Nomenclature of wöhlerite-group minerals

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Abstract

A nomenclature and classification scheme for wöhlerite-group minerals has been established. The general formula of minerals belonging to this group is given by $X_8(Si_2O_7)_2W_6$, where $X = Na^+$, $Ca^{2+}$, $Mn^{2+}$, $Ti^{4+}$, $Zr^{4+}$ and $Nb^{5+}$; and $W = F^-$ and $O^{2-}$. In addition, they may incorporate significant amounts of $Mg^{2+}$, $Fe^{2+}$, $Y^{3+}$ and $REE^{3+}$, where $REE$ are the lanthanides. The main structural feature of these minerals is the four-columns-wide octahedral walls, which are interconnected through corner sharing and via the disilicate groups. The wöhlerite-group minerals crystallise in different unit-cell settings and symmetries, depending on the cationic ordering in the octahedral walls and the relative position of the disilicate groups. Different combinations of $X$ and $W$ constituents should be regarded as separate mineral species. In the case of coupled heterovalent substitutions at different crystallographic sites, it is advised to use the site-total charge approach to determine the correct end-member composition. Due to their structural and chemical features, wöhlerite-group minerals can easily form crystals with several micro domains, showing different crystal structures and chemical compositions. In addition, the crystallisation of polytypes is relatively common, although they should not be regarded as distinct mineral species. To date, ten minerals belonging to the wöhlerite group are considered as valid species: baghdadite, burpalite, cuspidine, hiortdahlite, janhaugite, låvenite, moxuanxueite, niocalite, normandite and wöhlerite. Låvenite and normandite are isostructural and are respectively the Zr and Ti end-members of a solid-solution series. Marianoite is discredited, as it is corresponds to wöhlerite. The ideal formula of hiortdahlite is revised as $Na_3Ca_4(Ca_8Zr_3)Zr(Si_2O_7)_2O_2F_6$ with one cationic site characterised by a valency-imposed double site-occupancy. These changes have been approved by the IMA–CNMNC (Proposal 20–D).

Keywords: wöhlerite group, cuspidine, wöhlerite, marianoite, hiortdahlite, nomenclature

Introduction

The first mention of wöhlerite in the literature was made by Scheerer (1843) who studied the mineralogical paragenesis of the syenite pegmatites occurring on Lovøya island, Brevig area, Langesundsfjord, Norway. The chemical analysis performed by Scheerer (1843) indicated that wöhlerite is a silicate containing mainly Na, Ca, Zr and Nb, as well as minor amounts of Mg, Mn and Fe. At the time of wöhlerite’s discovery niobium was not officially approved as a distinct chemical element, and consequently Scheerer (1843) had erroneously reported niobium in wöhlerite as tantalum. Among the wöhlerite-group minerals, cuspidine was the first to have its crystal structure solved (Smirnova et al., 1955) and the wöhlerite group is sometimes mentioned as the cuspidine group in the literature (e.g. Merlino and Perchiazzi, 1988; Chakhmouradian et al., 2008). However, as wöhlerite, the first described species of the group the name should be wöhlerite group in accordance with Mills et al. (2009). Merlino and Perchiazzi (1988) demonstrated that the nature of the crystal structure of the wöhlerite-group minerals (WGM) permits the crystallisation in different unit-cell settings and the formation of polytypes. In addition, they identified ten different structure-types that are possible within the fixed cell dimension $a \approx b \approx 10.5$ Å and $c \approx 7.3$ Å. The WGM can form multi-domain crystals, as for instance in ‘guarinite’ from Monte Somma, Italy (Bellezza et al., 2012).

