

Original paper

Revision of Scheumann's classification of melilitic lamprophyres and related melilitic rocks in light of new analytical data

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Dykes of the Late Cretaceous to Early Tertiary (79.5±3.5 to 60.7±2.4 Ma) melilitic rock series of the Osečná Complex and the Devil's Walls dyke swarm, including ultramafic lamprophyres – polzenites – of Scheumann (1913) occur dispersed in the entire Upper Ploučnice River basin in northern Bohemia.

Polzenites and associated melilitic rocks are characterized by the mineral association of olivine + melilite ± nepheline, haüyne, monticellite, phlogopite, calcite, perovskite, spinels and apatite. New data on their mineral and chemical compositions from original Scheumann's localities (the Vesec, Modlibohov, Luhov types) argue against the abolition of the group of ultramafic lamprophyres and the terms 'polzenite' and 'alnöite' by the Le Maitre (2002) classification. Marginal facies and numerous flat apophyses of the lopolith-like body known as the Osečná Complex show an olivine micro-melilitolite composition (lamprophyric facies). The porphyritic texture, chemical composition and the presence of characteristic minerals such as monticellite and phlogopite point to their affinity with ultramafic lamprophyres – polzenites of the Vesec type. Melilite-bearing olivine nephelinites to olivine melilitites (olivine + clinopyroxene + nepheline + melilite ± haüyne and spinels with apatite) form a swarm of subparallel dykes known as the Devil's Walls.

The Scheumann's non-melilite dyke rock "wesselite", spatially associated with polzenites and often erroneously attributed to the polzenite group, is an alkaline lamprophyre of monchiquite to camptonite composition (kaersutite + phlogopite + diopside + olivine phenocrysts in groundmass containing clinopyroxene, phlogopite, haüyne, analcime, titanian magnetite, apatite ± glass/plagioclase). First K–Ar data show Oligocene ages (30.9±1.2 to 27.8±1.1 Ma) and an affinity to the common tephrite–basanite rock series.

Keywords: melilitic rocks, lamprophyres, polzenites, classification, geochemistry, Bohemian Massif

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1. Introduction

Rare ultramafic melilite-bearing olivine nephelinite to olivine melilitite volcanic rocks and their plutonic equivalents (mostly of the melteigite–ijolite–urtite series) and associated lamprophyric alnöite represent low-volume mantle-derived melts (Ishikawa et al. 2004; Melluso et al. 2011). Rocks of melilitic volcanic series occur in intraplate rift settings (Siberia – Egorov 1970; Zhabin and Surina 1970; the Balcones Province, Texas, and Tasmania – Brey 1978; Africa – Dawson et al. 1985; Bailey et al. 2005; Keller et al. 2006; Greenland – Bernstein et al. 2000; Madagascar – Melluso et al. 2011; Euro–Mediterranean region – Lustrino and Wilson 2007; Piromallo et al. 2008 and citations therein), at continental margins (the western Canary and Madeira islands are situated on oceanic lithosphere, whereas the eastern Canary Islands on transitional, oceanic/continental lithosphere – Hoernle and Schmincke 1993) and in ocean island settings (Hawaii – Wilkinson and Stolz 1983; Maaloe et al. 1992; French Polynesia – Chauvevel et al. 1992). Potassic to

ultrapotassic melilitic rocks of exceptional subduction-related orogenic settings are represented by the Roman Magmatic Province (Di Battistini et al. 2001 and citations therein) and a diatreme filling of polzenite (modlibovite) composition in California (Morgan et al. 1985).

Alnöite dykes are rare in rift-related settings, e.g., Alnö Island, Sweden (Eckermann 1961), Monteregian Hills, Canada (Hamois and Mineau 1991), Maimecha–Kotui Province, Siberian Platform (Egorov 1970; Zhabin and Surina 1970), Kola Peninsula (Beard et al. 1998; Rass 2008), Gardar Province, Greenland (Pearce et al. 1997), Sierra Subandinas, Argentina (Barbieri et al. 1997) but also in an island-arc setting on the Solomon Islands, Ontong Java Plateau (Ishikawa et al. 2004).

Melilitic rocks occur throughout the Central European Volcanic Province (CEVP) (Alibert et al. 1983; Keller 1984; Keller et al. 1990; Hegner et al. 1995; Wilson et al. 1995; Hegner and Vennemann 1997; Dunworth and Wilson 1998; Lustrino and Wilson 2007) including the Bohemian Massif. They are particularly common in western Bohemia and Saxony (Ulrych et al. 2000a, b;

2010; Seifert et al. 2008; Abratis et al. 2009; Krüger et al. 2013). Besides these usual melilitic volcanic rocks with substantial clinopyroxene, Scheumann (1913, 1922) recognized clinopyroxene-free lamprophyric dyke rocks in the Ploučnice River Region (PRR) in northern Bohemia and described them as polzenites. Pivec et al. (1986, 1998) and Ulrych et al. (1988, 2008) reported the presence of the hypabyssal Osečná olivine melilitolite intrusion, also free of clinopyroxene, from the same area.

Similar Late Cretaceous flat-lying sheets of ultramafic lamprophyres are also known from other localities of the Bohemian Massif. The sheets of monchiquite–ouachitite–aillikite series penetrate the Permo–Carboniferous sediments of central Bohemia (Ulrych et al. 1993), and a phlogopite-rich olivine clinopyroxenite (autotransformed alnöite) sheet intruded the boundary between Upper Permian and Upper Cretaceous sediments near Dvůr Králové nad Labem (Ulrych et al. 1996).

The present paper summarizes information on melilitic rocks from northern Bohemia (Ulrych et al. 2008) and presents new chemical analyses and K–Ar dating of melilitic and melilite-bearing dykes at the classical Scheumann’s localities.

2. Geological position

Late Cretaceous to Cenozoic intraplate volcanism in Central Europe belongs to the Central European Volcanic Province (CEVP) and concentrates to the European Cenozoic Rift System (Prodehl et al. 1995). The easternmost segment of this rift system is represented by the Ohře/Eger Rift (OR) in the Bohemian Massif. Peak volcanic activity related to this structure coincided with the maximum crustal extension and graben formation in the Late Eocene to Early Miocene; however, Cenozoic volcanic products in northern Bohemian Massif are not restricted to this event. Three phases of elevated volcanic activity can be distinguished (Ulrych et al. 2011): (i) pre-rift Upper Cretaceous to Palaeocene ultramafic melilitic rock series, (ii) syn-rift Oligocene to Early Miocene tephrite/basanite–phonolite/trachyte rock series, and (iii) sporadically occurring ultramafic rocks of the late-rift Late Miocene period.

The PRR is well known by the occurrence of Late Cretaceous to Palaeocene ultramafic intraplate volcanic rocks characterized by the presence of melilite. Scheumann (1913) defined a suite of clinopyroxene-free ultramafic lamprophyres from this region and named it after the Ploučnice River (*Polzen* in German) – polzenites.

The occurrences of polzenites and associated melilitic rocks in northern Bohemia concentrate along the SE marginal fault of the Ohře/Eger Rift graben. Strikes of their dykes (mostly NE–SW) are compatible with the stress field which governed the main episode of thrusting at the

NW–SE-striking Lusatian Fault (Adamovič and Coubal 1999; Ulrych et al. 2011). At this time, fractures parallel to the maximum principal stress component became dilated, permitting a rapid magma ascent through the crust.

Polzenites and associated melilitic rocks are represented by (1) dykes striking NE–SW, rarely ENE–WSW, several kilometres in length, and (2) the central intrusion of olivine melilitolite at Osečná (Pivec et al. 1986, 1998; Ulrych et al. 1988, 2008; Fig. 1). The Osečná intrusion is an asymmetrical, lopolith-shaped body, 20 to 60 m thick, emplaced in sediments of the Cenomanian to Turonian age (depth 0–200 m) over an area of 12.5 km² (see Fig. 2 in Ulrych et al. 2008). The dip angles of the Osečná saucer-shaped sill and flat-lying sheets are low (up to 30°) but steepen to as much as 70–80° on the edges of the saucer, forming finger-like apophyses (see also similar steepening from the Whin Sill, England and Midland Valley Sill, Scotland – Francis 1982). The intrusion reaches the surface only rarely in the form of marginal facies formed by fine-grained micro-melilitolite (micro-porphyrific melilitolite).

The Osečná Complex is spatially associated with dykes of olivine nephelinite to olivine melilitite compositions, dipping steeply to the NW (Wurm 1884). These comprise (from NW to SE) the Western Devil’s Wall, the Great Devil’s Wall and the Lesser Devil’s Wall (Fig. 1). Dykes of this composition are interspersed with a number of shorter dykes of polzenite which are of the same orientation and were obviously emplaced under the same tectonic stress field: the Kotel dyke (clinopyroxene “polzenite” – alnöite) is an example. The dyke of Pelousek Hill passes the village of Modlibohov and was most probably the site of the “Modlibohov-type” polzenites of Scheumann (1913). Svárovské návrší Hill at Osečná is a segment of a cone-sheet(?) dyke dipping at moderate angles. A NW-dipping micro-melilitolite dyke 700 m long between Vesec and Jiríčkov (Fig. 1) was mentioned by Scheumann (1913) but more attention was given to “Vesec-type” micro-melilitolite dykes at Děvín and Svojkov.

Farther west, between Mimoň and Osečná, the best examples of micro-melilitolite are the Great Ralsko dyke and the morphologically prominent Děvín dyke, also dipping steeply to the NW. The NE–SW-striking alnöite dyke is located within the subsided block of the OR near Luhov (Scheumann 1913). It crosses the top of Černý vrch Hill but provides no natural outcrops. Some of the westernmost dykes, also located within the OR, include the N-dipping micro-melilitolite dyke near Svojkov known as “Bockwener Mühle” (Scheumann 1913) and the NW-dipping alnöite dyke at Brenná known as “der Damm” (Wurm 1883; Scheumann 1913).

Occurrences of olivine melilitite and polzenite dykes continue to Saxony (Zeughausgang near Hinterhermsdorf – Pfeiffer 1978 and citations therein) and to Lusatia

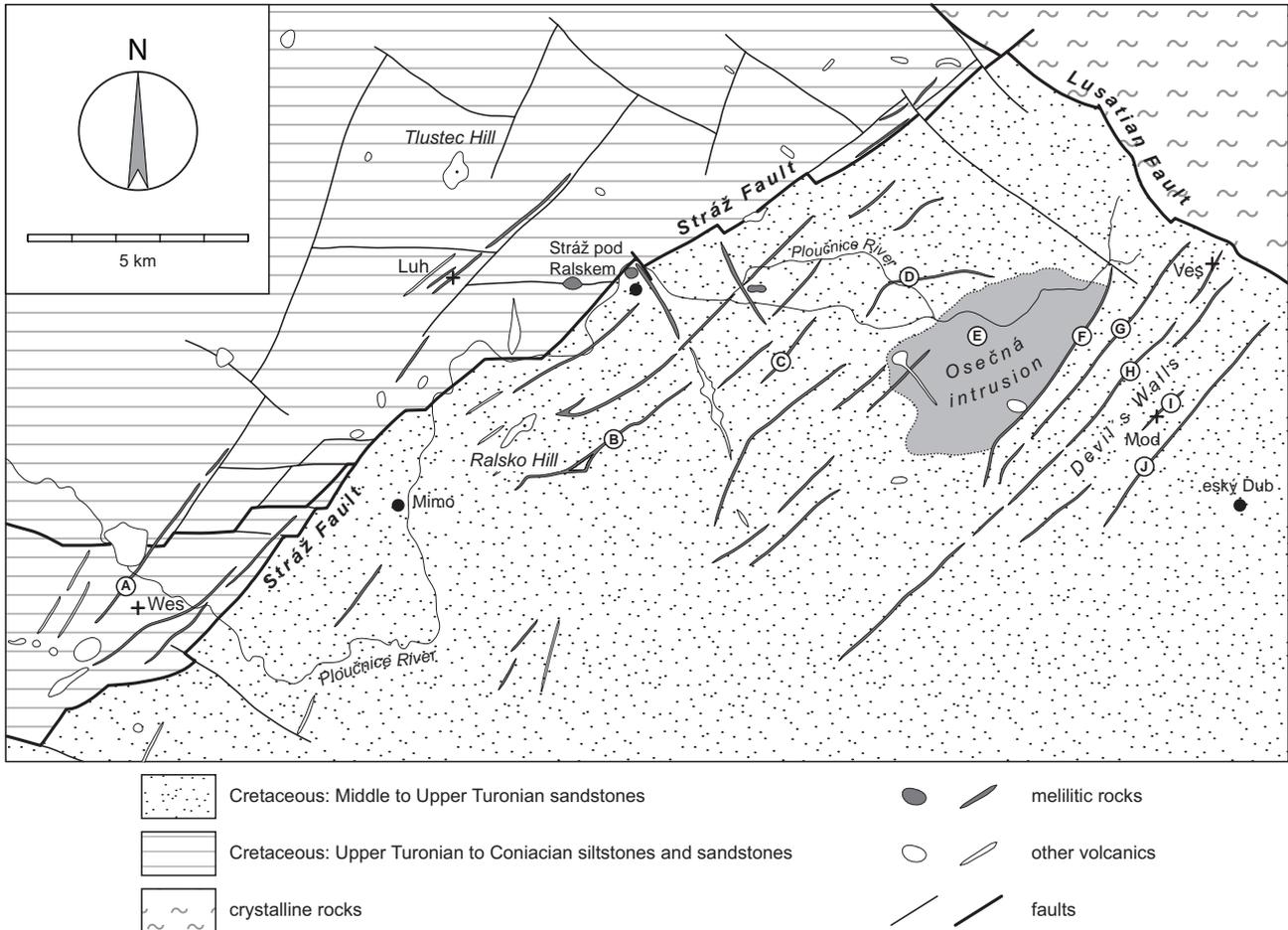


Fig. 1 A simplified geological map of northern Bohemia showing major post-Cretaceous faults and occurrences of Late Cretaceous and Cenozoic volcanic rocks in the Ploučnice River Region. Bodies of melilitic rocks: A – Brenná dyke “der Damm”, B – Great Ralsko dyke, C – Děvín dyke, D – Svárovské návrší dyke, E – Osečná intrusion, F – Western Devil’s Wall, G – Kotel dyke, H – Great Devil’s Wall, I – Pelousek Hill dyke, J – Lesser Devil’s Wall. Typical localities of melilitic rocks: Luh – luhite, Mod – modlibovite, Ves – vesecite. Wes – type locality of “wesselite” (Scheumann 1913, 1922).

(Görlitz area – Seifert et al. 2008 and citations therein). The Delitzsch Complex, occurring outside the Bohemian Massif near Leipzig, consists of Late Cretaceous ultramafic lamprophyres (alnöites and aillikites) and carbonatic rocks (Krüger et al. 2013).

