

The Romont tephrostratotype of the Rocourt Tephra, a widespread chronostratigraphic marker in western Europe

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ABSTRACT

In the loess stratotype of Belgium, in Romont quarry (Bassenge), minerals of a volcanic ash-fall deposit are dispersed in large quantities as a cryptotephra within a fossil humic horizon dating back to the end of the Weichselian Early Glacial. Chemical analysis of pyroxenes and amphiboles indicates that those minerals belong to the Rocourt Tephra as described in another four reference localities. The stratotype of this tephra is thus maintained in this quarry, as it is the only site of high chronostratigraphic value where it is still accessible.

RÉSUMÉ

Dans le stratotype des loëss de Belgique, dans la carrière du Romont (Bassenge), des minéraux d'une retombée volcanique sont dispersés en fortes quantités sous forme de cryptotéphra dans un horizon humifère fossile qui remonte à la fin du Début Glaciaire weichsélien. L'analyse chimique des pyroxènes et amphiboles démontre que ces minéraux appartiennent au Téphra de Rocourt tel qu'il a été décrit dans quatre autres localités de référence. Le stratotype de ce téphra est maintenu dans cette carrière car c'est le seul site à haute valeur chronostratigraphique où il est encore accessible.

KEYWORDS

Belgium,
Weichselian,
Humic Complex of
Remicourt,
cryptotephra,
pyroxenes,
amphibole

MOTS-CLÉS

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1. Introduction

The Rocourt Tephra (RT) is composed of explosively erupted ash-sized glass shards and of crystals, preserved in various host sediments in which they are too widely scattered and too fine (<350 μm) to be visible to the naked eye, i.e. it is a cryptotephra (Lowe, 2011). It was recognized by its concentration of volcanogenic mafic minerals (vmm) at about 40 sites in Belgium, western Germany, and south Netherlands (Fig. 1) (Juvigné et al., 2025). Volcanic glass shards have been detected at three sites in loess sections, Rocourt, Kesselt and Remicourt. They consist of blocky shards of hyaloclasts with a curvilinear surface and no vacuoles, which are typical of ash products from a major hydromagmatic eruptive event (Poulet et al., 2008). Based on its stratigraphic position, particularly in loess sequences, the age of the RT is bracketed between 70 and 80 ka (Juvigné et al., 2025).

In the Romont quarry, a cryptotephra has been discovered in a loess section and it was assigned to the RT based on its position in the stratigraphic sequence and its mineral association including clinopyroxenes, orthopyroxenes and amphiboles (Juvigné et al., 2008). The association of the vmm was established at all above cited sites where enough minerals were collected, and variations in relative frequencies were pointed out, particularly in Romont where the frequencies of amphiboles (5.4%) and orthopyroxenes (3.8%) are exceptionally low versus clinopyroxenes (90.8%).

The Romont quarry is located between Visé, Riemst and Maastricht on the left bank of the Geer valley (Fig. 2). Heidelberg Materials extracts Maastrichtian (Upper Cretaceous) chalk there. The overlying terrains consist of remnants of Lower and Middle Pleistocene Meuse terraces (Juvigné, 2024), themselves covered with loess. Outcrops of these terrains remain accessible around the perimeter of the quarry, and representative stratigraphic sequences are maintained by the operator for further scientific research. One of them has been defined as the stratotype for the loess sequence of Belgium (Haesaerts et al., 2011). Minerals likely to belong to the RT have been identified under the microscope in that section (Juvigné et al., 2008), which was therefore proposed to replace the previous tephrostratotype (Rocourt, backfilled in 2018).

The uncertainties of this optical technique, however, require geochemical identification to confirm the site's recognition as the tephrostratotype of the Rocourt Tephra.

Minerals and glass shards in the RT have been analysed by electron probe microanalyser at several sites (Rocourt, Kesselt, Remicourt and Walou), and a unique fingerprint was developed for mineralogical compositions (Poulet & Juvigné, 1993; Poulet et al., 2008). At the Romont site, the relative frequency of the vmm association is different from that of the other sites (see above). The aim of this study is to analyse the minerals of the cryptotephra of Romont to confirm or refute its inclusion in the RT.

2. Analytical procedures

The bulk sample was treated as follow: dispersion in boiling water, underwater sieving to isolate the useful fraction of 355/75 μm so that the loess (very rich in dense minerals) and the coarse fraction (without vmm) are eliminated, boiling in hydrogen peroxide to remove micro-particles of humus, and drying. Dense minerals (>2.8 g/cm^3) were separated with bromoform by repeating agitation-decantation-harvest cycles, until no more harvest was obtained (3 to 5 cycles). Aliquots were examined for identification with a petrographical microscope. In the fraction thus isolated, the vmm constitute the essential part of the minerals. Feldspars (light fraction) have not been studied because it is not possible to make sure that the feldspars belong to the RT rather than to the Oligocene marine sediments, a mound of which is present close to the stratotype. The search for volcanic glasses using density gradient and magnetic separation (Frantz) was unsuccessful despite processing a significantly larger sample of bulk sediment.