The new definition of the wöhlerite group has been approved by the Commission on New Minerals, Nomenclature and Classification (CNMNC) of the International Mineralogical Association (IMA) (Proposal 20–D; Miyawaki et al., 2020). The wöhlerite group includes mineral species that have the general formula $X_8(Si_2O_7)_2W_6$ (Table 1), where $X$ represents the cationic sites typically occupied by $Na^+$, $Ca^{2+}$, $Mn^{2+}$, $Fe^{2+}$, $Ti^{4+}$, $Zr^{4+}$ and $Nb^{5+}$; and where $W$ represents anionic sites with $F^-$, $OH^-$ and $O^{2-}$, which are not bonded to the silicate tetrahedra. The $X$ sites have the same general topology and consequently a specific chemical element will have a different $X$ site preference in different WGM species. The general formula proposed for WGM is similar to that of the rinkite group (seidozerite supergroup)
minerals (Sokolova and Cámara, 2017), however rinkite- and wöhlerite-group minerals have different structures. The crystal structure of WGM is characterised by four-columns-wide octahedral walls, which interconnect through corner sharing and via the disilicate groups to create a framework (Fig. 1). The cationic ordering in the walls and the relative position of the disilicate groups lead to different symmetries (monoclinic and triclinic). The crystal structure of the borate minerals warwickite and yuanfuliite show the same type of framework, with isolated triangular BO₃ groups replacing the disilicate groups (Bigi et al., 1991; Appel et al., 1999).

**Table 1.** List of minerals belonging to the wöhlerite group.

| Mineral and end-member formula | Type of Structure | S.G. | Unit-cell parameters | Ref. |
|--------------------------------|------------------|------|----------------------|------|
| Cuspidine Ca₈(Si₂O₇)₂F₄ | I 1 P2₁/α | 2 | a, Å 10.819 b, Å 10.485 c, Å 7.485 α, ° 90 β, ° 109.55 γ, ° 90 [1] |
| Lävenite Na₂CaMn₂Zr₂(Si₂O₇)₂O₂F₂ | I 1 P2₁/α | 2 | a, Å 10.83 b, Å 9.98 c, Å 7.174 α, ° 90 β, ° 108.1 γ, ° 90 [2] |
| Normandite Na₂CaMn₂Ti₂(Si₂O₇)₂O₂F₂ | I 1 P2₁/α | 2 | a, Å 10.798 b, Å 9.835 c, Å 7.090 α, ° 90 β, ° 108.08 γ, ° 90 [3] |
| Niocalite Ca₅(Si₂O₇)₂O₂F | I 1 P2₁ | 2 | a, Å 10.683 b, Å 10.431 c, Å 7.370 α, ° 110.1 β, ° 90 γ, ° 90 [4] |
| Janhugette Na₅Mn₃Ti₂(Si₂O₇)₂(OH)₂OF | I 1 P2₁/n | 4 | a, Å 10.668 b, Å 9.787 c, Å 13.931 α, ° 107.82 β, ° 90 γ, ° 90 [5] |
| Wöhlerite Na₂Ca₄ZrNb(Si₂O₇)₂O₃F | II 8 P2₁ | 2 | a, Å 10.823 b, Å 10.244 c, Å 7.290 α, ° 90 β, ° 109.00 γ, ° 90 [6] |
| Låvenite Na₂Ca₅Zn₂(Si₂O₇)₂O₂F | II 8 P2₁ | 2 | a, Å 10.1173 b, Å 10.4446 c, Å 7.2555 α, ° 90 β, ° 90.039 γ, ° 90 [7] |
| Burpalite Na₄Ca₂Zr₂(Si₂O₇)₂F₄ | III 6 P2₁/α | 2 | a, Å 10.432 b, Å 10.163 c, Å 7.356 α, ° 90 β, ° 90.96 γ, ° 90 [8] |
| Baghdadite Ca₅Zr₂(Si₂O₇)₂O₄ | III 6 P2₁/α | 2 | a, Å 10.9517 b, Å 10.9251 c, Å 7.3550 α, ° 109.369 β, ° 109.180 γ, ° 83.873 [9, 10] |
| Hiortdahlite Na₅Ca₄(Ca₀.5Zr₀.5)Zr₂(Si₂O₇)₂OF₂ | IV 4 P1 | 2 | a, Å 10.9527 b, Å 10.9289 c, Å 7.3592 α, ° 109.889 β, ° 83.416 [11] |
| Moxuanxueite Na₅Ca₄Zr₂(Si₂O₇)₂OF₃ | IV 4 P1 | 2 | a, Å 10.8459 b, Å 10.2260 c, Å 7.2727 α, ° 90 β, ° 90.332 γ, ° 90 [12] |

S.G. – space group. *see Merlino and Perchiazzi (1988).

