with other alluvial plain or down slope records. Six environmental phases (AUT.1 to AUT.6) were defined in this time frame spanning the first half of the Greenlandian.

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5.1.1. AUT.1: a cold steppic environment recording backswamp deposits

The phase AUT.1 covers sandy and clayey silts (SU 326, 327 and 328) deposited in low sedimentation dynamics. In the pollen assemblages, herbaceous plants such as Artemisia and Helianthemum are dominant and, in the carpological assemblages, taxa from basophilic grassland vegetation (such as Linum alpinum, Scabiosa columbaria and Knautia arvensis) show occurrences restricted to the phase AUT.1. The record of Linum alpinum in SU 326 is of significance in terms of palaeoclimatic interpretation, as this species only grows today in subalpine areas, above 1500 m a.s.l. (Julve, 1998). In the whole SU, the presence of coldtolerant molluscs in an assemblage dominated by Trochulus hispidus and the absence of thermophile species from woodland environments are consistent with such a reconstruction of cold climate conditions. In SU 327, the decline or disappearance of these cold-tolerant plants and molluscs suggests warmer conditions c. 11,600-11,500 cal yr BP. In the carpological analysis, *Pinus* is absent, as is *Juniperus*, which are both adapted to poorly developed and thin soils, as probably occurred on the valley's slope. Meanwhile Salix is the dominant tree, as it grew directly in the vicinity of the palaeochannel. In addition, characteristic plants of eroded sandy and stony soils (such as Arenaria serpylifolia and Linaria supina) together with the recorded fungal species Cenococcum geophilum, indicative of short environmental stress (Eide et al., 2006; Kroll, 1988; van Geel et al., 1989; Walker et al., 2003), suggest erosion of soil and organic matter in the floodplain. Sporadic fluvial inputs observed as overflow facies in SU 327 and SU 328 are in agreement with such biological features. The increasing proportions of the freshwater mollusc Valvata macrostoma, indicative of standing water, would be consistent with a decline in hydrodynamics. The notable abundances of *Urtica dioica*, *Chenopodium rubrum*, Ranunculus repens and Prunella vulgaris might suggest the presence of large herbivores (Bos et al., 2006) since coprophilous fungi (Sordariaceae and Sporormiella), which could be

indicators of grazing pressure (Gauthier et al., 2010; van Geel and Aptroot, 2006), have been observed in the pollen samples (E. Gauthier, personal communication). The slow change in vegetation in the Greenlandian at Autrecourt-et-Pourron might have favoured the Lateglacial megafauna, which eventually disappeared from the region with the subsequent disappearance of the steppes.

5.1.2. AUT.2: a well-vegetated abandoned channel-fill under cold conditions

The phase AUT.2 covers SU 329. *Pinus* forests developed gradually while a decline of *Betula* sp. was observed in both the palynological and carpological diagrams, suggesting a greater distance from wooded areas in the valley. A decrease in *Salix* sp. macro-remains was also detected in the floodplain, while Poaceae steppe, *Thalictrum* and *Filipendula* wet meadows were significantly well-represented. Plant macrofossils assemblages were strongly dominated by *Chara* oogonia and *Carex rostrata*, indicative of aquatic environments. This sudden change in the vegetation composition together with the development of organic mud in SU 329 suggest that channel abandonment was occurring away from the active river system. A water depth of two metres could be estimated on the basis of the formation of calcium carbonate encrusting the *Chara* oogonia (Pelechaty et al., 2013). Furthermore, the presence of aquatic and helophytic plants, such as *Myriophyllum spicatum* and *Carex rostrata*, indicate slightly colder average temperatures than during the end of phase AUT.1. The list of plant taxa has enabled an estimation of the minimum mean July temperatures to be between 8 and 10 °C (Bos et al., 2007).

5.1.3. AUT.3: development of marshland environment and fire activity

Whereas in the silts (SU 326 to SU 329), the moisture content was rather stable and provided a suitable environment for the preservation of organic material, the plant macrofossils in the peat layers (SU 330 to SU 332) were more deteriorated (Schaal and Naton, submitted).