The term “wesselite” was chosen by Scheumann (1913) for rocks of the monchiquite to camptonite rock series from the “Am Knobloschen Grund” dyke at Veselí/Wesseln SW of Mimoň. Their dykes strike mostly NNE–SSW in the Mimoň–Doksy area but show variable strikes further south and west. They have been reported from several places SW of studied area (Zahálka 1905; Ulrych et al. 1990, 1998; Kühn 1999).

3. Analytical methods

The study is based on two sets of samples. The previous dataset comes from an unpublished report for the

Uranium Industry Company and included major-element analyses (Ulrych et al. 1990). Trace-element concentrations of this sample set were acquired within the present study. The newly analysed set of samples comes from Scheumann’s type localities and other outcrops in the area.

New bulk analyses followed the methods described in detail by Johnson and Maxwell (1981) and Potts (1995) and included total digestion of the sample using HF–HClO₄ attack and/or alkaline sintering (both in Pt dishes and/or crucibles) and subsequent determination of major-element oxides by volumetric analyses (Al₂O₃, Fe₂O₃, FeO, MgO, CaO), spectrophotometric and/or inductively coupled plasma spectrometry (ICP-OES) analyses (P₂O₅ and TiO₂), and gravimetric analyses (SiO₂, H₂O). The Na₂O, K₂O and MnO contents were determined using flame atomic absorption spectrometry (FAAS) and/or ICP-OES. The accuracy was monitored analysing the BCR-2 reference material BCR-2 (USGS) and yielded

an average error (1σ) less than $\pm 5\%$ for all determined elements.

Bulk analyses of the previous set of samples were performed at the Faculty of Science, Charles University, Prague, using the methods described in detail above.

Trace-element concentrations in the new set of samples were determined using quadrupole-based ICP-MS X Series 2 (Faculty of Science, Charles University, Prague). Those in the previous set were determined on sector-field ICP-MS Thermo Element 2 at Institute of Geology, Academy of Sciences of the Czech Republic. The analytical protocol for both sets followed the methods of Strnad et al. (2005). In brief, 0.1–0.2 g of sample powder were decomposed in a mixture of HF–HClO₄ acids in teflon beakers and/or Pt crucibles. The remaining residue was thereafter dissolved using 2% HNO₃ and transferred to 100 ml volumetric flask. This solution was finally diluted for ICP-MS measurements. Instrumental calibration was performed using an aqueous multi-element calibration solutions and ¹¹⁵In as an internal standard for the correction of instrumental drift. The precision of the analyses was always better than $\pm 10\%$ (2σ) for X Series 2 ICP-MS and $\pm 5\%$ for Element 2 ICP-MS. The accuracy of $\pm 5\%$ (2σ) was obtained comparing the long-term reproducibility of the USGS reference material BCR-2 against the recommended values (Jochum and Nohl 2008).

The K–Ar age determinations were carried out according to the procedure described in Balogh (1985). Data were calibrated by interlaboratory standards LP-6, GL-O, HD-B1 and Asia 1/65, as well as atmospheric Ar (Odin et al. 1982).

The Sr–Nd isotope compositions were determined in the isotope laboratory at Universität München (Germany) according to the procedures outlined by Hegner et al. (1995). The ¹⁴³Nd/¹⁴⁴Nd ratios were determined with a Finnigan MAT 261 using a dynamic triple mass method and monitoring ¹⁴⁷Sm; Sm isotopes were determined in a static data collection mode. The ¹⁴³Nd/¹⁴⁴Nd ratios were normalized to ¹⁴⁶Nd/¹⁴⁴Nd = 0.7219 and ¹⁴⁷Sm/¹⁵²Sm = 0.56081. The ¹⁴³Nd/¹⁴⁴Nd ratio of the in-house Ames Nd standard solution was 0.512142 ± 12 ($n = 35$), corresponding to 0.511854 in the La Jolla Nd reference material. The ϵ Nd(t) values were calculated with the parameters of Jacobsen and Wasserburg (1980). Present-day ratios for the chondrite uniform reservoir (CHUR) were: ¹⁴⁷Sm/¹⁴⁴Nd =

0.1967, ¹⁴³Nd/¹⁴⁴Nd = 0.512638 (Jacobsen and Wasserburg 1980; ¹⁴³Nd/¹⁴⁴Nd re-normalized to ¹⁴⁶Nd/¹⁴⁴Nd = 0.7219). The ⁸⁷Sr/⁸⁶Sr ratios were determined by a dynamic double mass method, monitoring ⁸⁵Rb, and normalized to ⁸⁶Sr/⁸⁸Sr = 0.1194. The NIST 987 reference material yielded ⁸⁷Sr/⁸⁶Sr = 0.710230 ± 11 ($n = 22$).

4. Polzenite classification and petrographic characteristics

Scheumann (1913) recognized two types of clinopyroxene-free lamprophyres – polzenites – in the PRR: Vesec type (type locality of Vesec) and Modlibohov type (type locality of Modlibohov/Modlibov – Pelousek Hill). The original analyses of the Vesec type, however, came from Malý Bor–Svojkov/Klein Haida–Schwojka and Děvín/Děvín Hill near Hamr (Děvín type). Scheumann (1922) additionally distinguished the third type of “polzenite” – Luhov type (type locality of Luhov/Luh near Stráž pod Ralskem) which, however, corresponds to alnöite.

Of the above mentioned local names, only the general term polzenite was used in some classifications of lamprophyres (Streckeisen 1979; Le Maitre 1989; Rock 1991). Nonetheless, its understanding is often hindered by insufficient comprehension of the original German text (Scheumann 1913, 1922). In the recent classifications of Woolley et al. (1996) and Le Maitre (2002), terms like polzenite, alnöite and even the group of ultramafic lamprophyres are abolished, and melilitic lamprophyres are classified among common melilitic rocks (see the criticism by Tappe et al. 2005). Based on Tappe et al. (2005), we present a proposal of classification of ultramafic lamprophyres, including polzenites from the original localities (Tab. 1).

4.1. Ultramafic lamprophyres and associated melilitic and melilite-bearing volcanic rocks

4.1.1. Ultramafic lamprophyres

The original Scheumann’s definition of melilitic **lamprophyre** – **polzenite** (Scheumann 1913) emphasized its dyke form, porphyritic to doleritic texture, and in

Tab. 1 Proposed classification of ultramafic lamprophyres based on their modal compositions

Rock	Groundmass					Phenocrysts			
	Melilite	Nepheline	Sodalite group	Alkali feldspar	Carbonate	Clinopyroxene	Phlogopite	Olivine	Amphibole
Polzenite	MN	M	m	–	m	–	M/also in grdm	M	–
Alnöite	MN	m	–	–	m	M/also in grdm N	M/also in grdm N	M	–
Aillikite	–	–	–	–	MN	M	M/also in grdm	M	M/also in grdm
Damtjernite	–	mN*	–	mN*	m	M/also in grdm	M/also in grdm	–	–

M – major constituent, m – minor constituent, N – necessary, presence of only one “asterisk” phase, – absent, grdm – groundmass

particular, the absence of clinopyroxene and feldspar, but the presence of h aüyne and phlogopite (for modal composition of polzenites see tab. 1 in Pivec et al. 1998). Polzenite is thus a counterpart of the clinopyroxene-bearing melilitic, foid-free lamprophyric dyke rock – aln oite, defined by Rosenbusch (1887). The basic mineral association of polzenite is: olivine + melilite + nepheline + phlogopite + spinel + calcite ± monticellite, h aüyne (clinopyroxene), perovskite and apatite. Scheumann (1922) described the *polzenite series* formed by olivine-rich polzenite of the Vesec type, transitional member of the Modlibohov type, to olivine-free bergalite from Kaiserstuhl (S ollner 1913). A characteristic macroscopic feature of polzenites is the warty surface (Fig. 2a).

The Vesec type (vesecite) of polzenite was defined by Scheumann (1913) as a rock rich in olivine with monticellite rims and unevenly distributed phlogopite (Fig. 2b). The micro-porphyrific texture is characteristic; olivine and phlogopite micro-phenocrysts (0.1–0.8 mm) are set in a fine-grained groundmass with trachytic texture.

The Modlibohov type (modlibovite) of Scheumann (1913) represents an olivine- and phlogopite-rich rock with zoned foids (h aüyne core and sodalite rim: Ulrych et al. 1991). The texture is micro-porphyrific to doleritic with olivine and phlogopite (0.1–0.8 mm) micro-phenocrysts set in trachytic and poikilitic groundmass (Fig. 2c). Rare irregular clusters of clinopyroxene grains were detected in some modlibovite samples.

The Luhov type (luhite) is compositionally and texturally similar to modlibovite (Scheumann 1922) containing also minerals of the sodalite group (Ulrych et al. 1991). However, it has variable amounts of clinopyroxene micro-phenocrysts (up to 20 vol. %; 0.2–1 mm) forming irregular rims around olivine in association with carbonate (Fig. 2d). Scheumann (1922) classified it to h aüyne–melilite damtjernite. The Luhov type thus belongs to aln oite and not to polzenite (Johannsen 1949; Le Maitre 1989).

Ultramafic melilitic lamprophyres of the PRR are accompanied by other melilitic magmatic rocks forming (i) a hypabyssal olivine melilitolite sheet beneath Ose n a and (ii) a swarm of melilite-bearing olivine nephelinite to olivine melilitite dykes (the Devil’s Walls swarm).

4.1.2. Olivine melilitolite and olivine micro-melilitolite of the Ose n a Complex

The lopolith-like sheet of the Ose n a Complex is not strictly homogeneous. It is a hypabyssal equivalent of near-surface dykes. The central part is composed of olivine melilitolite: olivine + melilite + spinels ± nepheline, phlogopite, monticellite, perovskite and carbonate. The olivine melilitolite is a medium-grained porphyritic rock of lamprophyric character. Phenocrysts are formed by

olivine, melilite and phlogopite (1–5 mm) set in groundmass with ophitic to poikilophitic texture (see photo 2 in Ulrych et al. 1988).

The intrusion contains also rare dykes and pods of coarse-grained rocks such as melilitolite pegmatoids, ijolites, and glimmerites. They gradually pass into the parental olivine melilitolite (Ulrych et al. 2008).

The outermost parts of the melilitolite intrusion formed by its chilled margins and numerous apophyses are composed of olivine micro-melilitolite with micro-phenocrysts of olivine set in groundmass with trachytoid texture. Phenocrysts are formed by olivine with monticellite rims and phlogopite. The groundmass consists of melilite and nepheline ± sodalite–h aüyne ± carbonate; spinels, perovskite and apatite are common accessory minerals. The mineral association and textural characteristics of olivine micro-melilitolite are very similar to those of dykes of ultramafic lamprophyres – polzenites – of the Vesec type.

4.1.3. Melilite-bearing olivine nephelinite (melanephelinite) to olivine melilitite of the Devil’s Walls

The Devil’s Walls dykes are composed of rocks of microporphyritic **melilite-bearing olivine nephelinite** to rare **olivine melilitite** series. The microphenocrysts (0.1–1 mm) are composed of olivine and clinopyroxene, while the fine-grained holocrystalline groundmass (see photo 10 in Ulrych et al. 1998) consists of nepheline, melilite, olivine, clinopyroxene, phlogopite, spinels, apatite and rare perovskite.

4.2. Alkaline lamprophyres

Scheumann (1913) described the melilite-free lamprophyre as biotite–h aüyne basalt of the monchiquite and bekinkinite type, i.e., a melanocratic variety of theralite of Rosenbusch (1907), from “Am Knobloschen Grund” dyke at Vesel ı/Wesseln. The herein verified mineral association of this alkaline lamprophyre of coarse-porphyrific texture and holocrystalline to hemicrystalline groundmass from the original locality of Vesel ı (“wesselite”) is kaersutite (Fig. 2f) + titanian phlogopite + diopside + (serpentinized) olivine phenocrysts (1–10 mm) in groundmass containing clinopyroxene, phlogopite, h aüyne, analcime, titanian magnetite and apatite cemented by weakly anisotropic matter (2.4 wt. % Na₂O, 3.5 wt. % K₂O, 0.5 wt. % SO₃, 12.5 wt. % CO₂). The origin of the latter by decomposition of foids and/or glass is most probable. No melilite or its alteration products were identified. By modal composition, this rock corresponds to monchiquite (Le Maitre 2002). The modal composition of the rock is rather variable probably due to the zoning

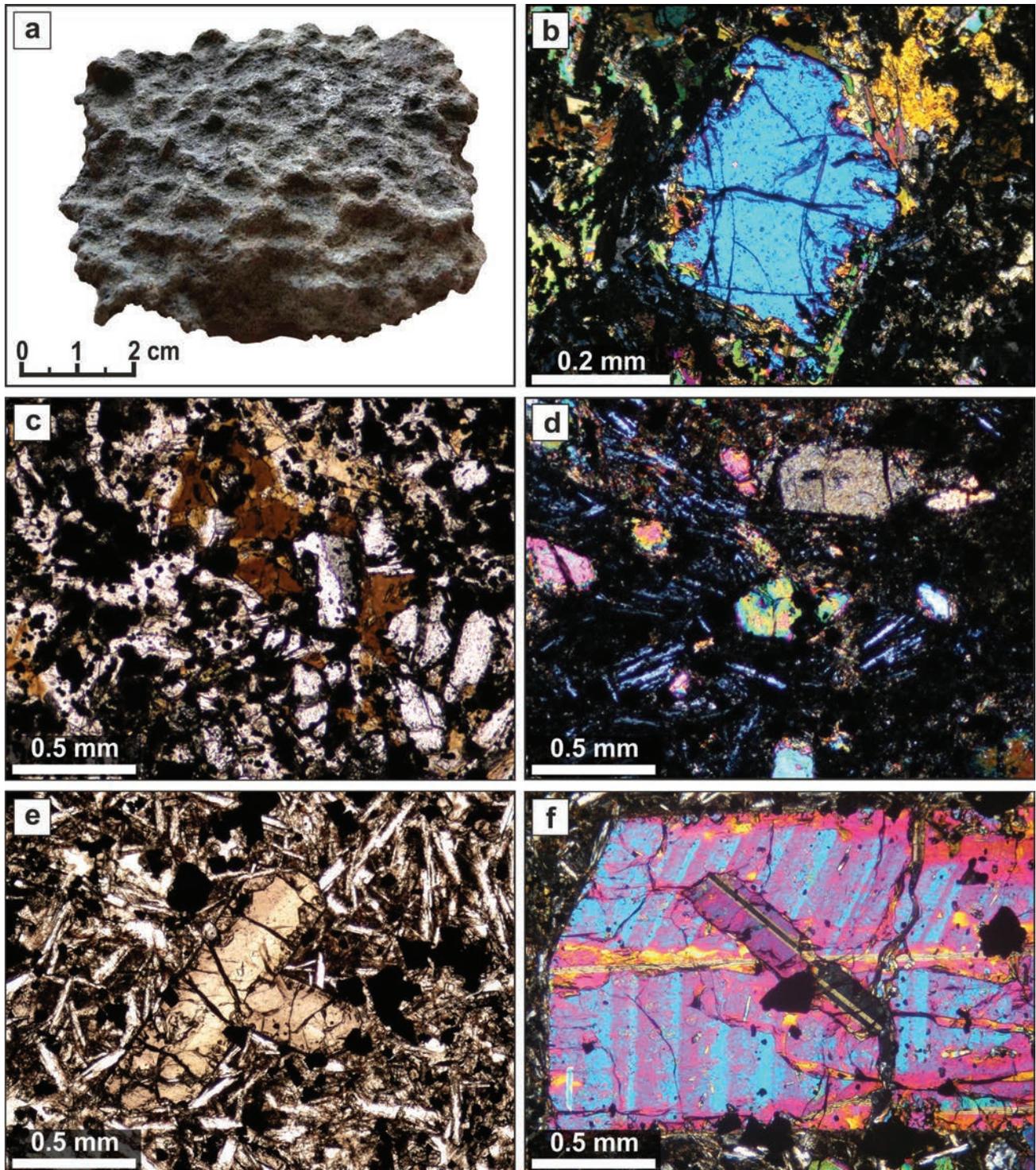


Fig. 2 Petrographic features of the studied lamprophyric rocks (photomicrographs: PPL = plane-polarized light; XPL = cross-polarized light). **a** – Characteristic warty surface of polzenites (locality Vesec); **b** – olivine with monticellite rim in polzenite of the Vesec type (locality Vesec; XPL); **c** – micro-porphyritic texture of the Modlibohov type polzenite characterized by olivine phenocrysts in a groundmass containing poikilitic phlogopite (locality Modlibohov; PPL); **d** – clinopyroxene-rich “polzenite” – alnöite of the Luhov type with laths of melilite-group minerals in groundmass (locality Luhov; XPL); **e** – texture of “wesselite” (camptonite) characterized by phenocrysts of kaersutite surrounded by pilotaxitic feldspar laths of groundmass (locality Veselí; PPL); **f** – twinned kaersutite crystal enclosed in a larger, strongly zoned grain of the same mineral (“wesselite”, locality Veselí; XPL).