Minerals have been analysed with a Jeol JXA-iHP200F electron microprobe (15 kV, 10 nA) at Camparis, University Paris-Sorbonne, France. For this purpose, $K\alpha$ lines were used. The chemical composition was determined using the following standards: garnet for Si, Ca, Fe and Mg, MnTiO_3 for Ti and Mn, Cr_2O_3 for Cr. The counting times were 10 s for peaks and 10 s

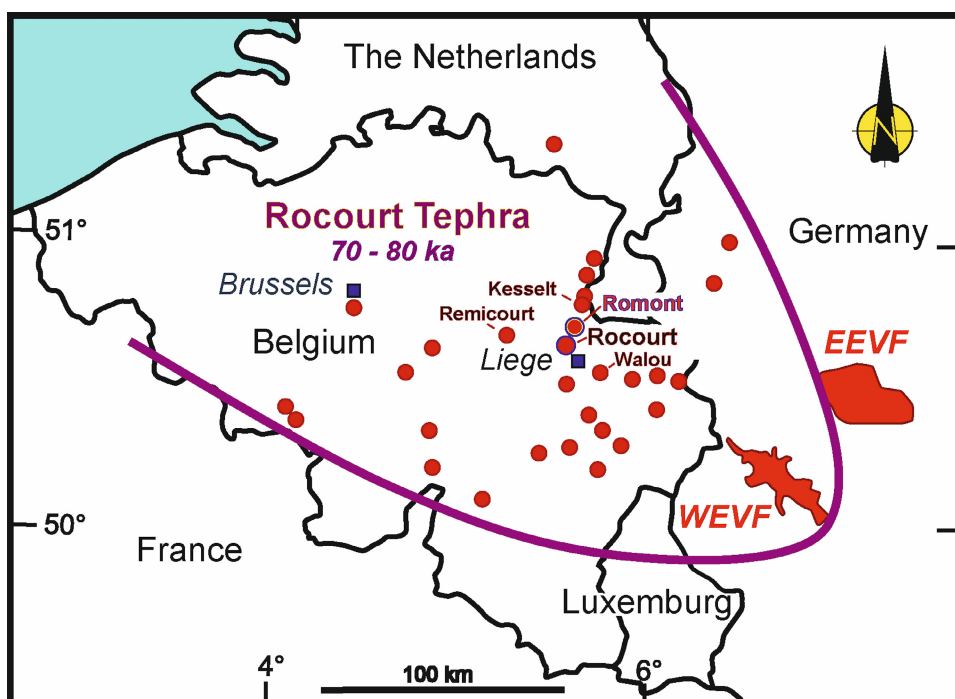


Figure 1. Geographical distribution of the Rocourt Tephra (RT), a widespread Quaternary stratigraphic marker in western Germany, Belgium, and south Netherlands (Juvigné et al., 2025). Red dots locate the tephra sites, the analysed sites being Rocourt, Remicourt, Romont (this work), Kesselt and Walou. West Eifel Volcanic Field (WEVF) where the RT emitting volcano should be located (Juvigné et al., 2024). Abbreviation: EEVF, East Eifel Volcanic Field.