This taphonomic bias, even if low, should be carefully considered in palaeoecological interpretations.

This environmental phase AUT.3 covers peats from SU 330. In the carpological diagram, aquatic plants from AUT.2 virtually disappeared and were replaced by plants from marshland environments, especially *Carex elata, Filipendula ulmaria* and *Thalictrum flavum*. The numbers of plant macrofossils from arboreal taxa were low. In addition, the biodiversity of terrestrial grasses sharply decreased. In the pollen diagram, at the transition between AUT.3 and AUT.4, *Pinus* sp. decline while *Betula* sp. and *Salix* sp. increase in proportion. This decrease in *Pinus* sp. was identified in only one sample. However, it was observed shortly before the disappearance of *Juniperus* in the pollen diagram and in connection with a peak in charcoal and carbonized seeds (Fig. 4).

5.1.4. AUT.4: a riparian vegetation development in an abandoned channel-fill

In SU 331, described as dark peat, the specific composition of marshland plants changed. In the carpological diagram, *Carex elata* and *Thalictrum flavum* were replaced by *Sparganium erectum* and *Filipendula ulmaria* as the main species. Large helophytic plants with rhizomes, such as *Typha* and *Sparganium*, developed in connection with the nutrient-rich peat. The proportions of arboreal pollen increased in the valley and, in the floodplain, *Salix* sp. showed a higher density of macro-remains. On the basis of their ecological requirements, the appearances of the tree species *Prunus spinosa* and *Rhamnus cathartica*, and the herbaceous forest plants *Stachys sylvatica* and *Luzula* sp. were favoured by the soil texture and its organic-rich content (Ellenberg, 1992; Julve, 1998). The abandoned channel was colonised by trees and grasses that composed the riparian vegetation.

5.1.5. AUT.5: development of an alluvial forest and hydrodynamic changes

This phase was defined according to the palaeobotanical assemblages of SU 332. *Betula* sp. and *Salix* sp. showed higher proportions in the pollen diagram. In the carpological sequence, *Betula* sp. was barely noticeable, while a high density of *Salix* sp. macro-remains was observed together with the first occurrence of *Populus*. The results reveal that an alluvial forest developed in the floodplain. Changes in the composition of the aquatic community (*Ranunculus aquatilis* and *Hippuris vulgaris* as the main species) provide evidence of hydrodynamic fluctuations. Therefore, we conclude that the abandoned channel was flooded by water estimated at 40 to 100 cm-depth. These wetter and more anoxic conditions favoured the conservation of the organic remains. The occurrence of *Typha latifolia* in the pollen assemblage is of climatic significance since this species requires minimum mean July temperatures of 13°C to develop (Bos et al., 2007).

5.1.6. AUT.6: a temperate woodland in an alluvial plain setting

During AUT.6, which covers SU 333, mesothermophilic taxa (*Corylus*, *Quercus*, *Ulmus*) characteristic of the temperate woodland develop, while *Pinus*, *Betula* and *Salix* decline. Heliophilous taxa (*Artemisia*, *Thalictrum*, *Helianthemum* and Chenopodiaceae) are virtually absent. The wetland considerably changed as well, with a mesohygrophilous community of *Eupatorium cannabinum* replacing the wet meadow composed of Cyperaceae. The swampy area shrank and only a few patches of sedges remained. The alluvial sedimentation also changed with massive clay deposition as observed in SU 333. These facies settled out from suspension in a backwater area of the floodplain, thus in the distal part of the fluvial distributary system. This upper layer of grey clay was strongly subjected to erosional factors, resulting in the carpological material being most likely moved and reworked, causing fragmentation and even destruction. Given the absence of fossil remains of aquatic plants and ichthyofauna, it can

be assumed that the channel was definitely abandoned (Bogaart et al., 2003; Martín-Closas and Gomez, 2004).