of the dyke or a presence of several subparallel dykes. It manifests in particular in the presence of plagioclase or glass/analcime. The **monchiquite** from Veselí (samples 49A and 1/13) and **camptonites** with a low proportion of ternary labradorite to andesine (Ulrych et al. 1998) from the Střelnice dyke at Doksy (sample X-1) and Liščí vrch Hill near Doksy (sample 42) are very similar in composition. Nevertheless, those from Veselí P-1* (Fig. 2e) and Svojkov (sample 35) represent the **leuco-camptonite** variety, and sample 49 is a **mela-camptonite**.

Scheumann (1922) assigned “wesselite” to monchiquite but associated it within the same rock series as polzenites. However, Tröger (1939), Wimmenauer (1973) and Rock (1991) classified “wesselite” directly to polzenites in spite of the absence of melilite.

5. Geochemical characteristics

5.1. Ultramafic lamprophyres, olivine melilitolite, and associated melilitic and melilite-bearing volcanic rocks

Concise geochemical characteristics of melilitic and melilite-bearing rocks of the PRR were presented by Ulrych et al. (1988, 2008) and Pivec et al. (1998). However, new geochemical and K–Ar geochronological data on polzenites from Scheumann’s original localities, and the associated specific alkaline lamprophyre “wesselite” (Ulrych et al. 1990) from this region have not been published yet (Tab. 2). A comparison of geological and, in particular, mineralogical, petrographic and geochemical characteristics of the individual melilitic and melilite-bearing rocks is presented in Tab. 3.

Lamprophyric dyke rocks, including hypabyssal olivine melilitolite, are characterized by very low SiO_2 and alkalis, while CaO and MgO contents are very high (Tab. 2). Melilite-bearing olivine nephelinite to olivine

melilitite is partly richer in SiO_2 , Al_2O_3 and alkalis but poorer in CaO, and MgO (see Tab. 2). Individual types of the melilitic and melilite-bearing rocks plot in the foidite field of the TAS diagram of Le Maitre (2002) (Fig. 3). All samples of these rocks, with the exception of the melilite-bearing olivine nephelinite, are larnite-normative (CIPW norm) and thus can be classified as olivine melilitite *sensu* Brey (1978). In the $\text{CaO} + \text{Na}_2\text{O} + \text{K}_2\text{O}$ wt. % vs. $\text{SiO}_2 + \text{Al}_2\text{O}_3$ wt. % diagram of Le Bas (1989), melilite-bearing rocks mostly fall into the melilitite field (Fig. 4).

According to the criteria of Frey et al. (1978), primary melts of mantle origin should have: (i) Mg\# [$\text{Mg\#} = 100 \text{Mg}/(\text{Mg} + \text{Fe}^{2+})$ where Fe^{2+} is calculated on the basis of $\text{Fe}^{3+}/\text{Fe} = 0.15$] between 68 and 75, (ii) high contents of compatible elements such as Ni (>300 ppm), Co (>50 ppm), Sc (>30 ppm), and Cr (>600 ppm) and (iii) upper mantle xenoliths. Based on these criteria, the melilitic and melilite-bearing volcanic rocks of the PRR represent near-primary melts ($\text{Mg\#} = 73\text{--}81$, Ni = ~130–670 ppm, Co = ~50–70 ppm, Sc = ~20–40 ppm and Cr = ~140–1300 ppm) in equilibrium with the primitive mantle peridotite which underwent only limited low-pressure fractional crystallization (Brey 1978; Alibert et al. 1983). Melilitic rocks have also high CaO/ Al_2O_3 ratios (1.3–2.4), thus exceeding those of typical ocean island and mid-ocean ridge basalts (OIB and MORB) which are both less than unity (Sun and McDonough 1989).

The strongest alteration effects are seen within the Osečná olivine melilitolite intrusion. The following types

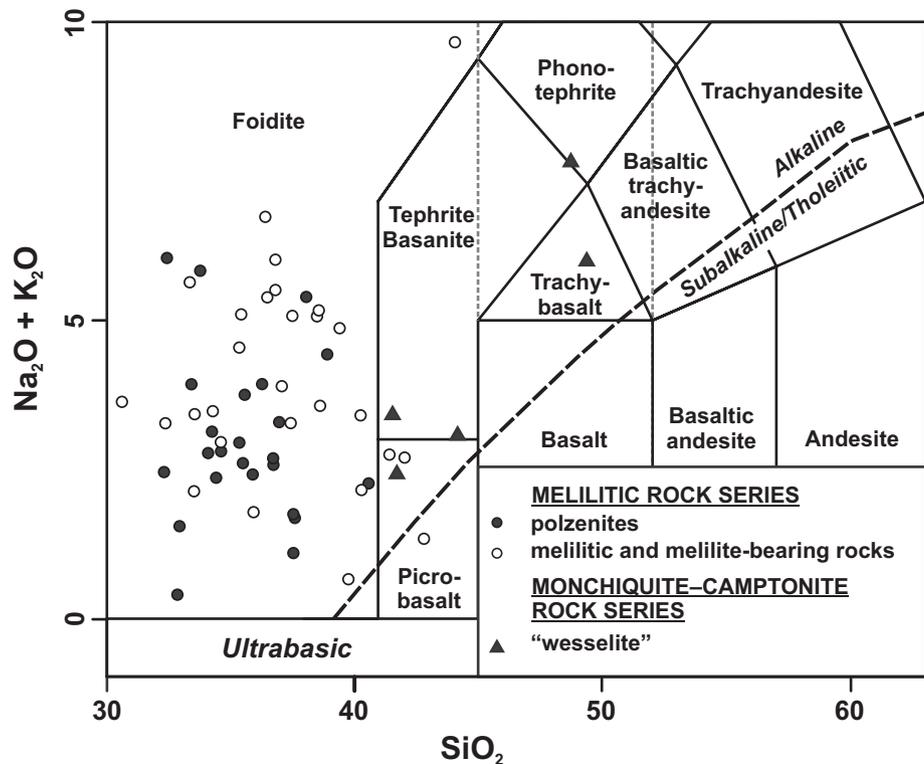


Fig. 3 Total alkali-silica (TAS) diagram (Le Maitre 2002) for the Late Cretaceous and Cenozoic volcanic rocks of the Ploučnice River Region. Data from Ulrych et al. (2008) and this work. Melilitic and melilite-bearing rock group include olivine melilitolite, olivine melilitolite pegmatoid, ijolite, olivine micro-melilitolite, melilite-bearing olivine nephelinite and olivine melilitite.

Tab. 2 Representative whole-rock major- and trace-element analyses of the melilitic and monchiquite–camptonite rock series of the Ploučnice River Region

Locality	Osečná borehole	Osečná borehole	Osečná borehole	Holičky borehole 047112 226 m	Černý vrch Hill at Luhov	Svojkov near Nový Bor	Brenná near Zákupy	Brenná near Zákupy	Vesec near Světlá p. J.	Děvín Hill near Hamr	Hamerský Špičák Hill near Hamr	Svárov Hill near Chrástná	Liberec-Rochlice trench	Pelousek Hill near Modlibohov	Černá Novina borehole near Hamr
Latitude °N	unpubl	unpubl	unpubl	50.680	50.708	50.716	50.649	50.648	50.703	50.693	50.689	50.705	50.755	50.678	50.679
Longitude °E	unpubl	unpubl	unpubl	14.897	14.750	14.595	14.627	14.630	14.983	14.855	14.851	14.891	14.074	14.970	14.868
Sample	87B	88	8	P-12*	25A	33	55	56	P-9*	57	58	22B	36	P-2*	65
Rock type	OM	PHL/PEG	OMM	OMM	POL-OMM	POL-OMM	POL-OMM	POL-OMM	POL-OMM	POL-OMM	POL-OMM	POL	POL	POL	CPOL
SiO ₂ (wt. %)	32.09	34.36	31.15	31.50	31.03	31.75	34.02	34.32	30.32	32.84	32.33	33.07	38.17	35.02	32.60
TiO ₂	2.84	1.74	2.28	2.57	2.13	2.20	2.08	2.17	1.97	2.23	2.23	2.36	2.36	2.10	2.75
Al ₂ O ₃	8.24	6.84	8.13	8.65	8.15	8.42	7.57	8.92	7.61	8.71	8.25	8.21	10.02	8.68	6.62
Fe ₂ O ₃	5.61	3.83	6.21	7.57	6.69	6.44	4.16	4.05	7.15	4.85	4.31	7.46	4.43	6.03	3.52
FeO	5.63	5.74	6.49	2.81	4.99	4.06	5.99	6.00	4.29	5.68	6.07	3.86	6.11	5.41	6.00
MnO	0.23	0.22	0.17	0.17	0.19	0.18	0.16	0.16	0.20	0.18	0.18	0.20	0.18	0.22	0.17
MgO	14.54	7.92	15.87	13.38	16.60	16.04	18.35	15.79	17.99	17.11	17.24	15.81	13.49	15.27	19.12
CaO	19.47	22.01	17.65	17.89	18.05	17.90	15.25	16.74	20.53	19.26	18.82	16.53	15.48	18.38	15.78
Na ₂ O	3.01	3.09	2.13	0.22	4.41	0.96	0.92	0.81	0.47	2.17	1.47	2.30	1.91	1.76	1.65
K ₂ O	2.78	3.51	1.59	1.98	1.70	2.20	1.31	1.40	0.70	1.41	1.39	1.71	0.85	1.39	1.17
P ₂ O ₅	1.67	5.11	1.01	0.95	1.73	1.52	0.87	0.95	1.02	1.46	1.60	1.52	1.07	1.10	1.32
H ₂ O ⁺	2.66	4.17	2.14	4.66	3.37	5.95	6.33	6.16	5.78	2.69	3.50	6.02	3.56	3.34	2.60
H ₂ O ⁻	0.04	0.10	0.66	1.92	0.26	0.54	0.98	0.85	0.32	0.33	0.20	0.26	0.84	0.20	1.50
CO ₂	0.61	0.87	4.09	5.39	0.81	1.17	1.60	0.93	1.02	0.50	1.23	0.91	1.42	0.44	4.52
F ₂	0.23	0.44	0.12	n.d.	0.13	0.21	0.16	0.26	n.d.	0.20	0.20	0.15	0.16	n.d.	0.37
S	n.d.	n.d.	0.16	n.d.	0.26	0.08	0.11	0.17	n.d.	0.10	0.13	0.09	0.10	n.d.	n.d.
Total	99.65	99.95	99.85	99.74	100.50	99.62	99.86	99.68	99.62	99.72	99.15	100.46	100.15	99.64	99.69
Ni (ppm)	280	74	277	216	367	345	470	336	276	460	302	355	265	321	332
Cr	512	25	785	700	596	463	788	625	800	885	430	645	556	559	497
Co	48.7	31.2	58.4	47.0	62.3	66.3	61.9	52.0	61.0	62.0	55.6	61.3	60.9	56.0	59.6
Sc	32.9	56.1	34.0	21.8	32.3	44.5	26.2	29.6	25.5	38.5	32.0	31.2	33.0	29.9	26.0
Rb	56.8	121	53.7	65.0	45.8	55.1	44.4	50.5	25.0	46.3	46.4	57.8	51.6	45.0	36.3
Sr	2012	2513	2046	1277	1764	1225	1741	2246	1897	1522	1226	1269	1770	2225	1110
Y	40	71	29	29	31	35	24	28	27	26	28	33	37	41	20
Zr	758	855	307	345	263	373	260	325	271	242	233	314	308	379	155
Nb	187	79	184	125	154	164	124	144	149	189	157	140	137	179	121
Cs	1.03	2.34	5.46	3.03	0.79	0.60	0.86	1.04	0.46	0.89	0.97	1.00	1.19	0.74	1.89
Ba	959	1391	1138	754	634	917	680	922	541	1459	1017	731	827	2115	698
La	119	26.4	127	83.6	127	141	94.9	98.1	150	99.9	109	115	107	158	85.6
Ce	220	31.0	246	177	235	252	178	179	300	182	201	219	200	280	164
Pr	24.8	3.1	27.8	19.9	26.0	27.4	19.3	19.4	31.0	20.8	22.3	24.7	22.2	27.5	18.4
Nd	92.6	11.8	10.3	77.2	95.2	98.8	73.6	73.5	110	80.1	84.5	91.0	81.5	98.3	71.5
Sm	15.7	3.0	16.7	13.4	15.4	15.5	12.0	12.3	17.2	12.9	14.1	15.0	13.1	16.5	11.8
Eu	4.80	1.66	5.10	3.85	4.59	4.76	3.62	3.85	4.87	3.74	4.27	4.45	3.73	4.94	3.51
Gd	14.3	5.2	14.8	13.9	13.8	14.0	10.7	11.2	18.0	10.3	12.5	13.6	11.6	18.3	10.2
Tb	1.88	0.90	1.81	1.62	1.73	1.75	1.34	1.44	1.82	1.28	1.58	1.67	1.48	2.10	1.25
Dy	9.33	7.15	8.56	6.28	8.12	8.36	6.39	7.13	6.35	6.44	7.55	8.25	7.25	8.38	5.81
Ho	1.61	1.77	1.27	0.97	1.24	1.33	1.00	1.16	0.97	1.07	1.19	1.28	1.19	1.41	0.86
Er	4.04	6.36	3.10	2.87	3.02	3.29	2.45	2.82	2.83	2.58	2.84	3.15	3.02	3.93	2.07
Tm	0.56	0.99	0.31	0.29	0.31	0.37	0.26	0.32	0.25	0.31	0.30	0.35	0.36	0.42	0.19
Yb	3.11	7.15	1.69	1.82	1.72	2.03	1.48	1.77	1.57	1.80	1.67	1.88	1.99	2.54	1.15
Lu	0.53	1.06	0.22	0.25	0.22	0.29	0.19	0.19	0.18	0.25	0.21	0.25	0.28	0.33	0.13
Hf	12.70	13.30	5.55	7.77	4.61	6.15	4.61	5.94	5.75	5.56	4.32	5.20	5.19	8.67	2.93
Ta	7.33	0.26	8.96	7.73	7.06	6.15	5.90	6.35	8.91	3.65	6.34	6.47	5.85	8.11	6.51
Th	11.8	0.33	14.5	10.2	12.7	14.7	10.7	10.6	19.8	12.1	11.9	10.6	15.1	19.5	9.41
U	4.77	3.21	3.25	2.49	3.16	4.11	2.24	2.83	3.76	3.87	3.06	3.03	3.37	5.63	2.34
Mg#	74.1	64.4	73.4	74.5	76.0	77.4	79.8	77.5	77.9	78.1	78.4	75.8	73.7	74.7	81.4
K/Rb	406	241	246	253	308	331	245	230	232	253	249	246	137	256	268
Rb/Sr	0.028	0.048	0.026	0.051	0.026	0.045	0.026	0.022	0.013	0.030	0.038	0.046	0.029	0.020	0.033
ΣREE	512	108	465	403	533	571	405	412	645	424	463	500	455	623	376
La _N /Yb _N	27.4	2.6	53.9	32.9	53.0	49.8	46.0	39.8	68.5	39.8	46.8	43.9	38.6	44.6	53.4
Eu/Eu*	0.96	1.27	0.97	0.86	0.94	0.97	0.96	0.98	0.84	0.96	0.96	0.93	0.91	0.87	0.95
Zr/Hf	59.7	64.3	55.3	44.4	57.0	60.7	56.4	54.7	47.1	43.5	53.9	60.4	59.3	43.7	52.9
Th/U	2.47	0.10	4.46	4.10	4.02	3.58	4.78	3.75	5.27	3.11	3.89	3.50	4.48	3.46	4.02
Nb/Ta	25.5	303.1	20.5	16.2	21.8	26.7	21.0	22.7	16.7	51.8	24.8	21.6	23.4	22.1	18.6
Nb/U	39.2	24.5	56.6	50.2	48.7	39.9	55.4	50.9	39.6	48.8	51.3	46.2	40.7	31.8	51.7
La/Nb	0.64	0.34	0.69	0.67	0.82	0.86	0.77	0.68	1.01	0.53	0.69	0.82	0.78	0.88	0.71
Ba/Nb	5.13	17.65	6.18	6.03	4.12	5.59	5.48	6.40	3.63	7.72	6.48	5.22	6.04	11.8	5.77