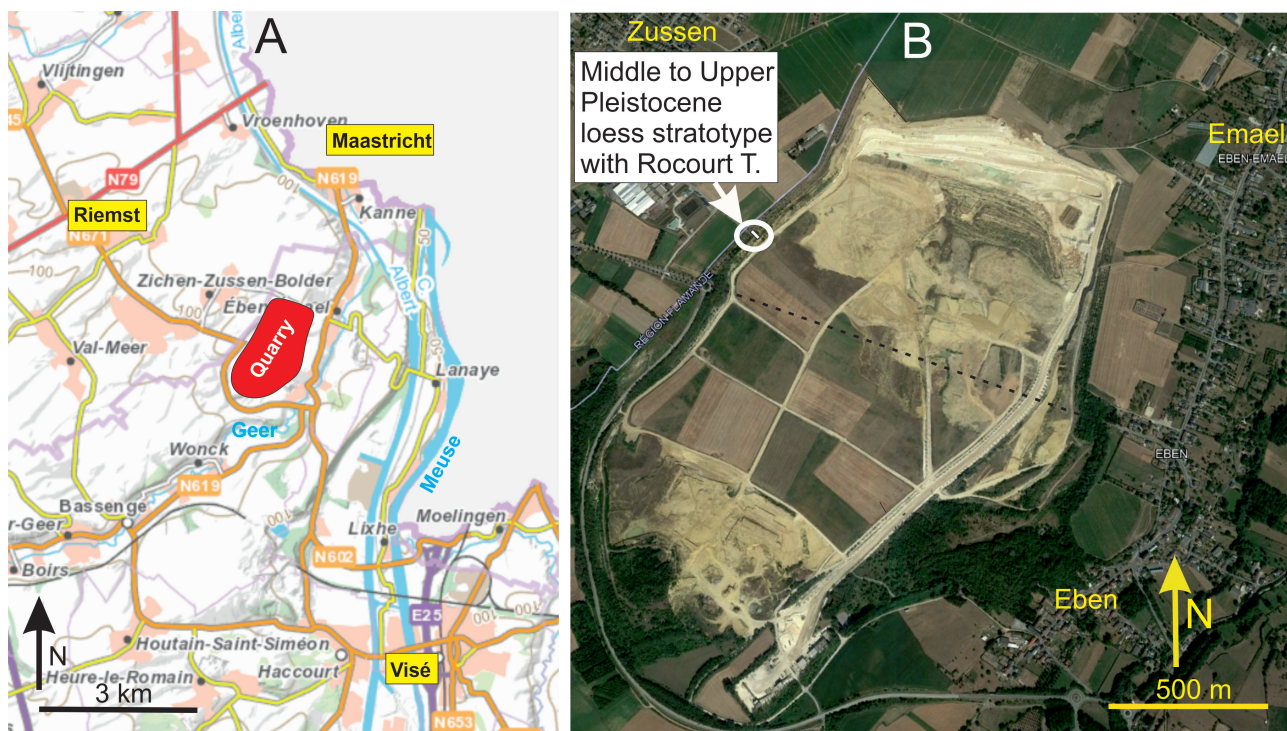


Figure 2. Romont quarry (Heidelberg Materials Romont & Lœn) where the loess stratotype containing the Rocourt Tephra is preserved for further scientific research. **A.** Location of the quarry. **B.** Location of the loess stratotype including the Rocourt Tephra at the edge of the quarry ($x = 239895$; $y = 165221$ (Belgian Lambert 72; EPSG 31370)) (progress status of the quarry in 2024). Abbreviation: Rocourt T., Rocourt Tephra.

for background, with a 5- μm defocused beam. As standards, combinations of natural and synthetic minerals were used. Data corrections were made using the PAP method correction of Pouchou & Pichoir (1991).

3. The Romont loess reference sequence and the RT sampling

The loess sequence in Romont has been first studied in the early 1990s (Juvigné, 1992). In 2007, an extension of the quarry gave access to a more complex sequence, encompassing most of the known litho- and pedostratigraphic markers of the Late Pleistocene loess deposits of Belgium and north-western Europe (Fig. 3), including the Rocourt Pedocomplex and the Humic Complex of Remicourt (HCR), covering the equivalent of the whole MIS 5 (Eemian interglacial and Weichselian Early Glacial; Haesaerts et al., 2016). In the HCR, the RT has been identified, and the location has been proposed as its new tephrostratotype (Juvigné et al., 2008). This thick succession of loess/palaeosols has then been described as the new stratotype for the loess cover from Middle Belgium, being defined as the Romont Group (Haesaerts et al., 2011).

At the sampling location, most of the classic stratigraphic markers of the Late Pleistocene loess reference sequence are present (Haesaerts et al., 2016), i.e. from bottom to top (soil classification according to IUSS Working Group WRB, 2006): (1) the Rocourt Pedocomplex, including three pedogeneses recorded by Bt to Bth horizons (luvisols to “greyzem”; Antoine et al., 2016; Haesaerts et al., 2016; Schirmer, 2016; see also Vancampenhout et al., 2013a, 2013b); (2) the Humic Complex of Remicourt (HCR; see below); (3) a thick deposit of laminated colluvium; (4) the Malplaquet soil present as a cambic Bw horizon interpreted as belonging to a cambisol (“boreal brown soil” of Antoine et al., 2016); (5) the Kincamp soil, a humic A

horizon probably developed under dense herb cover (Haesaerts et al., 2016); (6) the Nagelbeek tongued horizon (NTH) (Haesaerts et al., 1981); (7) the Brabantian loess (Fig. 3).

The Hesbayan loess, situated below the NTH in other parts of the quarry, is strongly eroded in the sampled section. This situation explains the absence of the Eltville Tephra (Pouquet & Juvigné, 2009) in the studied sequence.

The HCR is c. 50 cm thick and is composed of three units, which are, from bottom to top: a dark grey strongly bioturbated horizon (HCR-a); a humic loam (HCR-b); and finally, a dark grey horizon with oxidoreduction features (HCR-c) (Haesaerts et al., 1999, 2016). The bulk sample containing the cryptotephra was taken in April 2025 by one of us (EJ), in unit HCR-b within the stratotype. The sampled place was chosen after it was verified that it contains the highest concentration of vmm of the RT as shown in detail on a very close identical sequence by Juvigné et al. (2008). The decrease of frequency above and below the vmm peak is explained: (1) in the overlying laminated colluvium, by sedimentary reworking, evidenced by the fine laminations of the material eroding and reworking the HCR; (2) in the underlying horizon (HCR-a) by bioturbations, mainly by the action of burrowing animals. Thus, we can assume that the tephra fall-out took place after the formation of unit HCR-a but before the formation of unit HCR-c, a conclusion also reached in Remicourt, another site with detailed sampling and analyses of the RT (Juvigné et al., 2013).

4. Composition and significance of mafic minerals of the tephra

The electron probe microanalysis data comprise 42 analyses of clinopyroxenes, 5 of orthopyroxenes and 5 of amphiboles.