5.2 Palaeoenvironmental links with regional reference series

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407 5.2.1. A useful record of the first centuries of the Holocene (11,700-11,400 cal yr BP)

At Autrecourt-et-Pourron, the base of the alluvial sequence was dated from the Younger Dryas phase of the Lateglacial on the basis of radiocarbon dating. A multi-channel floodplain environment was proposed based on the evidence of a sinuous and a gravel-bed palaeochannel deposit (Fig. 7A). The Younger Dryas period was characterized by glaciers advancing, significant temperature reduction (below -20° C) and the replacement of pine forests by herbaceous steppe (Bos et al., 2018; Kasse et al., 2017; Renssen, 2001). The PalaeoMeuse River has evolved towards a low-energy fluvial system from the onset of the Holocene. Based on the section 3-6, fluvial sedimentation tends to be finer-grained than the sediment found with the preexisting channel. In numerous alluvial series from northern France (e.g. Antoine et al., 2000; Le Jeune et al., 2005; Orth et al., 2004; Pastre et al., 1997; Guélat and Richard, 2014) and northwestern Europe (e.g. Bogaart et al., 2003; Tebbens et al., 1999; Vandenberghe, 2002), the first centuries of the Holocene were marked by an incision phase. This erosive period was not recorded in the section 3-6 of Autrecourt-et-Pourron. In the Netherlands, while the transition between the Younger Dryas and the onset of the Holocene (Friesland phase sensu Behre, 1966) was reported as dating from c. 11,500 cal yr BP (Bos et al., 2007; Bos and van Geel, 2017), the Lateglacial/Holocene transition was interpreted as occurring around 11,700 cal yr BP in the GRIP and NGRIP Greenland ice cores (Björck et al., 1997; Rasmussen et al., 2014b). Autrecourt-et-Pourron thus provides a valuable insight of the transition between the Lateglacial and Holocene conditions in the Meuse valley. The cold-tolerant plants (such as *Linum alpinum*) and molluscs (such as Columella columella), observed at the base of phase AUT.1, are likely relics of colder Lateglacial conditions.

In the Dutch section of the middle Meuse River valley, the preservation of various biological archives in Younger Dryas deposits have enabled an accurate environmental reconstruction of the composition and distribution of vegetation (synthesised in Bos et al., 2007). Whether the absence of biological samples in the Lateglacial coarse sediments from Autrecourt-et-Pourron precludes any paleoenvironmental discussion, an open landscape most likely developed. One could assume that the relative instability of the river might have formed patches of gravel bars in the floodplain, while small-vegetated mid-channel islands developed between the wandering channels. With the warming temperatures of the Greenlandian and the changing style of the river, heliophilous herbs and trees became far more widespread in the floodplain (phase AUT.1, Fig. 7B). In the Netherlands and Belgium, the Friesland Phase was characterised by the full development of Betula woodland, while the open herbaceous vegetation of the Younger Dryas became scarce (Bos et al., 2007; van Geel et al., 1980). This phenomenon was not fully perceived in the carpological assemblages from Autrecourt-et-Pourron since Betula remains were quite low in comparison with the Dutch plant assemblages. It can be assumed that different edaphic conditions or geomorphological settings may explain the relative scarcity of Betula at Autrecourt-et-Pourron.

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5.2.2. Highly fluctuating humidity and temperature conditions over a short period (11,400-11,100 cal yr BP)