Tab. 2 Continued
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Locality	Brenná abandoned quarry	Černý vrch Hill near Luhov	Great Devil's Dyke	Mazova horka Hill near Věsec	Činkův kopec Hill near Janův Důl	Zábrdský kopec Hill near Osečná	Kuřivody near Mímoň trench	Veselí village near Brenná	Veselí "Am Knobloschen Grund"	Veselí "Am Knobloschen Grund"	Střelnice Dyke at Doksy	Liščí vrch Hill near Doksy	Svojkov near Nový Bor	Veselí "Am Knobloschen Grund"	Janův Důl near Osečná
Latitude °N	50.649	50.707	50.675	50.699	50.706	50.680	50.589	50.644	50.643	50.638	50.564	50.555	50.716	50.637	50.703
Longitude °E	14.625	14.743	14.946	14.985	14.954	14.924	14.838	14.632	14.637	14.641	14.671	14.644	14.595	14.639	14.954
Sample	67	P-10*	30	P-4*	P-7*	P-8*	P-5*	49	49A	1/13	X-1	42	35	P-1*	P-3*
Rock type	CPOL	CPOL	MON	MON	MON	MON	MON	MO	MO	MO	MO-CA	MO-CA	CA	CA	CA-BA (?)
SiO ₂ (wt. %)	36.01	33.75	37.98	39.48	38.86	39.28	35.86	40.92	39.81	38.94	39.90	42.30	47.17	47.78	40.57
TiO ₂	4.53	2.04	2.51	2.36	2.65	2.39	2.76	3.96	3.24	3.43	3.52	3.37	2.78	2.58	2.59
Al ₂ O ₃	10.78	8.38	9.14	9.54	9.72	8.61	10.18	11.80	13.68	13.08	11.90	12.78	16.14	16.03	14.34
Fe ₂ O ₃	10.14	5.57	4.11	4.94	6.48	6.53	7.49	2.91	6.83	6.62	5.62	6.20	3.99	4.79	6.57
FeO	7.39	5.17	6.89	6.01	3.57	4.23	4.39	4.92	4.86	5.07	5.75	4.10	5.10	4.72	5.60
MnO	0.22	0.20	0.18	0.18	0.18	0.20	0.21	0.08	0.19	0.16	0.17	0.15	0.15	0.16	0.22
MgO	7.92	17.12	16.48	17.75	9.82	13.82	14.14	15.08	10.62	11.01	10.70	9.00	4.18	4.73	7.79
CaO	15.79	18.38	13.37	13.23	16.78	14.28	12.84	14.54	13.82	13.17	14.53	13.77	9.11	9.38	13.06
Na ₂ O	0.44	1.62	3.05	2.48	1.26	1.60	0.49	1.57	0.80	0.89	1.37	1.84	4.01	2.82	1.36
K ₂ O	1.37	1.48	1.80	1.43	0.65	1.52	0.87	2.40	0.84	1.90	1.63	1.71	3.63	3.34	1.63
P ₂ O ₅	1.35	1.31	0.93	0.69	0.81	1.00	0.91	0.35	0.52	0.53	0.55	0.60	0.52	0.50	1.04
H ₂ O ⁺	3.12	3.68	2.04	0.89	4.21	3.17	6.29	1.38	2.93	3.38	2.80	1.82	2.21	2.38	3.54
H ₂ O ⁻	0.45	0.22	0.33	0.12	1.30	0.60	1.30	0.06	0.84	1.48	0.78	1.06	0.57	0.16	0.66
CO ₂	0.27	0.42	0.56	0.53	3.15	2.32	1.63	0.20	0.53	0.32	0.31	1.03	0.52	0.09	0.51
F ₂	0.14	n.d.	0.12	n.d.	n.d.	n.d.	n.d.	0.25	n.d.	n.d.	0.03	0.28	0.19	n.d.	n.d.
S	n.d.	n.d.	0.04	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	0.05	0.07	0.03	n.d.	n.d.
Total	99.92	99.58	99.53	99.73	99.59	99.70	99.57	100.42	99.51	99.98	99.61	100.08	100.30	99.58	99.62
Ni (ppm)	7	310	575	400	160	262	244	85	72	134	78	111	23	27	49
Cr	8	646	1190	984	595	582	706	242	175	262	184	172	65	75	114
Co	43.1	58.0	65.0	68.0	49.0	52.0	59.0	48.1	43.6	50.5	44.5	48.1	30.4	30.0	41.0
Sc	23.6	25.1	35.8	28.2	29.0	25.5	30.7	52.0	44.4	45.4	42.5	50.2	25.4	19.6	22.8
Rb	49.4	46.0	55.1	47.0	211	126	47.0	133	97.5	154	102	123	121	91.0	101
Sr	2123	1801	1005	813	1130	1135	1621	725	1017	854	1020	816	835	894	1045
Y	29	33	17	20	26	33	31	21	21	21	20	23	23	25	30
Zr	491	295	253	233	292	348	330	278	278	301	269	291	354	308	272
Nb	120	173	121	98	115	143	156	69	66.5	78.7	67.2	70.1	79.8	86.0	103
Cs	0.83	0.63	0.90	0.59	1.75	1.99	1.64	6.14	2.53	1.89	2.99	1.82	1.98	1.11	1.57
Ba	540	1229	988	687	949	1162	1900	936	560	1606	688	834	649	1146	886
La	84.7	126	51.2	52.1	79.7	97.7	103	47.6	49.3	51.3	47.2	53.5	60.2	61.0	68.7
Ce	175	249	97.0	99.5	151	203	202	98.0	102	104	99.8	108	115	118	137
Pr	21.1	25.5	11.5	11.6	16.6	21.5	21.3	11.6	12.1	12.5	11.9	12.7	12.7	13.1	15.5
Nd	82.6	91.6	45.9	44.9	61.4	80.7	77.0	47.2	49.4	50.6	48.2	51.7	49.1	48.2	59.0
Sm	14.4	15.6	8.63	8.70	10.9	14.2	13.2	8.60	9.10	9.27	8.87	9.40	8.50	8.60	10.6
Eu	4.23	4.40	2.41	2.57	3.33	4.17	3.93	2.80	2.74	2.74	2.64	2.98	2.68	2.50	3.15
Gd	12.7	16.2	8.25	8.33	11.1	14.7	14.0	7.80	8.09	8.61	7.96	8.56	7.91	8.92	10.9
Tb	1.61	1.83	0.99	1.02	1.25	1.66	1.60	0.99	1.08	1.03	1.00	1.11	1.05	1.07	1.34
Dy	8.05	6.80	4.34	4.29	5.11	6.88	6.27	5.08	5.43	4.77	4.91	5.62	5.52	4.74	5.83
Ho	1.36	1.13	0.72	0.68	0.92	1.17	1.02	0.86	0.93	0.82	0.88	0.96	0.98	0.84	1.04
Er	3.31	3.06	1.95	1.88	2.56	3.11	2.91	2.11	2.30	2.19	2.18	2.38	2.54	2.49	3.09
Tm	0.41	0.32	0.22	0.20	0.28	0.32	0.29	0.24	0.28	0.26	0.28	0.29	0.33	0.29	0.35
Yb	2.19	1.97	1.39	1.31	1.71	2.08	1.94	1.44	1.63	1.92	1.59	1.65	1.92	2.06	2.41
Lu	0.33	0.26	0.19	0.17	0.24	0.26	0.25	0.18	0.23	0.22	0.22	0.22	0.28	0.28	0.31
Hf	10.20	6.26	5.88	5.73	7.01	7.76	7.68	6.07	6.69	10.2	7.02	6.45	6.68	7.29	6.56
Ta	6.66	8.29	6.31	5.83	6.11	7.89	8.08	3.44	3.71	3.49	3.98	3.67	3.78	4.65	6.16
Th	7.94	13.9	6.87	5.99	9.83	10.6	13.1	4.84	6.89	12.3	7.98	5.59	8.75	10.0	7.33
U	2.12	3.76	1.89	1.62	2.40	3.02	3.74	1.20	1.85	1.45	1.92	1.44	2.20	2.48	1.84
Mg#	50.2	77.9	76.6	78.1	68.7	74.2	72.7	80.8	66.9	67.7	67.5	66.1	50.2	52.4	58.7
K/Rb	230	267	271	253	26	100	154	150	72	102	133	115	249	305	134
Rb/Sr	0.023	0.026	0.055	0.058	0.187	0.111	0.029	0.183	0.096	0.180	0.100	0.151	0.145	0.102	0.097
ΣREE	412	544	235	237	346	451	449	235	245	250	238	259	269	272	319
La _N /Yb _N	27.7	45.9	26.4	28.5	33.4	33.7	38.1	23.7	21.7	19.2	21.3	23.3	22.5	21.2	20.4
Eu/Eu*	0.94	0.84	0.86	0.91	0.92	0.87	0.88	1.03	0.96	0.92	0.94	1.00	0.98	0.87	0.89
Zr/Hf	47.9	47.1	43.0	40.7	41.7	44.8	43.0	45.8	41.6	29.5	38.3	45.1	53.0	42.2	41.5
Th/U	3.75	3.70	3.63	3.70	4.10	3.51	3.50	4.03	3.72	8.45	4.16	3.88	3.98	4.03	3.98
Nb/Ta	18.0	20.9	19.2	16.8	18.8	18.1	19.3	20.1	17.9	22.6	16.9	19.1	21.1	18.5	16.7
Nb/U	56.6	46.0	64.0	60.5	47.9	47.4	41.7	57.7	35.9	54.3	35.0	48.7	36.3	34.7	56.0
La/Nb	0.71	0.73	0.42	0.53	0.69	0.68	0.66	0.69	0.74	0.65	0.70	0.76	0.75	0.71	0.67
Ba/Nb	4.50	7.10	8.17	7.01	8.25	8.13	12.2	13.5	8.42	20.4	10.2	11.9	8.13	13.3	8.60

Explanations for Tab. 2:

type localities of Scheumann (1913) in bold

OM – olivine melilitolite, PHL/PEG – phlogopitite – glimmerite in OM pegmatoid, OMM – olivine micro-melilitolite “chilled margin” of OM, POL-OMM – polzenite – olivine micro-melilitolite (dykes of the vesecite type – Scheumann 1913), POL – polzenite (dykes of the modlibovite type – Scheumann 1913),

CPOL – clinopyroxene “polzenite” (dykes of the luhite type – Scheumann 1913), MON – melilite-bearing olivine nephelinite, MO – monchiquite (“wesselite”), CA – camptonite (“wesselite”), MO-CA – monchiquite–camptonite, CA-BA (?) – camptonite–basanite (?)