Clinopyroxenes show a broad compositional range from magnesian augite to fassaitic diopside (Fig. 4A). They are

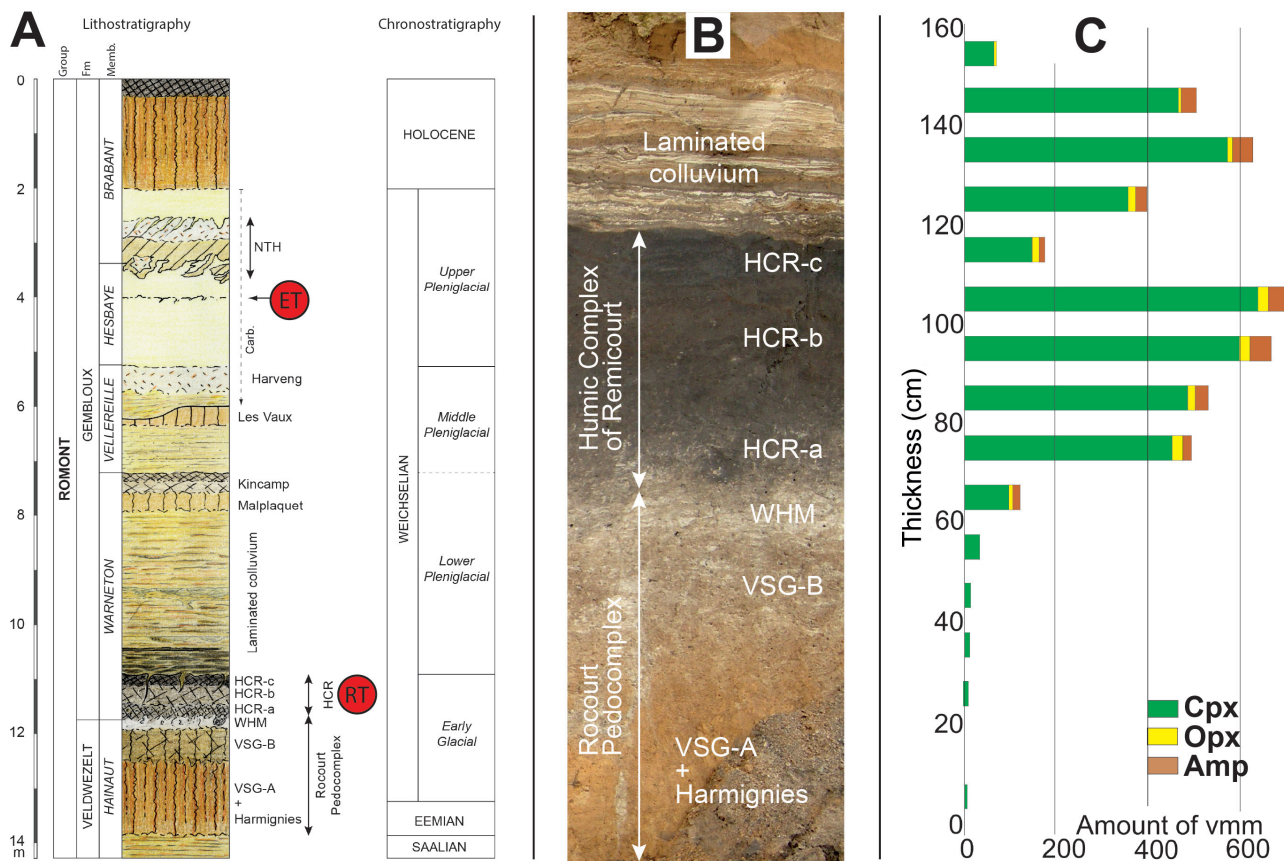


Figure 3. A. The synthetic loess sequence for the Late Pleistocene in the stratotype of the Romont quarry (Haesaerts et al., 2016). **B.** Section showing the stratigraphic context of the Rocourt cryptotephra (same vertical scale as C). **C.** Number of vmm per gram of the particle size fraction between 105 and 420 μm in the section studied by Juvigné et al. (2008) in the same quarry, where 15 samples were analysed above, within and below the layer of highest concentration. Abbreviations: Amp, amphiboles; Carb., carbonated sediment; Cpx, clinopyroxenes; ET, Eltville Tephra; Fm, Formation; HCR, Humic Complex of Remicourt; Memb., Member; NTH, Nagelbeek tongued horizon; Opx, orthopyroxenes; RT, Rocourt Tephra; WHM, Whitish horizon of Momalle; vmm, volcanogenic mafic minerals; VSG-A and VSG-B, Villers-Saint-Ghislain A and B soils.