At Autrecourt-et-Pourron, a sudden change was observed in the lithostratigraphy with the occurrence of peats c. 11,400 cal yr BP (Fig. 3). The development of these organic-rich sediments was observed in connection with a decline of *Betula*, an increase in Poaceae and the appearance of taxa from cold (*Ephedra*) and wet (*Equisetum*) environments during the phase AUT.2 (Fig. 5). In the phase AUT.3, despite rather poor conservation of plants macro-remains, a decline of aquatic plants was observed (Fig. 6) together with an increase in charcoal (Fig. 4), suggesting that a dry phase occurred c. 11,300 cal yr BP. One could hypothesize that the drying

of the valley bottom induced an increase in fire activity at the transition between phases AUT.3 and AUT.4. In Germany, an increase in fires has been reported as being connected with the presence of Mesolithic settlements (Bos and Urz, 2003). However, at Autrecourt-et-Pourron, there is no evidence to distinguish whether the fires were of natural or anthropogenic origin. Nevertheless, the drier conditions reconstructed around 11,300 cal yr BP were *per se favourable* to fires, as observed elsewhere in Europe (Doyen et al., 2015; Rius et al., 2014; Tinner and Lotter, 2001). During the first part of the phase AUT.4, marshland vegetation developed, but trees still showed poor assemblages until c. 11,100 cal yr BP (at 250 cm in Fig. 3).

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This short period, spanning 200 years, was thus marked by highly fluctuating humidity and temperature conditions. Each of these environmental phases lasted a few decades, or a century at the most.. In the Netherlands, this period has been considered as the Rammelbeek phase (between c. 11,400 and c. 11,250 cal yr BP), which was characterized by a decrease in Betula in favour of herbaceous steppe assemblages (Behre, 1978; Bos et al., 2007; van Geel et al., 1980). As a result, these changes in the plant cover led to the reconstruction of a dry climate phase with hotter summers and colder winters. Indeed, Autrecourt-et-Pourron provides a detailed and original record, far more complex than previously described for this period. The cold and wet phase described in AUT.2 (Fig. 7C) would be coincident with the "Preboreal Oscillation" (PBO), which was characterised by a gradual cooling over 100-150 years. This cold event, reported as the "11.4 ka event", was widely observed in the GRIP and NGRIP Greenland ice cores (Björck et al., 1997; Rasmussen et al., 2014b), in French and Swiss lakes (Arbogast et al., 2006; Magny, 2004; Magny et al., 2006; Magny and Ruffaldi, 1995), in Norway (Birks et al., 2000; Birks and Ammann, 2000), and in the British Isles (Lowe et al., 1999). The dry phase observed in AUT.3 and AUT.4 (Fig. 7D) would be coincident with an abrupt warming of 4 ± 1.5 °C observed in the Greenland ice cores (Kobashi et al., 2008). The authors suggested that the climate transition could have occurred within a few years given the

rapid increase in snow accumulation, and they compare this sudden warming with the Dansgaard–Oeschger (D/O events), which consisted of several abrupt warming phases that took place in a matter of decades (Bond et al., 2013).

Based on this high chronological resolution, Autrecourt-et-Pourron has become a regional key sequence in north-eastern France to investigate these Greenlandian short climate oscillations.

5.2.3. The wet Late Preboreal phase (11,100-10,700 cal yr BP)

At Autrecourt-et-Pourron, AUT.5 was marked by the development of alluvial forest and a new fluvial input in the palaeochannel (Fig. 7E), both indicative of wetter conditions. Our French series closely corresponds with the Dutch records from the Late Preboreal (11,250-10,700 cal yr BP), which was considered as a wet climate phase (Bos et al., 2007; van der Plicht et al., 2004). In addition, the appearance of *Populus* in AUT.5 is consistent with its colonisation during the Late Preboreal phase further downstream.

5.2.4. Temperate forest development of the Boreal phase

At Autrecourt-et-Pourron, the vegetation composition sharply changed in AUT.6, especially with the development of temperate woodlands (Fig. 7F). The development of *Corylus avellana* woodland, inferred from it being the main tree species in the pollen assemblage appeared as a strong marker of the Boreal phase in north-western Europe (e.g. Giesecke et al., 2011). In the Netherlands, the appearance of *Corylus avellana* was dated c. 10,700 cal yr BP (Bos et al., 2007). Similarly, the modelled age of the appearance of *Corylus avellana* at Autrecourt-et-Pourron is consistent with such an assumption (transition between SU 331 and SU 332 in Fig. 3). In addition, as in the Dutch Meuse River section, *Quercus* and *Ulmus* appeared shortly after *Corylus* (Fig. 5).