Mg# = 100 Mg / (Mg + Fe²⁺), for Fe³⁺/Fe = 0.15; n.d. – not determined, unpubl – unpublished

Analysts: wet analyses by P. Povondra and V. Vonásková (analyses with an asterisk) both from Faculty of Science, Charles University in Prague; trace-element analyses by ICP MS [J. Ďurišová, Institute of Geology, Acad. Sci. CR and L. Strnad, Faculty of Science, Charles University in Prague (analyses with an asterisk)]

Tab. 3 Comparison of geological, mineralogical, petrologic and geochemical characteristics of the melilitic and monchiquite–camptonite rock series of the Ploučnice River Region

	Olivine melilitolite	Olivine micro-melilitolite	P O L Z E N I T E S			Melilite-bearing ol. nephelinites to ol. melilitites	Monchiquite – camptonite	
	Osečná intrusion	Osečná intrusion margin	Vesec type	Modlibohov type	Luhov type (alnöite)	“Wesselite” type		
Age (Ma)*	65 to 61	64	80 to 68	70 to 63	63 to 61	64 to 59	31 to 28	
Geological body	sill, lopolith	marginal part of sill with finger-like apophyses	dykes and cone-sheets?	dykes	dykes	dykes	dykes	
Dip angle		apophyses up to 30°	70–80°	70–80°	?	80–90°	80–90°	
Thickness	20 to 60 m**		up to 5 m	up to 5 m	?	up to 5 m	up to 5 m	
Weathered surface			warty	warty	warty	smooth	smooth	
Texture	porphyritic	micro-porphyritic	porphyritic	porphyritic	porphyritic	micro-porphyritic	porphyritic	
Characteristic mineral association	clinopyroxene	no	no	rare	common	common	common	
	nepheline	common	common	common	common	common	analcime	
	sodalite/haüyne	very rare	very rare	very rare	rare	common	rare	
	monticellite	common	common	common	no	no	no	
	carbonates***	common	common	rare	rare	common	very rare	
Late-magmatic crystallization	phlogopite	common	common	common	common	rare	very rare	
	pegmatoids	common	rare	no	no	very rare	very rare	
	ijolites	rare	no	no	no	no	no	
Postmagmatic transformation	glimmerites	common	rare	no	no	no	no	
		medium to strong	very strong	low to medium	low to medium	very low	very low	
Characteristic chemical composition*	Mg#	73–76	73–75	73–80	73–76	71–81	69–78	50–81
	SiO ₂	31.5–34.9	31.2–31.9	29.5–34.3	33.1–35.0	32.6–36.0	36.3–39.5	38.9–42.3
	alkalis	3.3–5.8	2.2–4.9	2.1–6.1	2.8–5.3	2.8–4.2	2.7–6.3	1.6–3.4
	Al ₂ O ₃	7.8–8.8	7.2–8.7	6.5–8.9	8.2–10.0	6.9–9.1	8.4–10.2	11.8–13.7
	REE	430–510	350–460	410–640	340–620	380–540	220–500	240–260
	La _N /Yb _N	27–43	33–54	40–74	34–48	46–60	19–34	19–24
	K ₂ O, P ₂ O ₅	high	high	high	high	high	high	medium
	H ₂ O, CO ₂ , S	high	high	high	high	high	high	medium
	Ba, Sr	high	high	high	high	high	high	high
Isotopic composition*	(⁸⁷ Sr/ ⁸⁶ Sr) _i	0.7033–0.7042	0.7033	0.7042–0.7049		0.7042	0.7033–0.7034	0.7046–0.7061
	(¹⁴³ Nd/ ¹⁴⁴ Nd) _i	0.51274–0.51278	0.51276	0.51278–0.51279		0.51272	0.51281–0.51282	0.51279
	ε _{Nd}	3.6–4.4	3.9	4.3–4.6		3.2	4.9–5.2	3.3–3.5

* geochemical data from this publication and Ulrych et al. (2008)

** apparent thickness from boreholes

*** secondary > primary

Mg# = 100 Mg/(Mg + Fe²⁺), for Fe³⁺/Fe = 0.15

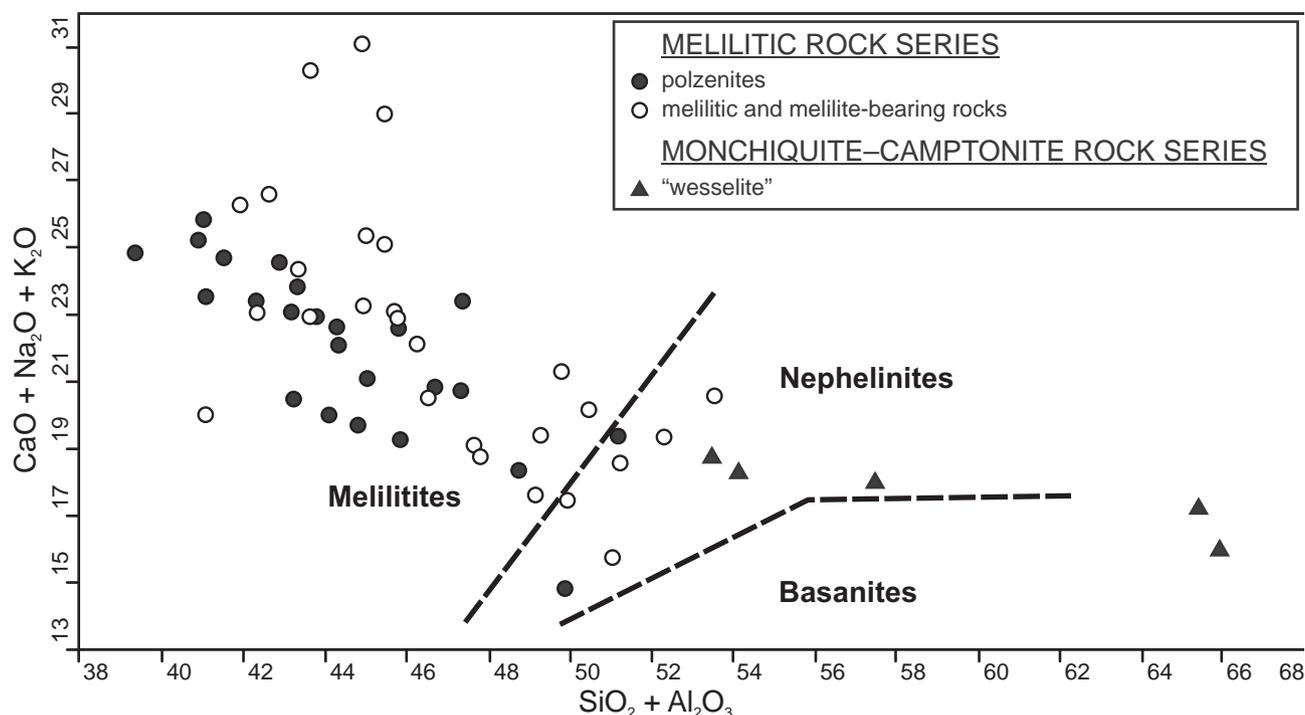


Fig. 4 A binary $\text{CaO} + \text{Na}_2\text{O} + \text{K}_2\text{O}$ vs. $\text{SiO}_2 + \text{Al}_2\text{O}_3$ (wt. %) diagram for the melilitic rock series and monchiquite–camptonite rock series (Le Bas 1989). Data from Ulrych et al. (2008) and this work.

of late-magmatic and post-magmatic enrichments have been recognized:

Ca-Na – (H_2O , P, F) which produced **melilitolite pegmatoids** containing melilite + nepheline + fluorapatite, (F, OH)-bearing titanian andradite, fluorite, wollastonite and thomsonite.

Na – (CO_2 , Zr) which mainly led to the production of **ijolites** (nepheline + Na-rich diopside + calcite, calzirtite and pectolite), see a similar paragenesis of pegmatoids in olivine nephelinites from western Bohemia (Ulrych et al. 2000a, 2005).

K – (Fe^{3+} , Ti, Zr, F, S) resulting in the formation of **glimmerites** formed by (Ti,Ba)-rich phlogopite + (Ti,Zr)-andradite, pyrite and rasvumite.

Nevertheless, the process of metasomatic transformation of melilitic rocks was complex. It included monticellitization, glimmeritization, garnetization and transformation of groundmass to a mixture of carbonates, zeolites, and pectolite (cebolitization). The alterations concentrated to intensively water-saturated sheets of olivine micro-melilitolite in Cretaceous sediments.

Geochemical data on the melilitolite pegmatoids and ijolite dykes presented by Ulrych et al. (2008) display lower contents of transition trace elements such as Cr, Ni and Co compared to the host olivine melilitolite (Tab. 2) and glimmerites. On the other hand, the concentrations of P, Sr, Ba, Y, REE, U, Zr and Nb are slightly elevated.

Primitive mantle-normalized multielement plots for melilitic rocks show enrichment in incompatible ele-

ments (Fig. 5). Melilite-bearing olivine nephelinites of the Devil's Walls dyke swarm have similar geochemical characteristics as those of olivine micro-melilitolites and polzenites of the Osečná Complex, although the concentrations of incompatible elements in the Devil's Walls dykes tend to be lower. The pronounced negative K anomaly in all samples is most characteristic.

Melilitic rocks are enriched in rare earth elements ($\Sigma\text{REE} \sim 220\text{--}640$ ppm) compared to primitive upper mantle (Sun and McDonough 1989) (Tab. 2, Fig. 6). They are similarly enriched in LREE relative to HREE with high La_N/Yb_N ratios. Ultramafic lamprophyres, olivine melilitolites and micro-melilitolites have mostly higher La_N/Yb_N ratios (27–68) than melilite-bearing olivine nephelinites La_N/Yb_N (20–38) although the differences among the bulk-rock compositions are subtle. No samples exhibit substantial negative Eu anomalies ($\text{Eu}/\text{Eu}^* = 0.84\text{--}0.98$).

5.2. Alkaline lamprophyres

Chemical compositions of “wesselite” (**monchiquite to camptonite**) dykes ($\text{Mg}\# = 50\text{--}68$ but also as high as 81!) (Scheumann 1913, 1922; Ulrych et al. 1990, 1998; Kühn 1999) from the PRR display partly variable chemical characteristics (see Tab. 2) and plot prevalently to basanite in the TAS diagram (Fig. 3). The **monchiquite** from Veselí (samples 49A and 1/13) shows medium $\text{Mg}\#$ values of 67–68. **Camptonites** from the Střelnice dyke at Doksy

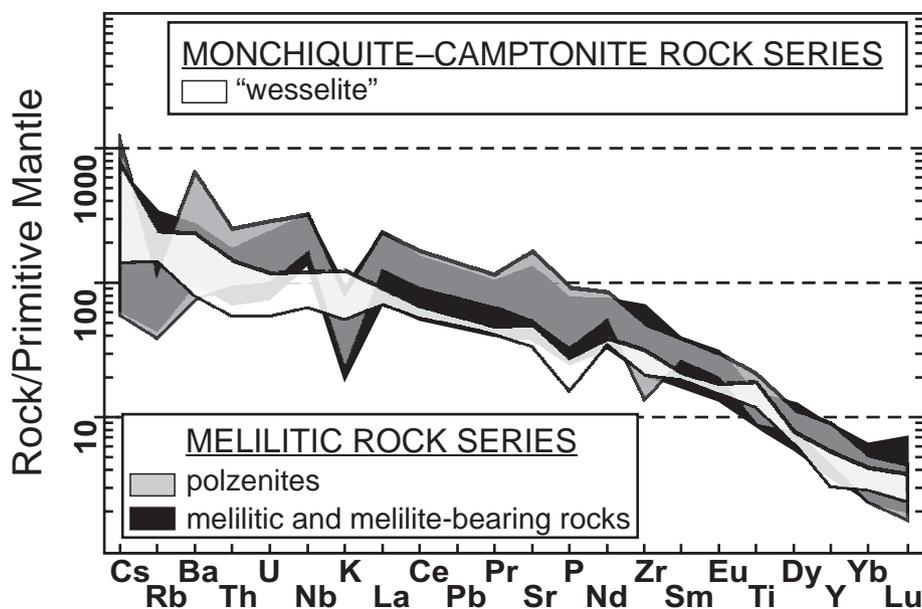


Fig. 5 Primitive mantle-normalized (Sun and McDonough 1989) trace-element patterns for volcanic rocks of the Ploučnice River Region. Shaded fields represent the compositional ranges of the melilitic rock series (lamprophyres – polzenites and other melilitic and melilite-bearing rocks) and the monchiquite–camptonite rock series (“wesselites”). Data from Ulrych et al. (2008) and this work.

(X-1) and Liščí vrch Hill near Doksy (sample 42) are characterized by similar Mg# (66–67). Samples from Veselí P-1* (Fig. 2e) and Svojkov (sample 35) are leucocratic and display very low Mg# of 50–52, while that from Veselí (sample 49) is melanocratic with very high Mg# (81).

Primitive mantle-normalized multielement plots for alkaline lamprophyres of the “wesselite” type show a degree of enrichment in incompatible elements similar to that in the compared melilitic rocks (Fig. 5) but lack a negative K anomaly.

The alkaline lamprophyres are moderately enriched in LREE relative to HREE ($La_N/Yb_N = 20\text{--}24$; Tab. 2, Fig. 6). On average, the enrichment in LREE abundances of these rocks is lower than that of the compared melilitic rocks.

5.3. K–Ar age determinations

New K–Ar age determinations (80–61 Ma, for errors see Tab. 4) of the polzenite types of vesecite (80–68 Ma), modlibovite (70–63 Ma) and luhite (63–61 Ma) from the original localities (Tab. 4) expand the hitherto presented Late Cretaceous to Palaeocene age span for the melilitic volcanic rock series of the PRR (68–59 Ma; Ulrych et al. 2008).

However, K–Ar data for original “wesselite” (monchiquite) dyke sample 1/13 are 30.9 Ma (phlogopite) to 27.8 Ma (kaersutite), while the age of the “wesselite” (camptonite) dyke sample P-1* (whole-rock data) is 23.2 Ma, see Tab. 4.

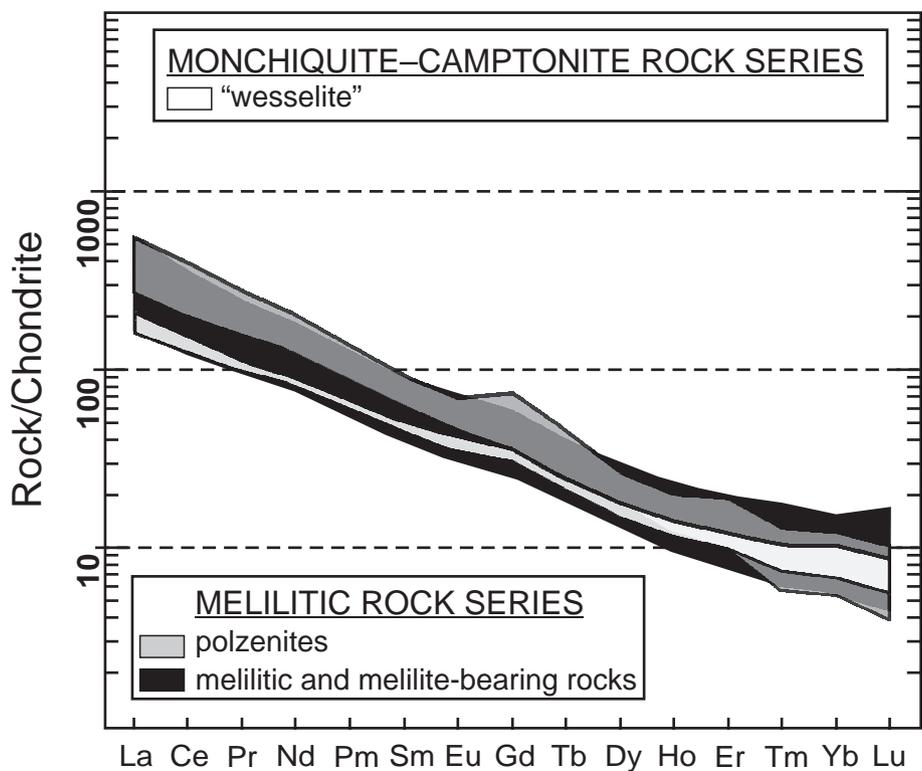


Fig. 6 Chondrite-normalized (Boynton 1984) REE patterns for the studied volcanic rocks. Shaded fields represent the compositional ranges of the melilitic rock series (lamprophyres – polzenites and other melilitic and melilite-bearing rocks) and the monchiquite–camptonite rock series (“wesselites”). Data from Ulrych et al. (2008) and this work.