References: [1] Saburi et al. (1977); [2] Mellini (1981); [3] Perchiazzi et al. (2000); [4] Mellini (1982); [5] Annehed et al. (1985); [6] Mellini and Merlino (1979); [7] Merlino et al. (1990); [8] Biagioni et al. (2010); [9] Merlino and Perchiazzi (1985); [10] this work; [11] Qu et al. (2020); [12] Chakhmouradian et al. (2008).

**Historical synopsis**

**Cuspidine**

Cuspidine, ideally Ca₈(Si₂O₇)₂F₄ (Z = 2), was described by Scacchi (1876) from Monte Somma, Somma–Vesuvius complex, Italy. Cuspidine occurs in different geological environments such as skarns (Tilley, 1947; Taner et al., 2013), tuff ejecta (Federico and Peccerillo, 2002), pegmatitoid facies of venanzite (Bellezza et al., 2004a), calc-silicate xenoliths (Owens and Kremser, 2010), natrocarbonatite (Mitchell and Belton, 2004) and alkaline rocks (Andreeva et al., 2007).
Cuspidine is monoclinc, $P2_1/a$, with $a = 10.906$, $b = 10.521$, $c = 7.518$ Å and $\beta = 109.90^\circ$. The first structural investigation was provided by Smirnova et al. (1955) who considered the structure as an array of chains of edge-sharing Ca(O,F)$_6$ octahedra running parallel to the c axis, by analogy to the structure of ilvaite, epidote and tilleyite. Subsequent refinement of the cuspidine structure by Saburi et al. (1977) concluded that the coordination environments of the Ca sites are not solely octahedral but vary between six-, seven- and eight-fold. There are four Ca sites in total, with an average bond distance of 2.367, 2.404, 2.428 and 2.449 Å. In each column of the wall, the sites alternate between being small or large. Bellezza et al. (2004a) reported the incorporation of up to 0.22 Zr atoms per formula unit (apfu) and 0.32 Na apfu in cuspidine, following the substitution mechanism $2Ca^{2+} + F^{-} \leftrightarrow Na^{+} + Zr^{4+} + \text{O}^{2-}$. According to their structural model, Zr is incorporated on the small octahedral site lying in the outer columns of the wall ($X_1$), whereas Na is incorporated on the large site ($X_3$) lying in internal columns and connected by edge-sharing to the Zr-bearing octahedra (Fig. 2). Note that Taner et al. (2013) reported cuspidine from the Güneyce Ikizdere Region in Turkey, with an unusually low F content (1.36 apfu), which may correspond to a hydroxide equivalent. Finally, Kızıltan et al. (2018) reported from the Hatrurim complex, Israel, a ‘uranian cuspidine’ containing up to 0.64 U apfu, and only 0.98 F apfu. The oxidation reported from the Hatrurim complex, Israel, a with an unusually low F content (1.36 apfu), which may correspond to the Zr-bearing octahedra (Fig. 2).

Låvenite and normandite

Låvenite, ideally Na$_2$Ca$_2$Mn$_2$Zr$_2$(Si$_2$O$_7$)$_2$O$_2$F$_2$, was described by Brogger (1884) from nepheline syenite pegmatites occurring on the Läven island, Langesundsfjord area, Norway. Låvenite is now reported from many alkaline localities around the world: Igaliku complex, Greenland (Jones and Larsen, 1985; Friis et al., 2010); Los Archipelagos, Boggiani complex, Paraguay (Comin-Chiaramonti et al., 2016); Itatiaia complex, Brazil (Melluso et al., 2017); and Burpala massif, Russia (Vladykin and Sotnikova, 2017).