At the onset of the Boreal, the palaeochannel was reduced to a very shallow stream with stagnant water. In the French Ardennes, paleoenvironmental studies in fluvial contexts show significant hiatus for the Boreal phase (Lefèvre et al., 1993). As such, well-preserved sequences are still lacking to better characterise the fluvial dynamics and palaeoenvironmental development during this phase.

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6. Conclusions

A multiproxy investigation of a Greenlandian palaeochannel located in north-eastern France has provided a good reconstruction of the river dynamics and environmental changes in the Meuse River. The Meuse River showed several narrow and sinuous channels over the Late Glacial, probably flowing in a mobile gravel-bed river system. At the onset of the Holocene, such a depositional setting, based on the well-developed sequence of Autrecourt (section 3-6), recorded a slowly infilling sequence before leaving behind an abandoned and vegetated channel on the floodplain. Three steps are proposed as the response of the Preboreal climate variations: 1. The Early Preboreal (11,700-11,400 cal yr BP): silty inputs characterise the beginning of the Preboreal at 11,700 cal yr BP. Subsequently, the flow regime of the channel slowly declined for 300 years. Aquatic plants colonised in and around the area of the channel, which became marshy. The speed of the current was slow enough for floating vegetation to colonise, which suggests calm conditions. The slow infilling dynamics were interrupted by the "11.4 ka event". 2. The "11.4 ka event": during this very brief phase, lasting less than a century, the temperature dropped sharply by 4°C. At the same time, groundwater levels remained very high. The water in the channel remained deep and calm and the lowland vegetation continued to grow. Separated from the main active river, the water-saturated area was covered with floating vegetation. Amphiphytes, which benefit from high water levels, colonised. The "11.4 ka event" was immediately followed by a rapid rise in temperature at around 11,300 cal yr BP, and channel cut-off occurred suddenly. During this phase, lasting about a century or less, the water table level decreased, the land dried up, and the wet vegetation disappeared and gave way to extensive grasslands. As a result, these large areas of dry grass were able to catch fire

3. The Late Preboreal (11,100-10,700 cal yr BP): after these two oscillations of cold and warm conditions, the vegetation dynamics were unstable for a century. For the next four hundred years, the relatively high water table allowed aquatic and riparian vegetation to colonise and the abandoned channel had a new fluvial input. Nevertheless, during the Boreal (10,700/10,600 cal yr BP), the channel was abandoned.

According to studies in the southern Netherlands, Bohncke et al. (1991) concluded that during the Preboreal the vegetation phases were not so much related to thermal phenomena, but to humidity variations and water conditions governing the observed palynological changes. The carpological analysis of Autrecourt-et-Pourron not only confirms this hypothesis but also adds detail to it as plant macrofossils provide information for characterizing river flows and water levels in palaeochannels. The data presented in this study shows that the identification of a large number of wetland species is a major step for improving the understanding and accuracy of past climate and hydrological phenomena in North-East France (Schaal, 2019b), but also may be useful for predicting future patterns.