distributed in three groups (Fig. 4B–D) which are distinguishable by their contents in Al^{IV} , Al^{VI} , Ti, and Na in the formulae units (Table 1). The first group consists of few Mg-rich and Cr-rich diopside enriched in Al^{VI} at the expense of Al^{IV} with low Ti content. According to Juvigné et al. (2024), these clinopyroxenes are high pressure xenocrysts extracted from peridotitic and pyroxenitic mantle xenoliths. In fact, they have the same composition as the mantle clinopyroxenes of xenocrysts and nodules of lherzolite and pyroxene from the lavas of the Eifel volcanoes (Duda & Schmincke, 1985; Witt-Eickschen et al., 1998). The second group comprises megacrysts of clinopyroxenes discriminated by their relatively high Al^{VI} , Ti and Na contents. They originated from early crystallization of primitive magma stored in deep crustal reservoirs. The third group includes phenocrysts distinguished by their high Ti and Ca contents with a fassaitic trend. They have crystallized at low pressure in the upper crust magma chamber of the emitting volcano. The same megacrysts and phenocrysts were analysed in the Eifel lavas and interpreted in the same way by Becker (1977), Duda & Schmincke (1978, 1985), Stosch & Lugmair (1986), Shaw & Eyzaguirre (2000) and Shaw (2024).

Orthopyroxenes are magnesian enstatite (Table 2). Like clinopyroxene xenocrysts, these orthopyroxenes have the composition of mantle pyroxenes from xenocrysts and nodules of lherzolite and pyroxenite analysed in lavas from the Eifel volcanoes (Duda & Schmincke, 1985; Witt-Eickschen et al., 1998).

Amphiboles are determined as Ti-rich pargasite (Locock,

2014) (Table 3). Most of them are xenocrysts that originated from upper mantle to lower crust cumulates and veins of amphibole-bearing pyroxenites according to the data of Witt-Eickschen et al. (1998) and Shaw & Eyzaguirre (2000) on xenolithic amphiboles of the Eifel volcanoes (Fig. 5, areas Acx and Hvp). One may be a lava megacryst or phenocryst according to the data of Duda & Schmincke (1978, 1985) and Wörner & Schmincke (1984).

5. Discussion and conclusion

The Rocourt Tephra, a cryptotephra including clinopyroxenes, orthopyroxenes and amphiboles, has been described in the Humic Complex of Remicourt in several sites on the Belgian loess belt since 1954 (Gullentops, 1954; see also Haesaerts et al., 2016, and Juvigné et al., 2025). Electron probe analyses performed on mafic minerals from three of these loess sites (Rocourt, Kesselt and Remicourt) confirmed its unique geochemical fingerprint (Pouclet et al., 2008).

The same mineralogical association found in the Humic Complex of Remicourt in the Romont quarry and studied under the polarising microscope (Juvigné et al., 2008) was analysed through microprobe. The results presented in this paper show that the association of the different pyroxenes is strictly similar to those obtained on minerals from these three loess sites (Rocourt, Kesselt and Remicourt) as well as on the cryptotephra from Walou Cave (Pouclet & Juvigné, 1993; Pouclet et al., 2008), which has a very comparable stratigraphic position (Pirson, 2011; Pirson