The crystal structure of låvenite was refined on samples from the Lovozero alkaline massif, Russia (Simonov and Belov, 1960), Langesundsfjord, Norway (Mellino, 1981) and Los Archipelagos, Guine (Biagioni et al., 2012); Cerro Boggiani complex, Paraguay (Comin-Chiaramonti et al., 2016); Itatiaia complex, Brazil (Melluso et al., 2017); and Burpala massif, Russia (Vladykin and Sotnikova, 2017). The crystal structure of låvenite was refined on samples from the Lovozero alkaline massif, Russia (Simonov and Belov, 1960), Langesundsfjord, Norway (Mellino, 1981) and Los Archipelagos, Guine (Biagioni et al., 2012). Låvenite is monoclinic, $P2_1/a$, with $a = 10.83$, $b = 9.98$, $c = 7.17$ Å and $\beta = 108.1^\circ$. These structural data, as well as the chemical data published elsewhere (see the references listed above) indicate clearly the occurrence of cationic substitution on the four $X$ sites (Fig. 2). The larger $X_2$ and $X_4$ sites are dominated by Ca and Na, respectively. Note that $X_2$ usually contains a high amount of Na leading to the mix occupancy close to Ca$_{0.60}$Na$_{0.40}$. The smallest $X_1$ site is dominated by Zr, and the main substitution observed is Zr$^{4+} \leftrightarrow$ Ti$^{4+}$. The last site, $X_3$, has an intermediate size ($\approx 2.23$ Å) and is occupied by a mix of Ca, Fe, Mn and Zr. In låvenite, Mn$^{2+}$ is dominant on $X_3$ though the high Ca contents reported in some samples may indicate that Ca could also be dominant, thus leading to a new end-member composition (Na$_2$Ca$_2$Zr$_2$(Si$_2$O$_7$)$_2$O$_2$F$_2$). The W2 site, bonded to the $X_3$ and $X_4$ sites, is fully occupied by F$^-$, while the W1 site, bonded to the $X_1$, $X_3$ and $X_4$ sites, is populated by O$^2-$ which is partially substituted by F$^-$.

Normandite, ideally Na$_4$Ca$_2$Mn$_2$Ti$_2$(Si$_2$O$_7$)$_2$O$_2$F$_2$, is the titanium analogue of låvenite described initially from the Poudrette quarry, Mont Saint-Hilaire, Quebec, Canada (Chao and Gault, 1997). Note that a mineral with similar physical properties and composition had been reported prior to the normandite description from the Khibiny massif and Lovozero massif, Russia (Vlasov, 1966); Tenerife, Canary Islands (Ferguson, 1978) and Tamazeght, Morocco (Khadem Allah, 1993). Perchiazzi et al. (2000) refined the crystal structure of normandite from the Poudrette quarry and Amdrup Fjord, Greenland. Normandite is monoclinic, $P2_1/a$, with $a = 10.799$, $b = 9.801$, $c = 7.054$ Å and $\beta = 108.08^\circ$. The normandite structure confirmed the structural model and the cation distribution established in låvenite. Perchiazzi et al. (2000) also noted that their samples had an excess of Na and Ca with respect to the expected value of 4 apfu in total, and also had an excess of high-charge cations (Zr and Ti). At the same time, the sum of cations (Mn, Fe and Mg) located on the $X_3$ site is significantly below 2 apfu, suggesting that the excess of Ca and Zr is hosted on the $X_3$ site with an average bond distance of $\approx 2.20$ Å (Fig. 2).

Baghdadite

Baghdadite, Ca$_4$Zr$_2$(Si$_2$O$_7$)$_2$O$_4$, is the only wöhlerite-group mineral without any F or OH groups. First reported from melilitte skarn in contact with diorite, Qala–Dizeh region, Iraq (Hermezi et al., 1986), it has since been described from skarns, calc-silicate marbles and hornfels (Jamviet et al., 1997; Matsubara and Miyawaki, 1999; Shiraga et al., 2001; Galuskin et al., 2007; Galuskina et al., 2010). The main chemical substitution occurring in baghdadite is the homovalent substitution Zr$^{4+} \leftrightarrow$ Ti$^{4+}$. Baghdadite is monoclinic, $P2_1/a$, with $a = 10.432$, $b = 10.163$, $c = 7.356$ Å and $\beta = 90.96^\circ$ (Biagioni et al., 2010). In addition to its chemical composition, the crystal structure of baghdadite is also unique for WGM as it shows the edge-sharing of two ZrO$_6$ octahedra in the internal columns of the wall (Biagioni et al., 2010) (Fig. 2). This structural feature is at odds with the Pauling’s fourth rule, which states that high-valence cations tend to not share polyhedron elements (Pauling, 1929). In all other WGM the high-valence cations (Y$^{3+}$, Ti$^{4+}$, Zr$^{4+}$ and Nb$^{5+}$) do not share any ligands.