Tab. 4 K–Ar ages for the studied volcanic rocks

Sample	Locality	Rock type	K (wt.%)	⁴⁰ Ar (rad) 10 ⁻⁶ ccSTP/g	⁴⁰ Ar (rad) (%)	Age ± 1σ (Ma)
Melilitic rock series						
POL-57	Děvín Hill near Hamr	Polzenite – vesecite	0.971	3.067×10 ⁻⁶	47.7	79.5 ± 3.5
P*-2	Modlibohov	Polzenite – modlibovite	1.318	3.636×10 ⁻⁶	40.9	69.6 ± 3.0
P*-10	Luhov	Clinopyroxene “polzenite” – luhite (aillikite)	1.102	2.672×10 ⁻⁶	43.9	61.3 ± 2.6
P*-4	Mazova horka Hill near Vesec	melilite-bearing olivine nephelinite	1.142	2.795×10 ⁻⁶	60.7	61.9 ± 2.4
Monchiquite–camptonite rock series						
1/13-1	Veselí, “Am Knobloschen Grund”	phlogopite in monchiquite – “wesselite”	7.617	9.234×10 ⁻⁶	79.5	30.9 ± 1.2
1/13-2	Veselí, “Am Knobloschen Grund”	kaersutite in monchiquite – “wesselite”	1.569	1.711×10 ⁻⁶	74.6	27.8 ± 1.1
P*-1	Veselí, “Am Knobloschen Grund”	camptonite – “wesselite”	2.993	2.713×10 ⁻⁶	45.3	23.2 ± 1.0
P*-3A	Janův Důl near Osečná	matrix of camptonite – basanite (?)	1.649	1.494×10 ⁻⁶	26.5	23.1 ± 1.2
P*-3B	Janův Důl near Osečná	kaersutite in camptonite – basanite (?)	1.246	1.401×10 ⁻⁶	22.9	28.7 ± 1.6

Vesecite type locality – Děvín Hill, modlibovite type locality – Modlibohov, luhite type locality – Luhov; type localities of Scheumann (1913) in bold

The results correspond well with new K–Ar data for the camptonite–basanite (?) dyke from Janův Důl locality, sample P*-3: 28.7 Ma for kaersutite phenocrysts and 23.1 Ma for groundmass (Tab. 4).

6. Discussion

6.1. Problems of the polzenite and associated rock classification

Regarding the classification of polzenites, there are currently two equivalent approaches. According to the first view, polzenites represent only lamprophyric, i.e. volatile-rich, facies (“lamprophyre clan”) of melilite-bearing group of rocks *sensu* Mitchell (1994) (see Woolley et al. 1996; Le Maitre 2002). This is based on the assumption that the term lamprophyre has no genetic significance and the recognition of a lamprophyre facies is proposed as a means of conveying the concept that some members of a cogenetic petrological series crystallized under volatile-rich conditions (Mitchell 1994). The textural, modal and chemical similarity of polzenites (in particular of the Vesec type) and marginal olivine micro-melilitolite facies of the Osečná intrusion support this view. Nevertheless, the actual Osečná olivine melilitolite sheet can represent a body of lamprophyre or a lamprophyric facies of olivine melilitolite.

According to the second view, polzenites belong to a separate group of ultramafic lamprophyres (Rock 1987, 1991; Le Maitre 1989) as they meet all criteria required by Le Maitre (1989) and Rock (1991) for lamprophyres. However, the polzenites were considered a (more felsic) variant of alnöite by some authors (Tappe et al. 2005). The original Scheumann’s (1913) definition based, contrary to alnöite, on clinopyroxene-free mineral composition was not taken in consideration. Tappe et al. (2005)

introduced a critical modification to the Le Maitre (2002) classification, reintroducing the ultramafic lamprophyres as inequigranular textured rocks with olivine and phlogopite macrocrysts and/or phenocrysts. Nevertheless, they recognized only three end-members: alnöite (essential groundmass phase melilite), aillikite (essential groundmass primary carbonate) and damtjernite (essential groundmass nepheline and/or alkali feldspar) (Tab. 1). With respect to the petrography of clinopyroxene-free polzenite (the Vesec type; typical Mg# = 76–80), this polzenite can be considered a valid end-member of the ultramafic lamprophyre group in accord with original Scheumann’s (1913) definition (Tab. 1).

The similarity of polzenite dykes of the Vesec type and the olivine micro-melilitolite of the Osečná intrusion in porphyritic texture, modal and chemical compositions is striking (see Tab. 3). Notable is also the considerably close mean composition of the volatile-rich porphyritic medium-grained olivine melilitolite of the Osečná intrusion with high phlogopite content (~14 vol. % – Pivec et al. 1998) and polzenites of the Vesec type and olivine micro-melilitolites with only half amount of poikilitic phlogopite (~7 vol. % – Pivec et al. 1998).

Pivec et al. (1986) and Ulrych et al. (1988) considered the Modlibohov type a typical polzenite despite minor clinopyroxene contents (up to 2.5 vol. %). The term “clinopyroxene polzenite” for the Luhov type (clinopyroxene up to 20 vol. %) of Scheumann (1922) is not correct as its modal composition corresponds to alnöite *sensu* Rosenbusch (1887). Scheumann (1913) saw the fundamental differences between polzenite and alnöite in the absence of clinopyroxene and the presence of haüyne in the former rock type. However, only samples of the Vesec type contain no clinopyroxene. The modal composition testifies for a continuously increasing presence of clinopyroxene at the expense of melilite and partly also phlogopite in the modlibovite–luhite series. The Luhov

type alnöites are characterized by higher SiO_2 contents (by ~ 2 wt. %) and lower MgO and CaO contents (by ~ 2 wt. %) compared to the pyroxene-free Vesec type.

The present classifications, including that of Le Maitre (1989) and Rock (1991), do not consider these characteristics (especially the absence of clinopyroxene!) at all. The presence of clinopyroxene and Na-bearing foids is not considered diagnostic for the discrimination between polzenite and alnöite in these classifications, and nepheline is not considered typical of polzenite. Nevertheless, nepheline is the most characteristic mineral of the studied polzenites (Scheumann 1913, 1922; Ulrych et al. 1991; Pivec et al. 1998). The presence of magmatic carbonate is characteristic of alnöite but minor primary carbonate should be present in polzenites as well.

Vesecite is the oldest and most characteristic polzenite type in the PRR. Pfeiffer (1994) published a K–Ar whole-rock age of 71.3 Ma for the famous Zeughausgang olivine melilitite (polzenite) dyke in Saxony. Alnöite of the Delitzsch Complex yielded a Rb–Sr phlogopite age of 73 ± 2 Ma (Krüger et al. 2013). Ultramafic lamprophyre from Ebersbach, Lusatia shows substantially higher ages (K–Ar on phlogopite: 130 ± 5 Ma – Renno et al. 2003a and Ar–Ar on phlogopite: 126.64 ± 0.27 Ma – Renno et al. 2003b). The new K–Ar dating of “wesselite” samples confirms an affinity of the monchiquite–camptonite series to Oligocene volcanic rocks of the tephrite/basanite–phonolite/trachyte rock series (42 to 16 Ma) of the syn-rift period of the northwestern Bohemian Massif (Ulrych et al. 2011).

6.2. Nature, source and evolution of magma

Geological and geochemical characteristics of melilitic and melilite-bearing rocks of the PRR point to a single rock series (Tab. 3). The limited volume and high $\text{Mg}\#$ (73–81) of this **melilitic rock series** suggest an origin by low degrees of fertile (carbonate-enriched) lithospheric mantle melting with residual garnet in a peridotitic source. This is consistent with high La_N/Yb_N values (Tab. 3). Some of the samples reach the $\text{Mg}\#$ up to 80–81 characteristic of mafic cumulates (Rhodes 1981).

A specific type of olivine-free, low- $\text{Mg}\#$ (50), Fe_2O_3 - (10.1 wt. %) and TiO_2 -rich (4.5 wt. %) clinopyroxene-bearing (alnöitic) rock comes from Brenná (sample 67). More evolved melilitites described from Tanzania with lower $\#Mg$ (~ 60), low Ni (~ 100 ppm), but rich in Ti, Ca, Na, K, and poor in Al are interpreted as the result of olivine fractionation of parental nephelinite melt (Dawson et al. 1985).

In contrast, the younger **monchiquite–camptonite dyke series** is characterized by substantially lower $\text{Mg}\#$ (50–67) and lower contents of compatible elements (in particular Ni and Cr) compared to those of the melilitic rock series. They are comparable with the Oligocene basanite lava flows of the České středohoří Mts. ($\text{Mg}\# = 41\text{--}58$) that originated in Ohře/Eger continental rift setting (Ulrych et al. 2002).

Wilson et al. (1995) proposed that olivine melilitites represent characteristic near-primary partial melts of the thermal boundary layer at the base of the lithosphere. The deep Saxothuringian–Moldanubian contact in the Bohemian Massif is marked by a lithospheric thinning to about 80–90 km beneath the (western) Ohře Rift (Babuška and Plomerová 2010; Geissler et al. 2010). According to the model of the Hessian Depression of Wedepohl (1987) the olivine nephelinite/melilitite magma was generated by partial melting at similar depths of *c.* 90 km. The

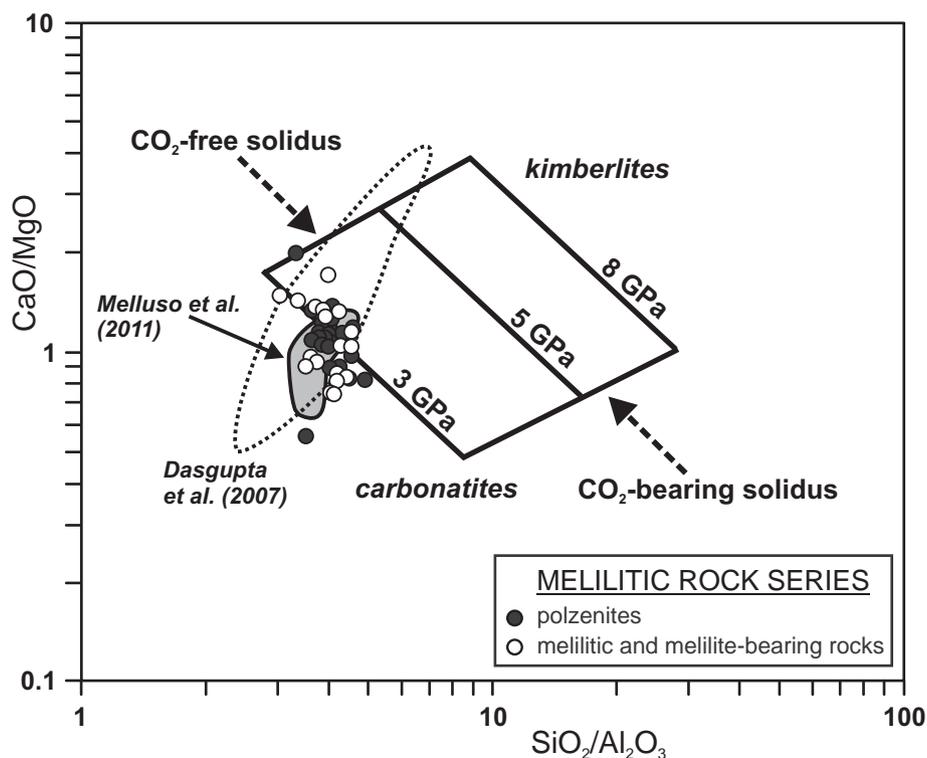


Fig. 7 Rocks of the melilitic rock series of the Ploučnice River Region in the CaO/MgO vs. $\text{SiO}_2/\text{Al}_2\text{O}_3$ diagram after Gudfinnson and Presnall (2005) and the compositional ranges of experimental melts at various pressures and volatile conditions. The experimental compositions of melilitic glasses of Dasgupta et al. (2007) and natural melilitic rocks from Madagascar (Melluso et al. 2011) are shown for comparison. Data from Ulrych et al. (2008) and this work.

Tab. 5 Rb–Sr and Sm–Nd isotopic data for camptonite–monchiquite “wesselite” dykes.

Sample	Locality	Rock	Age (Ma)	Rb (ppm)	Sr (ppm)	$^{87}\text{Rb}/^{86}\text{Sr}$	$^{87}\text{Sr}/^{86}\text{Sr}(\text{m})$	$^{87}\text{Sr}/^{86}\text{Sr}(\text{t})$	Nd (ppm)	Sm (ppm)	$^{147}\text{Sm}/^{144}\text{Nd}$	$^{143}\text{Nd}/^{144}\text{Nd}(\text{m})$	$^{143}\text{Nd}/^{144}\text{Nd}(\text{t})$	$\epsilon_{\text{Nd}}(\text{t})$
49	Veselí	monchiquite	30.9	133	725	0.531	0.704826 ± 10	0.7046	47	8.6	0.1102	0.512793 ± 7	0.512771	3.5
1/13	Veselí	monchiquite	30.9	154	854	0.522	0.706082 ± 13	0.7061	51	9.3	0.1112	0.512791 ± 6	0.512769	3.3

origin of the melilitic magma characterized by extreme silica undersaturation and high Ca content probably involved incongruent melting of clinopyroxene in presence of substantial amounts of H_2O and CO_2 at $P > 26$ kbar (Morgan et al. 1985).

The principal role of CO_2 in the genesis of melilitic volcanic rocks has been established by Brey et al. (1978). The release of CO_2 during magma ascent, due to its well-known low solubility at low pressures, could account for the geochemistry, mineralogy and physical properties of such magmas (Melluso et al. 2011). A comparison between the bulk-rock compositions and experimental results in the $\text{CaO-MgO-Al}_2\text{O}_3\text{-SiO}_2\text{-CO}_2$ system points to a similarity with near-solidus melts in a moderately CO_2 -rich garnet-bearing mantle at ~ 3 GPa (Fig. 7 – Gudfinnson and Presnal 2005). The experimental results indicate a likely depth of melting of at least 90–105 km to produce the variety of magmas. This corresponds with the estimated depths of *c.* 80–90 km of the mantle lithosphere base beneath the Ohře Rift in the western part of the Bohemian Massif (Babuška and Plomerová 2010). Melilitic rocks from Madagascar (Melluso et al. 2011) and experimental melilitite glasses of Dasgupta et al. (2007) share the similar position in Fig. 7 with moderately CO_2 -rich conditions.

The K troughs in primitive mantle-normalized diagrams in Fig. 5 are significant in the interpretation of magma sources (see Melluso et al. 2011 and citations therein). Le Roex et al. (2003) ascribed this feature to the presence of amphibole or phlogopite as residual phases during low-degree melting, characteristic in particular of melilitic rocks (kimberlites).