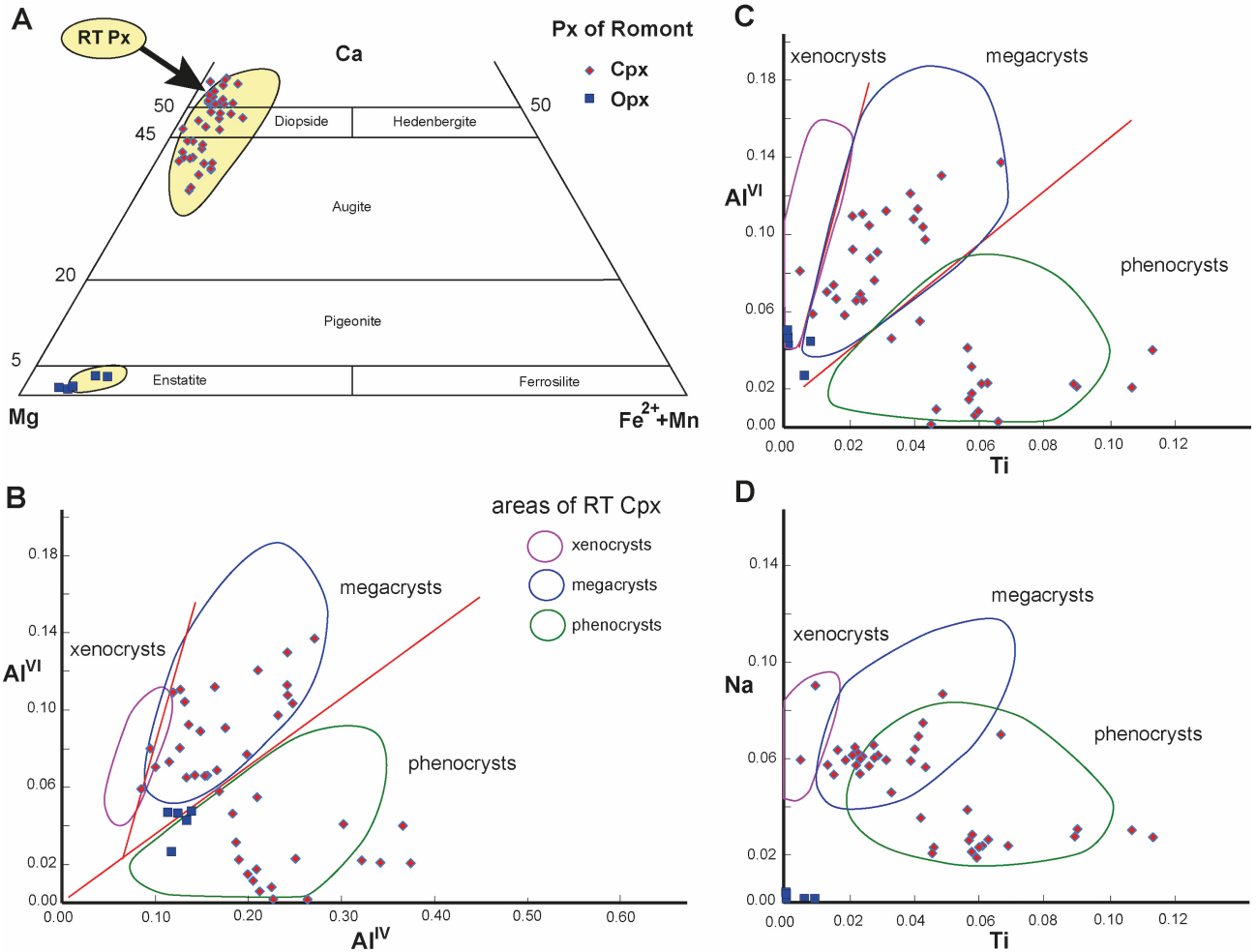


Figure 4. Comparison of the Romont pyroxenes (purple dots) with the compositional areas of the RT pyroxenes redrawn after Juvigné et al. (2024). **A.** Mg–Ca–Fe²⁺ + Mn diagram and the nomenclature of Morimoto (1988). **B., C., and D.** Covariation diagrams of Al^{VI}, Al^{IV}, Ti and Na of the formulae units. The pyroxenes are distributed in three groups: xenocrysts, megacrysts, and phenocrysts. Abbreviations: Cpx, clinopyroxenes; Px, pyroxenes; Opx, orthopyroxenes; RT, Rocourt Tephra.

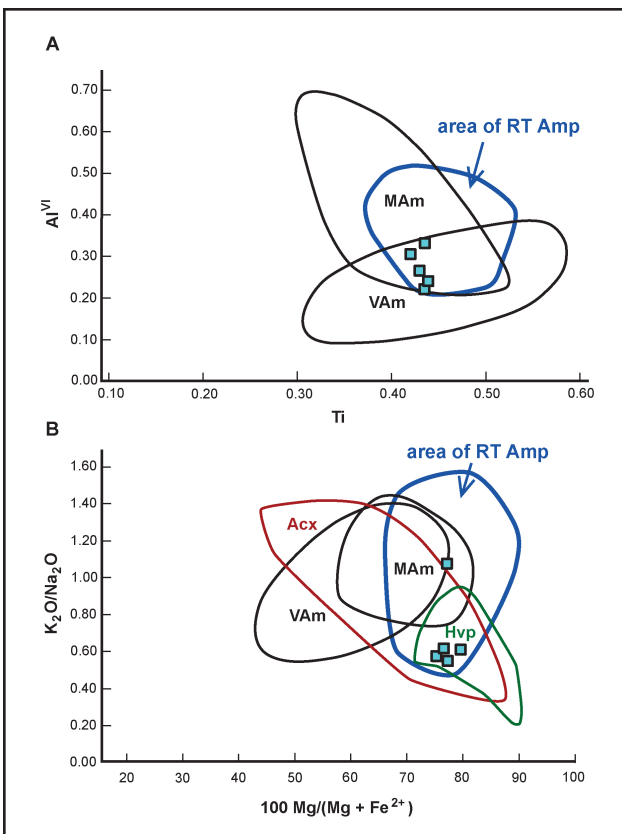


Figure 5. Ti versus Al^{VI} (**A**) and 100Mg/(Mg+Fe²⁺) versus K₂O/Na₂O (**B**) diagrams of the amphiboles. Area of the RT amphiboles after Pouclet et al. (2008). MAm, megacrystic amphiboles and VAm, amphibole phenocrysts of the Eifel lavas; Acx and Hvp, amphiboles from pyroxenite xenoliths and from veins in peridotite, respectively. Areas redrawn after the data of Duda & Schmincke (1978), Wörner & Schmincke (1984), Witt-Eickschen et al. (1998) and Shaw & Eyzaguirre (2000). Abbreviations: Amp, amphiboles; RT, Rocourt Tephra.

Table 1. Selected chemical analyses of clinopyroxenes.