Buralpite

Buralpite, Na$_4$Ca$_2$Zr$_2$(Si$_2$O$_7$)$_2$F$_4$, was found for the first time at a fenitised sandstone in the contact zone of the Burpalinskii alkaline massif, North Transbaikal, Russia (Merlino et al., 1990). It is reported in only a few other localities around the world: Umbozero mine, Lovozero massif, Russia; Vesle Aroya, Langesundsfjorden, Norway; and Nanna pegmatite, Igaliku, Greenland (Friis et al., 2010). Chemical data on burpalite are scarce and the published data on the type material indicates a composition close to the end-member formula.

Buralpite is monoclinic, pseudo-orthorhombic, $P2_1/a$, with $a = 10.117$, $b = 10.445$, $c = 7.255$ Å and $\beta = 90.04^\circ$ (Merlino et al., 1990). Site occupancies indicate that the $X_1$ and $X_2$ sites are fully occupied by Zr and Ca, respectively. The larger $X_3$ and $X_4$ sites are mainly populated by Na with minor substitutions of Ca. Bond valence analysis confirms the presence of only F on the $X_2$ site, and the replacement of a small amount of F by O on the $X_1$ site is bonded to the Zr polyhedron (Fig. 2).

Structural investigations performed by Merlino et al. (1990) also indicate that some crystals of burpalite contain domains with a lävenite-type structure. Buralpite- and lävenite-type
Fig. 2. Schematic and idealised representation of the cationic distribution in the walls of cuspidine, baghdadite, burpalite, låvenite, niocalite, janhaugite and wöhlerite (and marianoite, see text). Normandite has the same cationic distribution as låvenite with $X_1 = Ti$. 

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structures are related, as they are two distinct ordered members in a family of order–disorder structures. In addition, a so-called ‘orthorhombic lavenite’ was reported from the Burpalinski massif (Portnov et al., 1966; Portnov and Sidorenko, 1975). This mineral has the same chemical composition as burpalite, but with a B-centred orthorhombic cell, \(a = 21.01, b = 10.05\) and \(c = 7.23\) Å, and is polysynthetically twinned on (100). That unit-cell however, can be transformed to monoclinic, \(a = 11.11, b = 10.05, c = 7.23\) Å and \(\beta = 108.99^\circ\), twinned on (100), analogous to the unit-cell of lavenite. Consequently ‘orthorhombic lavenite’ is a polype of burpalite (burpalite-\(\text{L}_2\)), corresponding to a twinned maximum degree of order (MDO) polytype.

Merlino et al. (1990) suggested that isotopic series may occur between burpalite and baghdadite through the substitution \(\text{Na}^+ + F^- \leftrightarrow \text{Ca}^{2+} + \text{O}^{2-}\). However, a burpalite–baghdadite series is unlikely because in burpalite the \(\text{ZrO}_6\) octahedra are lying in the outer columns of the walls, while in baghdadite they are in the central columns (Fig. 2). Therefore, a transformation of burpalite into baghdadite requires a complete reordering of the cationic octahedral sites.

Wöhlerite

Wöhlerite, ideally \(\text{Na}_2\text{Ca}_5\text{ZrNb}((\text{Si}_2\text{O}_7)_2\text{O}_3\text{F})\), was reported initially by Scheerer (1843) from the syenite pegmatites occurring on Lovøya island, Brevig area, Langesundsfjord, Norway. Wöhlerite is an abundant mineral throughout the Langesundsfjord (Andersen et al., 2016; Larsen, 2010; Sunde et al., 2018), though also from other syenites and carbonatites around the world (e.g. Mariano and Roeder, 1989; Keller and Williams, 1995; Bellezza et al., 2012; Biagioni et al., 2012; Melluso et al., 2017; Guarino et al., 2019).