Variations in the chemical and isotopic compositions of melilitic magmas of northern Bohemia are related to lithospheric thinning or incipient rifting (Ulrych et al. 2006) – the pre-rift period of Ulrych et al. (2008) – and point to local differences in their sources. The primary mineral association of olivine + melilite + spinels \pm clinopyroxene is universal and characteristic of ultramafic melilitic rocks of the Osečná Complex and the associated Devil’s Walls swarm.

Negligible, if any, fractionation can be inferred from the high Mg# values of 73–81 for the melilitic rock series (Ulrych et al. 2008 and new data). Only olivine melilitolite of the Osečná Complex was substantially affected by late-magmatic crystallization and postmagmatic fluids, causing metasomatic transformation (see Section 5.1). Glimmerites with Mg# of 74–76 are chemically similar

to the parental olivine melilitolite while pegmatites (Mg# = 64–66) and ijolites (Mg# = 64) represent evolved metasomatic products enriched in incompatible elements (Ulrych et al. 2008).

The Nd–Sr isotopic data of the melilitic rock series of Ulrych et al. (2008) indicate similar, yet heterogeneous, mantle sources particularly with respect to $^{87}\text{Sr}/^{86}\text{Sr}$ ratios (~ 0.7033 to ~ 0.7049). An alternative interpretation of the relatively high initial $^{87}\text{Sr}/^{86}\text{Sr}$ ratios is a late- and/or post-magmatic alteration of the rocks, lowering their Rb/Sr ratios. The high positive initial ϵ_{Nd} values (+3.2 to +5.2) are interpreted to indicate melting of depleted, although moderately heterogeneous mantle sources precluding significant melt contamination by evolved continental crust. New Nd–Sr data on the monchiquite–camptonite rock series (Tab. 5) show a fair variation in initial $^{87}\text{Sr}/^{86}\text{Sr}$ ratios (~ 0.7046 to ~ 0.7061) corresponding most probably to a substantial postmagmatic alteration of the dykes, nevertheless a minor crustal contamination cannot be excluded. The positive initial ϵ_{Nd} values (+3.3 to +3.5) correspond well to those known from the melilitic rock series.

The presence of monticellite in silica-poor polzenite of the Vesec type and olivine micro-melilitolites of the PRR indicates a transition toward more silica-undersaturated compositions, where clinopyroxene tends to be minor or absent, due to the low silica activity of the primitive magmas characterized by higher Mg# (Melluso et al. 2011).

Rocks of the monchiquite–camptonite dyke series show signs of fractionation. Presence of phenocrysts of kaersutite and phlogopite in this series indicates that the primary magma was enriched in H_2O and other volatile components but impoverished in incompatible elements compared to the melilitic rock series.

The whole-rock major- and trace-element contents imply that the rocks of melilitic dyke association were derived from subcontinental lithospheric mantle sources. The contents of incompatible trace elements in the melilitic rock series are high and variable (Sr = 810–2200 ppm, Ba = 630–2100 ppm), significantly higher than those of average continental crust (Sr \sim 320 ppm, Ba \sim 460 ppm; Rudnick and Gao 2003).

The OIB-like incompatible trace-element ratios, such as high Nb/U (32–68), low La/Nb (0.4–1.0) and variable Ba/Nb (4–12), suggest a limited lithospheric mantle contamination (see Hofmann 1988; Sun and McDonough 1989; Melluso et al. 2011). For the generation of such a melt it is necessary to speculate on an upper mantle source enriched in incompatible elements, together with

a small degree of partial melting of the source material. The minor positive Nb anomaly both in the melilitic and monchiquite–camptonite rock series might indicate Nb fractionation during the passage of primary asthenospheric magma through a metasomatized lithospheric mantle (Hofmann et al. 1986).

Evidence for such metasomatized upper mantle may be provided by xenoliths of glimmerite to mica clinopyroxenite in the polzenite dykes. To the contrary, dunite to harzburgite xenoliths in melilite-bearing olivine nephelinite most likely represent a depleted mantle which underwent moderate to high-degree partial melting. Garnet peridotite, eclogite, norite and ferro-dunite xenoliths are entrained from the local crystalline basement and occur in vent breccias only (Ulrych et al. 2000c). High Ca contents in the melilitic rocks point to a clinopyroxene-rich veined mantle *sensu* Lloyd et al. (1991).

Rocks of the monchiquite–camptonite series, in comparison to the melilitic rock series, are characterized by low contents of compatible elements such as Ni (~20–110 ppm), Co (~30–50 ppm), Sc (~20–50 ppm) and Cr (~70–260 ppm). The contents of incompatible elements, e.g. Sr (~700–1000 ppm) and Ba (~600–1100 ppm) in the rocks of this series are high and variable, though lower than those in the melilitic rock series. Chemical analyses of the “wesselite” from the original locality demonstrate an inhomogeneity and probably a zoned development of the dyke and/or a presence of several dykes (Tab. 2). The incompatible element ratios and the Nb anomaly are largely similar to those of the melilitic rocks series.

The presence of two age-contrasting melilitic (Late Cretaceous to Palaeocene) and tephrite–basanitic (Oligocene) series in northern Bohemia has no analogy either in the Bohemian Massif (Ulrych et al. 2011) or elsewhere in the Circum-Mediterranean anorogenic Cenozoic igneous province of Lustrino and Wilson (2007).

7. Conclusions

- The Cenozoic Central European Volcanic Province includes two main diachronous volcanic rock series in the northern part of the Bohemian Massif: (i) the Late Cretaceous to Early Tertiary ultramafic melilitic rock series including ultramafic clinopyroxene-free lamprophyres – polzenites (80–61 Ma), and (ii) the Oligocene monchiquite–camptonite dyke series (31–28 Ma) free of melilite, belonging to the tephrite–basanite rock series, which was erroneously genetically ascribed to the melilitic series.
- Rare occurrences of ultramafic melilitic and melilite-bearing rocks concentrate to the Osečná Complex associated with the Devil’s Walls swarm. These rock suites occur in the outer parts of the Ohře/Eger Rift at

the junction with the Lusatian Fault Zone in northern Bohemia.

- Olivine melilitolite of the Osečná intrusion with a marginal facies of micro-melilitolite are porphyritic rocks with attributes of lamprophyres (textural characteristics, presence of phlogopite, carbonate, and richness in incompatible elements). Similar modal and chemical compositions of the Vesec type of polzenite dykes and micro-melilitolite of the Osečná intrusion correspond fully to the concept of the lamprophyric facies. Nevertheless, polzenites share all characteristics quoted for (ultramafic) lamprophyres in the literature.
- The steeply dipping dykes associated with the Osečná intrusion are composed of a melilitic ultramafic lamprophyre with volatile-rich mineral association – polzenite. Clinopyroxene-free polzenites of the Vesec type are micro-porphyritic, with phenocrysts being represented by olivine with monticellite rims set in groundmass with poikilitic phlogopite and abundant perovskite. Polzenites of the Modlibohov type are rich in phlogopite, zoned foid of haüyne–sodalite composition, poor in olivine with minor clinopyroxene content, passing to the (clinopyroxene-rich) alnöite of the Luhov type. Carbonates are present in variable amounts (1–10 vol. %) in all types of melilitic rocks.
- The Devil’s Walls swarm is temporally, spatially and genetically associated with the Osečná Complex. Steeply dipping dykes consist predominantly of melilite-bearing olivine nephelinite with rare transitions to olivine melilitite.
- Melilitic and melilite-bearing rocks of the PRR show very high Mg# (73–81). Polzenites of the Vesec type and olivine micro-melilitolites have mean values of 78, thus corresponding rather to mafic cumulates. Peridotite xenoliths are present in all these rocks. Glimmerite to mica clinopyroxenite xenoliths in polzenites represent samples of a metasomatized upper mantle.
- Geological, mineralogical, petrological and geochemical evidence from the melilitic rock suite points to the presence of a singular rock series originated from a common mantle magma source. They can be interpreted as low-degree partial melts of a heterogeneous, clinopyroxene-veined mantle re-fertilized by metasomatic phlogopite ± dolomite and other phases rich in incompatible elements. Mantle metasomatism was probably related to carbonatitic magmatism associated with incipient Neoidic rifting of the lithosphere in the northern and northwestern parts of the Bohemian Massif.

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References

- ABRATIS M, MUNSEL D, VIERECK-GÖTTE L (2009) Melilitite und Melilith-führende Magmatite des sächsischen Vogtlands: Petrographie und Mineralchemie. *Z geol Wiss* 37: 41–79
- ADAMOVIČ J, COUBAL M (1999) Intrusive geometries and Cenozoic stress history of the northern part of the Bohemian Massif. *Geolines* 9: 5–14
- ALIBERT C, LETERRIER R, ALBARÈDE F (1983) The transition from alkali basalts to kimberlites: isotope and trace element evidence from melilitites. *Contrib Mineral Petrol* 82: 176–186
- BABUŠKA V, PLOMEROVÁ J (2010) Mantle lithosphere control of crustal tectonics and magmatism of the western Ohře (Eger) Rift. *J Geosci* 55: 171–186
- BAILEY K, LLOYD F, KEARNS S, STOPPA F, EBY N, WOOLLEY A (2005) Melilitite at Fort Portal, Uganda: another dimension to the carbonate volcanism. *Lithos* 85: 15–25
- BALOGH K (1985) K/Ar dating of Neogene volcanic activity in Hungary. Experimental technique, experience and methods of chronological studies. Unpublished ATOMKI Report D/1, Debrecen, pp 277–278
- BARBIERI M, GHIARA MR, STANZIONE D, VIKLAR LM, PEZ-ZUTI NE, SEGAL SJ (1997) Trace-element and isotope constraints on the origin of ultramafic lamprophyres from Los Alisos (Sierra Subandinas, Northern Argentina). *J South Amer Earth Sci* 10: 39–47
- BEARD AD, DOWNES H, HEGNER E, SABLUKOV SM, VETRIN VR, BALOGH K (1998) Mineralogy and geochemistry of Devonian ultramafic minor intrusions of the southern Kola Peninsula, Russia: implication for the petrogenesis of kimberlites and melilitites. *Contrib Mineral Petrol* 130: 288–303
- BERNSTEIN S, LESLIE AG, HIGGINS AK, BROOKS CK (2000) Tertiary alkaline basic volcanics in the Nunatak Region, North-East Greenland: new discoveries and their similarities to the maymechites of Siberia. *Lithos* 53: 1–20
- BOYNTON WV (1984) Cosmochemistry of the rare earth elements: meteorite studies. In: HENDERSON P (ed) *Rare Earth Element Geochemistry*. Elsevier, Amsterdam, pp 63–114
- BREY G (1978) Origin of olivine melilitites – chemical and experimental constraints. *J Volcanol Geotherm Res* 3: 61–88
- CHAUVEL C, HOFMANN AW, VIDAL P (1992) HIMU–EM: the French Polynesian connection. *Earth Planet Sci Lett* 110: 99–119
- DASGUPTA R, HIRSCHMANN MM, SMITH ND (2007) Partial melting experiments of peridotite + CO₂ at 3 GPa and genesis of ocean islands basalts. *J Petrol* 47: 647–671
- DAWSON JB, SMITH JV, JONES AP (1985) A comparative study of bulk rock and mineral chemistry of olivine melilitites and associated rocks from East and South Africa. *Neu Jb Mineral, Abh* 152: 143–175
- DI BATISTINI G, MONTANINI A, VERNIA L, VENTURELLI G, TONARINI S (2001) Petrology of melilite-bearing rocks from Montefiascone Volcanic Complex (Roman Magmatic Province): new insight into the ultrapotassic volcanism of Central Italy. *Lithos* 59: 1–24
- DUNWORTH EA, WILSON M (1998) Olivine melilitites of the SW German Tertiary volcanic province: mineralogy and petrogenesis. *J Petrol* 39: 1805–1836
- ECKERMANN H VON (1961) The petrogenesis of the Alnö alkaline rocks. *Bull Geol Inst Univ Uppsala* 40: 25–36
- EGOROV LS (1970) Carbonatites and ultrabasic–alkaline rocks of the Maimecha–Kotui region, N. Siberia. *Lithos* 3: 341–359
- FRANCIS EH (1982) Magma and sediment – I. Emplacement mechanism of late Carboniferous tholeiite sills in northern Britain. *J Geol Soc, London* 139: 1–20
- FREY F, GREEN DH, ROY SD (1978) Integrated models of basalt petrogenesis: a study of quartz tholeiites to olivine melilitites from southeastern Australia utilizing geochemical and experimental petrological data. *J Petrol* 19: 463–513
- GEISSLER W H, SODOUDI F, KIND R (2010) Thickness of the central and eastern European lithosphere as seen by S receiver functions. *Geophys J Int* 181: 604–634
- GUDFINN SON GH, PRESNALL DC (2005) Continuous gradations among primary carbonatitic, kimberlitic, melilititic, basaltic, picritic and komatiitic melts in equilibrium with garnet lherzolite at 3–8 GPa. *J Petrol* 46: 1645–1659
- HAMOIS L, MINEAU R (1991) Geochemistry of the Île Cadieux monticellite alnöite, Quebec, Canada. *Can J Earth Sci* 28: 1050–1057