Major oxides (weight percent)	Xenocrysts		Megacrysts					Phenocrysts				
SiO ₂	51.06	52.83	50.23	49.47	48.36	48.00	50.17	45.97	43.41	44.46	48.82	48.43
TiO ₂	0.17	0.34	0.68	1.01	1.57	1.45	0.98	2.02	3.78	3.20	1.51	2.07
Al ₂ O ₃	4.82	3.42	5.29	6.45	7.65	8.12	5.40	7.87	8.95	8.24	6.10	5.21
FeOt	3.04	3.01	5.03	5.36	4.29	5.26	5.71	5.85	6.29	6.57	4.62	5.93
Cr ₂ O ₃	1.32	1.61	0.73	0.46	0.83	0.59	0.40	0.37	0.31	0.17	0.57	0.16
MnO	0.17	0.11	0.10	0.14	0.08	0.08	0.18	0.09	0.09	0.12	0.13	0.04
MgO	17.59	17.06	17.88	16.42	14.63	15.01	17.24	13.50	12.62	12.66	14.69	14.86
CaO	19.23	20.11	18.26	19.58	21.62	20.10	17.14	23.35	23.18	23.50	22.87	22.41
Na ₂ O	0.84	1.30	0.84	0.86	0.80	0.90	0.90	0.54	0.42	0.43	0.50	0.40
K ₂ O	0.01	0.02	0.04	0.04	0.04	0.00	0.01	0.06	0.00	0.03	0.06	0.02
Total	98.26	99.80	99.09	99.79	99.87	99.51	98.13	99.62	99.05	99.37	99.89	99.52
Fe ₂ O ₃ *	2.15	2.09	4.13	4.30	3.00	3.67	2.13	6.39	5.71	5.94	3.33	3.64
FeO	1.11	1.13	1.31	1.49	1.59	1.95	3.79	0.09	1.15	1.22	1.63	2.65
* Papike et al. (1974)												
Formulae on the basis of 6 oxygens												
Si	1.872	1.912	1.831	1.800	1.768	1.757	1.852	1.698	1.626	1.659	1.791	1.791
Al ^{IV}	0.128	0.088	0.169	0.200	0.232	0.243	0.148	0.302	0.374	0.341	0.209	0.209
T	2.000	2.000	2.000	2.000	2.000	2.000	2.000	2.000	2.000	2.000	2.000	2.000
Al ^{VI}	0.081	0.058	0.058	0.076	0.097	0.108	0.087	0.041	0.020	0.021	0.055	0.017
Ti ^{VI}	0.005	0.009	0.019	0.028	0.043	0.040	0.027	0.056	0.106	0.090	0.042	0.057
Fe ³⁺	0.059	0.057	0.113	0.118	0.083	0.101	0.059	0.178	0.161	0.167	0.092	0.101
Cr	0.038	0.046	0.021	0.013	0.024	0.017	0.012	0.011	0.009	0.005	0.017	0.005
Fe ²⁺	0.028	0.030	0.031	0.037	0.043	0.050	0.089	0.003	0.034	0.037	0.047	0.074
Mg	0.789	0.800	0.758	0.728	0.710	0.684	0.725	0.712	0.669	0.681	0.748	0.745
M1	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.000
Fe ²⁺	0.006	0.004	0.009	0.008	0.097	0.010	0.087	0.000	0.002	0.001	0.003	0.007
Mg	0.173	0.121	0.214	0.162	0.043	0.135	0.027	0.032	0.036	0.024	0.055	0.075
Mn	0.005	0.003	0.003	0.004	0.083	0.002	0.059	0.003	0.003	0.004	0.004	0.001
Ca	0.756	0.780	0.713	0.763	0.024	0.789	0.012	0.924	0.930	0.940	0.899	0.888
Na	0.060	0.091	0.059	0.061	0.043	0.064	0.089	0.039	0.031	0.031	0.035	0.028
K	0.000	0.001	0.002	0.002	0.710	0.000	0.725	0.003	0.000	0.002	0.003	0.001
M2	1.000	1.000	1.000	1.000	1.000	1.000	1.000	1.001	1.001	1.001	1.000	1.000
Total	4.000	4.000	4.000	4.000	4.000	4.000	4.000	4.001	4.001	4.001	4.000	4.000
Mg/Mg+Fe ²⁺	0.966	0.964	0.960	0.951	0.942	0.932	0.890	0.996	0.951	0.949	0.941	0.909

& Juvigné, 2011). Amphiboles from Romont are also similar to those analysed in these four sites. While we have not found any spinel nor glass, based on its stratigraphic location and the chemical signature of the pyroxenes, there is no doubt that the Romont cryptotephra is a fallout deposit of the Rocourt Tephra. The loess stratotype preserved in the Romont site can therefore be confirmed as the new tephrostratotype of the Rocourt Tephra. Its stratigraphic position inside the Humic Complex of Rémicourt fits with the other occurrences of the tephra in the loess sequences that form the type region.

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Table 2. Results of chemical analyses of orthopyroxenes. Abbreviation: Opx, orthopyroxenes.