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Tables and figures

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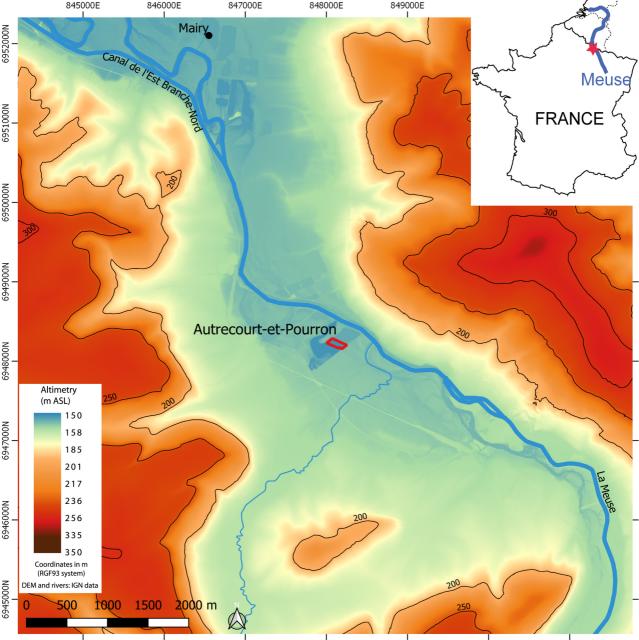
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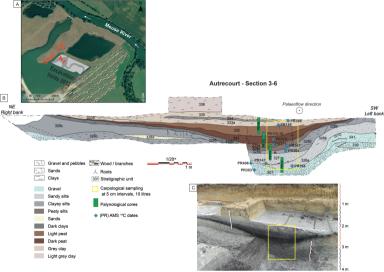
Table 1. AMS radiocarbon dates, depth and dated material of Autrecourt-et-Pourron section 820 821 3-6 as indicated in Fig. 2. The shaded line was not considered in in the age model. Table 2. Stratigraphy of the channel-fill sequence of Autrecourt (section 3-6). 822 Table 3. Molluscs: list of species and number of specimens. 823 824 Fig. 1. Location of Autrecourt-et-Pourron sites (red rectangle), IGN-France map with altitude. 825 Fig. 2. Section 3-6 stratigraphy based on 2012 excavations at Autrecourt-et-Pourron. A -Location of 3-6 profile on the Meuse River valley. **B** - Detailed Cross-section 3-6 of 826 827 Autrecourt-et-Pourron. 301: Alluvial plain with overbank fines deposits and pedogenesis, 339: Sandy gravels, 341: Gravelly sands with calcareous granules, 326a: Homogeneous grey 828 sandy silts, 326b: Grey Sandy silts, 327: Grey sandy silts and with wood debris, 328a&b: 829 830 Light grey clayey silts with sandy beds, 328c: Light grey clayey silts, slightly sandy, 328d: Homogeneous yellowish sands, 329a: Brown silts, slightly organic and bioturbated, 329b: 831 832 Grey clayey silts, 329c: Mottled clayey silts with root traces, 330a: Light-brown peat, 330b: 833 Grey clayey silts, 331: Dark-brown peat, 332: Light-brown peat affected by desiccation cracks topward, 333a: Grey to greenish-grey clay, 333b: Mottled light clay with sandy beds, 834 334: Dark-grey humic horizon, 335: Gray clay, 336: Light-grey clay with hydromorphic 835 features. C – Section's picture, yellow rectangle = sediment sampling area. 836 Fig. 3. Age-depth model in calibrated yr BP and lithology of the Autrecourt-et-Pourron, 837 section 3-6 sequence. Radiocarbon dates (Table 1) were calibrated using Oxcal 4.3 838 (©Christopher Bronk Ramsey 2018) based on the Northern Hemisphere terrestrial IntCal13 839 calibration curve (Reimer et al., 2013). The absolute chronology based on an age-depth model 840 was produced with a 2σ confidence envelope using the CLAM package (Blaauw, 2010). The 841

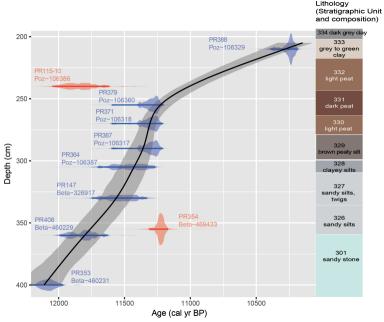
Bayesian age-depth model is presented with its median (middle grey line) and its associated