- HEGNER E, VENNEMANN TW (1997) Role of fluids in the origin of Tertiary European intraplate volcanism: evidence from O, H, and Sr isotopes in melilitites. *Geology* 25: 1035–1038
- HEGNER E, WALTER H-J, SATIR M (1995) Pb–Sr–Nd isotopic compositions and trace element geochemistry of megacrysts and melilitites from the Urach volcanic field: source composition of small volume melts under SW Germany. *Contrib Mineral Petrol* 122: 322–335
- HOERNLE K, SCHMINCKE H-U (1993) The petrology of the tholeiites through melilite nephelinites on Grand Canaria, Canary Islands: crystal fractionation, accumulation, and depths of melting. *J Petrol* 34: 573–597
- HOFMANN AW (1988) Chemical differentiation of the Earth: the relationship between mantle, continental crust, and the oceanic crust. *Earth Planet Sci Lett* 90: 297–314
- HOFMANN AW, JOCHUM KP, SEUFERT M, WHITE WM (1986) Nb and Pb in oceanic basalts: new constraints on mantle evolution. *Earth Planet Sci Lett* 79: 33–45
- ISHIKAWA A, MARUYAMA S, KOMIYA T (2004) Layered lithosphere mantle beneath the Ontong Java Plateau: implications from xenoliths in alnöite, Malaita, Solomon Islands. *J Petrol* 45: 2011–2044
- JACOBSEN SB, WASSERBURG GJ (1980) Sm–Nd isotopic evolution of chondrites. *Earth Planet Sci Lett* 50: 139–155
- JOCHUM KP, NOHL U (2008) Reference materials in geochemistry and environmental research and the GeoReM database. *Chem Geol* 253: 50–53
- JOHANNSEN A (1949) *A Descriptive Petrography of the Igneous Rocks IV*. University Press, Chicago, pp 1–523
- JOHNSON WM, MAXWELL JA (1981) *Rock and Mineral Analysis*. Wiley, New York, pp 1–489
- KELLER J (1984) Der jungtertiäre Vulkanismus Südwestdeutschlands: Exkursionen in Kaiserstuhl und Hegau. *Fortschr Mineral* 62: 2–35
- KELLER J, BREY G, LORENZ V, SACHS P (1990) Pre-conference excursion 2A: Volcanism and petrology of the Upper Rhinegraben (Urach–Hegau–Kaiserstuhl). IAVCEI International Volcanological Congress, Mainz 1990, pp 1–31
- KELLER J, ZAYTSEV A, WIEDENMANN D (2006) Primary magmas at Oldoinyo Lengai: the role of olivine melilitites. *Lithos* 91: 150–172
- KRÜGER JC, ROMER RL, KÄMPF H (2013) Late Cretaceous ultramafic lamprophyres and carbonatites from the Delitzsch Complex, Germany. *Chem Geol* 353: 140–150
- KÜHN P (1999) The “Střelnice” dyke in Doksy (north Bohemia). *Bull mineral-petrolog Odd Nár Muz (Praha)* 7: 169–172
- LE BAS MJ (1989) Nephelinitic and basanitic rocks. *J Petrol* 30: 1299–1312
- LE MAITRE RW (1989) *A Classification of Igneous Rocks and Glossary of Terms*. Blackwell Scientific, Oxford, pp 1–193
- LE MAITRE RW (2002) *Igneous Rocks. A Classification and Glossary of Terms*. 2nd Edition. Cambridge University Press, Cambridge, pp 1–236
- LE ROEX AP, BELL DR, DAVIS, P (2003) Petrogenesis of group I kimberlites from Kimberley, South Africa: evidence from bulk-rock chemistry. *J Petrol* 44: 2261–2286
- LLOYD FE, HUNTINGDON AT, DAVIES GR, NIXON PH (1991) Phanerozoic volcanism of southwest Uganda. A case for regional K and LILE enrichment of the lithosphere beneath a domed and rifted continental plate. In: KAMPUNZU AB, LUBALA RT (eds) *Magmatism in Extensional Structural Settings*. Springer, Berlin, pp 23–72
- LUSTRINO M, WILSON M (2007) The Circum-Mediterranean anorogenic Cenozoic igneous province. *Earth Sci Rev* 81: 1–65
- MAALOE S, JAMES D, SMEDLEY P, PETERSEN S, GARMANN LB (1992) The Koloa volcanic suite of Kaunai, Hawaii. *J Petrol* 33: 761–784
- MELLUSO LM, LE ROEX AP, MORRA VM (2011) Petrogenesis and Nd-, Pb-, Sr-isotope geochemistry of the Cenozoic olivine melilitites and olivine nephelinites (“ankaraites”) in Madagascar. *Lithos* 127: 505–521
- MITCHELL RH (1994) The lamprophyre facies. *Mineral Petrol* 51: 137–146
- MORGAN JW, CZAMANSKE GK, WANDLESS GA (1985) Origin and evolution of the alkalic ultramafic rocks in the Coyote Peak diatreme, Humboldt County, California. *Geochim Cosmochim Acta* 49: 749–759
- ODIN GS AND 35 AUTHORS (1982) Interlaboratory standards for dating purposes. In: ODIN GS (ed) *Numerical Dating in Stratigraphy*. Wiley & Sons, Chichester, pp 123–149
- PEARCE NGJ, LENG MJ, EMELEUS CH, BEDFORD CM (1997) The origins of carbonatites and related rocks from the Gronnedal-Ika nepheline syenite complex, South Greenland: C–O–Sr isotope evidence. *Mineral Mag* 61: 515–529
- PFEIFFER L (1978) Beitrag zur Petrochemie der sächsischen Tertiärvulkanite. *Freiberg Forsch H C* 333: 1–64
- PFEIFFER L (1994) Der Tertiäre Magmatismus im Erzgebirge und in den benachbarten Gebieten der Tschechischen Republik. *Beih Eur J Mineral* 6: 179–228
- PIROMALLO C, GASPERINI D., MACERA P, FACCENNA C (2008) A late Cretaceous contamination episode of the European–Mediterranean mantle. *Earth Planet Sci Lett* 268: 15–27
- PIVEC E, POVONDRA P, RUTŠEK J, ULRYCH J (1986) Petrology and geochemistry of the Osečná intrusion in the Ještěd foothills (Northern Bohemia). *Acta Montana* 74: 23–31
- PIVEC E, ULRYCH J, HÖHNDORF A, RUTŠEK J (1998) Melilitic rocks from northern Bohemia: geochemistry and mineralogy. *Neu Jb Mineral, Abh* 1998: 312–339
- POTTS PJ (1995) *A Handbook of Silicate Rock Analysis*. Blackie Academic and Professional, London, pp 1–622
- PRODEHL C, MUELLER S, HAAK V (1995) The European Cenozoic rift system. In: OLSEN KH (ed) *Continental Rifts: Evolution, Structure, Tectonics*. Developments in Geotectonics 25: Elsevier, Amsterdam, pp 133–212

- RASS IT (2008) Melilite-bearing and melilite-free rock series in carbonatite complexes: derivatives from separate primitive melts. *Canad Mineral* 46: 951–969
- RENNO AD, STANEK KP, LOBST R, PUSHKAREV Y (2003a) A new lamprophyre species from the Klunst quarry (Ebersbach, Lusatia, Germany) – geochemical and petrological implications. *Z geol Wiss* 31: 1–20
- RENNO AD, HACKER BR, STANEK KP (2003b) An Early Cretaceous (126 Ma) ultramafic alkaline lamprophyre from the Quarry Klunst (Ebersbach, Lusatia, Germany). *Z geol Wiss* 31: 31–36
- RHODES JM (1981) Characteristics of primary basaltic magmas. In: *Basaltic Volcanism Study Project: Basaltic Volcanism of the Terrestrial Planets*. Pergamon Press, New York, pp 128–147
- RUDNICK RL, GAO S (2003) Composition of the continental crust. In: RUDNICK, RL (ed) *The Crust. Treatise on Geochemistry 3*: Elsevier–Pergamon, Oxford, pp 1–64
- ROCK NMS (1987) The nature and origin of lamprophyres: an overview. In: FITON JG, UPTON BGJ (eds) *Alkaline Igneous Rocks*. Geological Society London Special Publications 30: 191–226
- ROCK NMS (1991) *Lamprophyres*. Blackie and Sons Ltd., Glasgow and London, pp 1–285
- ROSENBUSCH H (1887) *Mikroskopische Physiographie der Mineralien und Gesteine*. Vol. II Massige Gesteine, 2nd edition. Schweizerbart'sche Verlag, Stuttgart, pp 1–877
- ROSENBUSCH H (1907) *Mikroskopische Physiographie der Mineralien und Gesteine*. Vol II Massige Gesteine, 4th edition. Schweizerbart'sche Verlag, Stuttgart, pp 1–716
- SCHUEMANN KH (1913) Petrographische Untersuchungen an Gesteinen des Polzengebietes in Nord-Böhmen. *Abh kön Sächs Gesel Wiss math–phys Kl* 32: 607–776
- SCHUEMANN KH (1922) Zur Genese alkalisch-lamprophyrischen Ganggesteine. *Zbl Mineral Geol Paläontol* 2: 495–545
- SEIFERT W, BÜCHNER J, TIETZ O (2008) Der „Melilithbasalt“ von Görlitz im Vergleich mit dem Melilithit vom Zeughausgang: Retrospektive und neue mineral-chemische Ergebnisse. *Z geol Wiss* 36: 155–176
- SÖLLNER J (1913) Über den Bergalith, ein neues melilitreiches Ganggestein aus dem Kaiserstuhl. *Mitt Badisch geol Landesanst* 7: 2
- STÖRMER JC, NICHOLLS J (1978) XLFrac: a program for interactive testing of magmatic differentiation models. *Comput and Geosci* 87: 51–64
- STRECKEISEN A (1979) Classification and nomenclature of volcanic rocks, lamprophyres, carbonatites and melilitic rocks. *Neu Jb Mineral, Abh* 134: 1–14
- STRNAD L, MIHALJEVIČ M, ŠEBEK O (2005) Laser ablation and solution ICP-MS determination of rare earth elements in USGS BIR-1G, BHVO-2G and BCR-2G glass reference material. *Geostand Geoanal Res* 29: 303–314
- SUN SS, McDONOUGH WF (1989) Chemical and isotopic systematics of oceanic basalts: implications for mantle compositions and processes. In: SAUNDERS AD, NORRIS MJ (eds) *Magmatism in the Ocean Basins*. Geological Society London Special Publications 42: 313–345
- TAPPE S, FOLEY SF, JENNER GA, KJARSGAARD BJ (2005) Integrating ultramafic lamprophyres into the IUGS classification of igneous rocks: rationale and implications. *J Petrol* 46: 1893–1900
- TRÖGER WE (1939) Über Theralith and Monchiquit. *Zbl Miner Geol Paläont Abt A*: 80–94
- ULRYCH J, POVONDRA P, RUTŠEK J, PIVEC E (1988) Melilitic and melilite-bearing subvolcanic rocks from the Ploučnice river region, Czechoslovakia. *Acta Univ Carol, Geol* 195–231
- ULRYCH J, POVONDRA P, HUSPEKA J, PIVEC E, RUTŠEK J (1990) Chemical composition of melilitic volcanics of northern part of the Bohemian Massif. Report of the Projects II-4-1/07 and II/4/4/05. Institute of Geosciences, Faculty of Science, Charles University, Prague, pp 1–123 (in Czech)
- ULRYCH J, PIVEC E, POVONDRA P, RUTŠEK J (1991) Rock-forming minerals of polzenite and cognate melilitic rocks from northern Bohemia, Czechoslovakia. *Acta Univ Carol, Geol Fediuk Vol 2*: 139–163
- ULRYCH J, PIVEC E, ŽÁK K, BENDL J, BOSÁK P (1993) Alkaline and ultramafic carbonate lamprophyres in Central Bohemian Carboniferous basins, Czech Republic. *Mineral Petrol* 48: 65–81
- ULRYCH J, POVONDRA P, PIVEC E, RUTŠEK J, BENDL J, BILIK I (1996) Alkaline ultramafic sill at Dvůr Králové nad Labem, Eastern Bohemia: petrological and geochemical constraints. *Acta Univ Carol, Geol* 53–79
- ULRYCH J, PIVEC E, LANG M, LLOYD FE (2000a) Ijolitic segregations in melilite nephelinite of Podhorní vrch volcano, Western Bohemia. *Neu Jb Mineral, Abh* 175: 317–348
- ULRYCH J, CAJZ V, PIVEC E, NOVÁK JK, NEKOVAŘÍK Č, BALOGH K (2000b) Cenozoic intraplate alkaline volcanism of Western Bohemia. *Stud Geophys Geod* 44: 346–351
- ULRYCH J, PIVEC E, POVONDRA P, RUTŠEK J (2000c) Upper-mantle xenoliths in melilitic rocks of the Osečná Complex, North Bohemia. *J Czech Geol Soc* 45: 79–93
- ULRYCH J, SVOBODOVÁ J, BALOGH K (2002) The source of Cenozoic volcanism in the České Středohoří Mts., Bohemian Massif. *Neu Jb Mineral, Abh* 177: 133–162
- ULRYCH J, LLOYD FE, BALOGH K, HEGNER E, LANGROVÁ A, LANG M, NOVÁK JK, ŘANDA Z (2005) Petrogenesis of alkali pyroxenite and ijolite xenoliths from the Tertiary Loučná–Oberwiesenthal Volcanic Centre, Bohemian Massif in the light of new mineralogical, geochemical and isotopic data. *Neu Jb Mineral, Abh* 182: 57–79
- ULRYCH J, PEŠEK J, ŠTĚPÁNKOVÁ-SVOBODOVÁ J, BOSÁK P, LLOYD FE, VON SECKENDORFF V, LANG M, NOVÁK JK (2006) Permo–Carboniferous volcanism in late Variscan continental basins of the Bohemian Massif (Czech

- Republic): geochemical characteristic. *Chem Erde* 66: 37–56
- ULRYCH J, DOSTAL J, HEGNER E, BALOGH, K, ACKERMAN L (2008) Late Cretaceous to Paleocene melilitic rocks of the Ohře/Eger Rift in northern Bohemia, Czech Republic: insights into the initial stages of continental rifting. *Lithos* 101: 141–161
- ULRYCH J, JELÍNEK E, ŘANDA Z, LLOYD FE, BALOGH K, HEGNER E, NOVÁK JK (2010) Geochemical characteristics of the high- and low-Ti basaltic rocks from the uplifted shoulder of the Ohře (Eger) Rift, Western Bohemia. *Chem Erde* 70: 319–333
- ULRYCH J, DOSTAL J, ADAMOVIČ J, JELÍNEK E, ŠPAČEK P, HEGNER E, BALOGH K (2011) Recurrent Cenozoic volcanic activity in the Bohemian Massif (Czech Republic). *Lithos* 123: 133–144
- WEDEPOHL KH (1987) Kontinentaler Intraplatten-Vulkanismus am Beispiel der tertiären Basalte der Hessischen Senke. *Fortschr Mineral* 65: 19–47
- WILKINSON JFG, STOLZ AJ (1983) Low-pressure fractionation of strongly undersaturated alkaline ultrabasic magma: the olivine–melilite–nephelinite at Moiliili, Oahu, Hawaii. *Contrib Mineral Petrol* 83: 363–374
- WILSON M, ROSENBAUM JM, DUNWORTH EA (1995) Melilitites: partial melts of the thermal boundary layer? *Contrib Mineral Petrol* 119: 181–196
- WIMMENAUER W (1973) Lamprophyre, Semilamprophyre und anchibasaltische Ganggesteine. *Fortschr Mineral* 51: 3–67
- WOOLLEY AR, BERGMAN CC, EDGAR AD, LE BAS MJ, MITCHELL RH, ROCK NMS, SCOTT-SMITH BH (1996) Classification of lamprophyres, lamproites, kimberlites, and the kalsilitic, melilitic and leucitic rocks. *Canad Mineral* 34: 175–186
- WURM F (1883) Über das Vorkommen von Melilithbasalt zwischen Böhmisch-Leipa und Böhmisch-Aicha. *Sitzber Kön Böhm Gesell Wiss (Prag)* 2: 1–8
- WURM F (1884) Die Teufelsmauer zwischen Oschitz und Böhm.-Aicha. *Nordböhmischer Excursions-Club, Česká Lípa*, pp 1–35
- ZAHÁLKA B (1905) Eruptive rocks from the Mělník and Mšeno neighbourhoods. *Věst Král České Spol Nauk v Praze III*: 1–79 (in Czech with German abstract)
- ZHABIN AG, SURINA NP (1970) Petrology of dikes, sills and pipes of the Maymecha–Kotuyi Province. *Nauka, Moscow*, pp 1–202 (in Russian)