Major oxides (weight percent)	Opx1	Opx2	Opx3	Opx4	Opx5
SiO ₂	54.38	54.86	54.29	54.90	54.60
TiO ₂	0.00	0.00	0.00	0.24	0.31
Al ₂ O ₃	4.67	4.26	4.41	3.55	4.04
FeOt	6.07	6.14	7.01	7.85	8.59
Cr ₂ O ₃	0.37	0.37	0.44	0.31	0.19
MnO	0.11	0.21	0.13	0.14	0.21
MgO	34.17	34.20	33.66	32.26	31.41
CaO	0.56	0.52	0.55	1.98	1.86
Na ₂ O	0.03	0.03	0.01	0.01	0.01
K ₂ O	0.01	0.03	0.01	0.00	0.04
Total	100.37	100.61	100.51	101.24	101.25
Fe ₂ O ₃ *	3.28	2.89	3.34	2.44	1.96
FeO	3.12	3.54	4.00	5.66	6.82
* Papike et al. (1974)					
Formulae on the basis of 6 oxygens					
Si	1.860	1.874	1.862	1.886	1.883
Al ^{IV}	0.140	0.126	0.138	0.114	0.117
T	2.000	2.000	2.000	2.000	2.000
Al ^{VI}	0.048	0.045	0.041	0.030	0.047
Ti ^{VI}	0.000	0.000	0.000	0.006	0.008
Fe ³⁺	0.084	0.074	0.086	0.063	0.051
Cr	0.010	0.010	0.012	0.008	0.005
Fe ²⁺	0.042	0.048	0.054	0.080	0.097
Mg	0.816	0.823	0.807	0.812	0.792
M1	1.000	1.000	1.000	1.000	1.000
Fe ²⁺	0.047	0.053	0.061	0.083	0.100
Mg	0.926	0.919	0.914	0.840	0.823
Mn	0.003	0.006	0.004	0.004	0.006
Ca	0.020	0.019	0.020	0.073	0.069
Na	0.002	0.002	0.001	0.001	0.001
K	0.001	0.001	0.001	0.000	0.002
M2	1.000	1.000	1.000	1.000	1.000
Total	4.000	4.000	4.000	4.000	4.000
Mg/Mg+Fe ²⁺	0.951	0.945	0.938	0.910	0.891

Table 3. Results of chemical analyses of amphiboles. Abbreviations: Amp, amphiboles; Prg, pargasite.

Major oxides (weight percent)	Amp 1	Amp 2	Amp 3	Amp 4	Amp 5
SiO ₂	39.98	39.27	38.86	39.77	41.20
TiO ₂	3.89	3.88	3.73	3.85	4.02
Al ₂ O ₃	13.64	13.48	14.25	13.74	13.23
FeO (a)	8.34	8.05	7.67	7.84	7.18
Fe ₂ O ₃ (a)	0.76	1.01	1.23	1.03	1.36
Cr ₂ O ₃	0.04	0.00	0.00	0.08	0.13
MnO	0.16	0.09	0.12	0.15	0.11
MgO	14.15	14.69	14.29	14.80	15.54
CaO	11.56	12.12	12.02	11.91	11.75
Na ₂ O	2.49	2.46	2.12	2.51	2.35
K ₂ O	1.42	1.52	2.28	1.37	1.42
H ₂ O+ (a)	2.02	2.01	2.01	2.03	2.07
Total	98.44	98.58	98.57	99.09	100.37
Formulae on the basis of 22 oxygens (a)					
Si	5.943	5.850	5.800	5.874	5.976
Al ^{IV}	2.057	2.150	2.200	2.126	2.024
T	8.000	8.000	8.000	8.000	8.000
Ti	0.435	0.434	0.419	0.428	0.438
Al ^{VI}	0.332	0.217	0.306	0.266	0.238
Cr	0.005	0.000	0.000	0.010	0.014
Fe ³⁺	0.085	0.113	0.138	0.115	0.149
Fe ²⁺	1.008	0.973	0.957	0.921	0.800
Mg	3.136	3.262	3.179	3.259	3.360
C	5.001	4.999	4.999	4.999	4.999
Mn ²⁺	0.020	0.012	0.014	0.018	0.014
Fe ²⁺	0.029	0.030	0.000	0.047	0.071
Ca	1.841	1.934	1.922	1.885	1.826
Na	0.111	0.024	0.064	0.050	0.089
B	2.001	2.000	2.000	2.000	2.000
Na	0.608	0.685	0.548	0.670	0.572
K	0.269	0.289	0.435	0.259	0.263
A	0.877	0.974	0.983	0.929	0.835
Sum T,C,B,A	15.879	15.973	15.982	15.928	15.834
Mg/Mg+Fe ²⁺	0.75	0.76	0.77	0.77	0.79
Species	Ti-rich Prg	Ti-rich Prg	Ti-rich Prg	Ti-rich Prg	Ti-rich Prg
(a) calculation using the spreadsheet of Locock (2014)					

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Author contributions

EJ, AP and SP designed the study. EJ performed the sample collection. JMB analysed the samples. EJ and AP wrote the introduction and prepared Figure 1. SP and EJ wrote the stratigraphy and data sampling and prepared Figures 2 and 3. AP wrote the analytical chapter with part of the discussion and prepared Figures 4 and 5 and all the tables. All authors reviewed the manuscript.

Data availability

All chemical analyses of the clinopyroxenes are available in the Supplementary material.

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