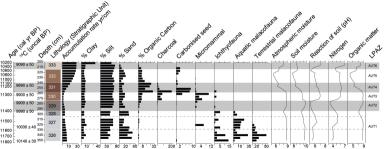
95 % confidence interval (grey area). The calibrated age range of dated samples is indicated 843 in blue and the dates excluded from model in red (outlying). Two dates were rejected after 844 age-depth modelling and outlier analysis, primarily because of age reversals. 845 Fig. 4. Diagram with grain size (%), organic carbon (%), charcoal (number/litre), carbonized 846 seed (number/litre) and aquatic and terrestrial fauna (number/litre) data. Ecological values 847 (moisture, reaction, nitrogen and organic matter), of the identified taxa at the species level, 848 were averaged and weighted by the abundance of plant fossil remains. Depth below ground 849 850 level. LPAZ: Local Pollen Assemblage Zone. Fig. 5. Simplified pollen diagram (%). Depth below ground level. Exaggeration curve x5. 851 LPAZ: Local Pollen Assemblage Zone. 852 Fig. 6. Carpological diagram showing the abundances of plant remains/litre of sediment. 853 Depth below ground level. LPAZ: Local Pollen Assemblage Zone. 854 855 Fig. 7. Conceptual diagram of palaeoenvironment dynamics of the Autrecourt-et-Pourron

channel during the Early Holocene. Exaggerate scales.

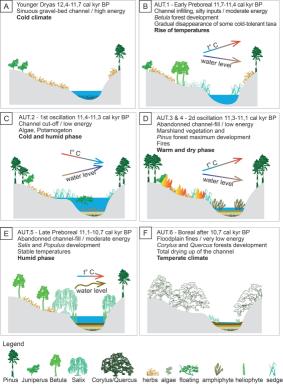












Autrecourt sample	Depth cm	Laboratory N°	Dated material	δ ¹³ C (‰)	Age ¹⁴ C yr B.P. without d13C correction	Conventional Age ¹⁴ C yr B.P. with d13C correction	Calibrated age (cal yr B.P. 2σ) INTCAL13 (95.4%)	Calibrated age (cal yr B.P. 2σ) INTCAL13 (68.2%)	Calibrated age (cal yr B.C. 2σ) INTCAL 13 (95.4%)
2018-PR388-US333	210	Poz-106329	waterlogged <i>Eupatorium cannabinum</i> seeds (174 fgt)		9,090 ± 50		10,396 - 10,182	10,271 - 10,200	8,447 - 8,233
2012-PR148-US333a	215	Beta-326918	waterlogged root	-29.3	4,750 ± 40	4,680 ± 40	5,577 - 5,5315	5,467 - 5,323	3,628 - 3,366
2018-PR115-10-US332	240	Poz-10386	waterlogged Hippuris vulgaris (4) + Carex (1) seeds		10,160 ± 50		12,053 - 11,511	11,974 - 11,750	10,104 - 9,562
2018-PR379-US331	255	Poz-106360	waterlogged Thalictrum flavum seeds (20)		9,890 ± 50		11,597 - 11,202	11,336 - 11,230	9,648 - 9,253
2018-PR371-US330a	270	Poz-106318	waterlogged Thalictrum flavum seeds (20)		9,900 ± 50		11,600 - 11,208	11,385 - 11,236	9,651 - 9,259
2018-PR367-US329	290	Poz-106317	waterlogged Carex rostrata seeds (30)		9,900 ± 50		11,600 - 11,208	11,385 - 112,36	9,651 - 9,259
2018-PR364-US328	305	Poz-106387	waterlogged Carex seeds (20)		9,990 ± 50		11,705 - 11,263	11,603 - 11,322	9,756 - 9,314
2012-PR147-US327	330	Beta-326917	charred wooden twigs	-28.2	10,030 ± 40	9,980 ± 40	11,620 - 11,264	11,620 - 11,398	9,671 -9,315
2017-PR354-US326	355	Beta-469433	waterlogged Linum alpinum seeds (3)	-26.0	9,810 ± 50	9,790 ± 50	11,287 - 11,126	11,242 - 11191	9,338 - 9,177
2012-PR406-US326a	360	Beta-460229	waterlogged wooden twigs	-27,6		10,140 ± 30	11,998 - 11626	11,941 - 11,715	10,049 -9,977
2012-PR353-US303	400	Beta-460231	waterlogged wooden twigs	-27.9		10,310 ± 30	12,377 - 11,962	12,156 - 12,011	10,428 - 10,013

Stratigraphic units (SU)	Description (field & grain-size)	escription (field & grain-size) Thickness (in m.)					
336	Light-grey clay with hydromorphic features		(depositional setting) Alluvial plain with overbank fines				
335	Grey clay	0.05	deposits and pedogenesis (humic				
334	Dark-grey humic horizon	0.85	horizon and redoxic soil) (FF : floodplain				
333 a	Grey to greenish-grey clay		fines sensu Miall, 1996)				
333 b	Mottled light clay with sandy beds						
332	Light-brown peat affected by desiccation cracks topward	0.3	Vegetated swamp deposits with Histosol development (3 soils) (CH-FF sensu				
331	Dark-brown peat (reworked to the North-East direction)	0.22	Miall, 1996)				
330 a	Light-brown peat						
330 b	Dark-grey clay (progressive transition with 330a)	0.12					
329 a	Brown silts, slightly organic (peat) and bioturbated	0.2	Vegetated swamp deposits (a) with				
329 b	Grey clayey silts	0.2	overbank fines (b) and incipient soil (c) (CH-FF sensu Miall, 1996)				
329 c	Mottled clayey silts with root traces		(CITTT SCISO Wildin, 1330)				
328 a & b	Light grey clayey silts with sandy beds						
328 c	Light grey clayey silts, slightly sandy		Backswamp to overbank deposits in a				
328 d	Homogeneous yellowish sands	0.5	channelized area (CH-FF sensu Miall,				
327	Grey sandy silts and with wood debris		1996)				
326 a	Homogeneous grey sandy silts						
326 b	Grey Sandy silts						
341	Gravelly sands with calcareous granules		Lateral accretion macroform & cross-				
339	Sandy gravels		bedded silty sands (Lateral accretion:				
301	Cross-bedded polygenic gravels with silty sand beds organized in fining upwards (Clast size: 2 – 50 mm, subrounded clasts, moderately sorted, openwork texture and normal grading)	1 min.	LA; Sandy bedforms: SB sensu Miall, 1996)				

	Sedimentary units	SU326				SU327					SU328	
	Taxa/Samples	Pr354	Pr355	Pr356	Pr357	Pr358	Pr359	Pr360	Pr361	Pr362	Pr364	Pr365
	Clausilia rugosa	3	2	12	6	3	4	2				1
	Cochlicopa lubrica		1	3								
	Columella columella		1									
	Euconulus fulvus			1								
	Nesovitrea hammonis		2				1					
	Oxyloma elegans	4	2	23	8	4				1		
Terrestrial	Punctum pygmaeum			1								
fauna	Pupilla alpicola	2										
	Pupilla muscorum	2	14	23	7	7	7	2				1
	Slugs		3				1	2	6			2
	Succinella oblonga					3	8	3				7
	Trochulus hispidus	39	53	99	39	15	22	16	3		3	
	Vallonia costata	1	2	5								
	Vallonia sp.	1	4	2	1	1	2	1				
	Vertigo pygmaea		1	1								
	Ancylus fluviatilis	1		2	1							
	Anisus sp.	1	1	2			1			1		2
Freshwater	Pisidium amnicum							1	1	3	2	
fauna	Pisidium subtruncatum	1	15	24	11	4	22	18	6	14	2	
	<i>Radix</i> sp.	1	2	11						2		13
	Valvata macrostoma	1	8	17	2	4	3	8	23	63	25	26
	Total number of specimens	57	108	229	75	41	70	52	35	90	32	52
	Volume studied (in litres)	2	2	2	2	2	2	2	2	4	4	4