Wöhlerite is monoclinic, \(P2_1/\alpha\), with \(a = 10.823, b = 10.244, c = 7.290\) Å and \(\beta = 109.00^\circ\) (Mellini and Merlino, 1979). Shibayeva and Belov (1962) and Golsyhev et al. (1973) performed the first structure refinements on wöhlerite and showed the presence of four-columns-large octahedral walls interconnected by corner sharing and \(\text{Si}_2\text{O}_7\) diorthosilicate groups. Mellini and Merlino (1979) provided a structure refinement of wöhlerite from Brevig, Norway, confirming the space group and showing that the structure is based on four independent Si sites, four Ca sites, two Na sites, one \(\text{Zr}\) site and one Nb site (Fig. 2). The bond-variation analysis indicates that only one anionic site is dominated by a monovalent anion. Biagioni et al. (2012) have refined the crystal structure of wöhlerite from Los Archipielagos, Guinea, which contains more Mn and F, and less Nb than the Norwegian material.

Chemical data provided by Mariano and Roeder (1989) on wöhlerite from different localities indicate that the chemical composition of wöhlerite is relatively consistent, and they note that the largest variations are observed for the Nb, Ti and F contents. According to the structural model they establish the coupled substitution of Nb by another chemical element. This is confirmed by subsequent structure refinements in which the Nb site is populated by a significant amount of Mn or Mg (Bellezza et al., 2012; Biagioni et al., 2012), Andersen et al. (2010) and Sunde et al. (2018) showed only minor chemical variations in wöhlerite from different localities in the Larvik plutonic complex in Norway.

Janhaugite

Janhaugite, ideally \(\text{Na}_3\text{Mn}_3\text{Ti}_2((\text{Si}_2\text{O}_7)_2\text{O}_3\text{F})\text{OF}\), is an extremely rare mineral, with Mn content in janhaugite (up to 2.4 apfu) is the highest recorded for any WGM. Electron microprobe analytical (EMPA) data show the presence of roughly one F apfu. The Mn content in janhaugite (up to 2.4 apfu) is the highest recorded for any WGM. Electron microprobe analytical (EMPA) data show the presence of roughly one F apfu. Infrared spectroscopy confirmed the presence of OH groups, and the splitting of the \(\nu\text{O–H}\) stretching frequencies (3550, 3510 cm\(^{-1}\)) may indicate that OH groups are distributed on two different sites. Andersen et al. (2010) and Sunde et al. (2018) showed that the structure coupled with the \(\nu\text{O–H}\) stretching frequencies (3550, 3510 cm\(^{-1}\)) may indicate the presence of OH groups. Nevertheless, the WGM. The Mn content in janhaugite (up to 2.4 apfu) is the highest recorded for any WGM. Electron microprobe analytical (EMPA) data show the presence of roughly one F apfu. Chemical data provided by Mariano and Roeder (1989) on wöhlerite from different localities indicate that the chemical composition of wöhlerite is relatively consistent, and they note that the largest variations are observed for the Nb, Ti and F contents. According to the structural model they establish the coupled substitution of Nb by another chemical element. This is confirmed by subsequent structure refinements in which the Nb site is populated by a significant amount of Mn or Mg (Bellezza et al., 2012; Biagioni et al., 2012), Andersen et al. (2010) and Sunde et al. (2018) showed only minor chemical variations in wöhlerite from different localities in the Larvik plutonic complex in Norway.

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**Wöhlerite**

**Janhaugite**
Hiortdahlite

Hiortdahlite, Na$_2$Ca$_4$(Ca$_{0.5}$Zr$_{0.5}$)Zr(Si$_2$O$_7$)$_2$OF$_3$, one of the oldest WGM known, was described from the Langodden pegmatite (Brøgger, 1890a, 1890b). Hiortdahlite is observed in several localities across the world and has been studied from the Korgeredaba alkaline Massif, Tuva, Russia (Kapustin and Bykova, 1965); Jingera, Australia (Eggleton et al., 1979) and Los Archipelago, Guineá (Biagiioni et al., 2012).

Hiortdahlite is triclinic, $P\bar{1}$, $a = 11.015$, $b = 10.941$, $c = 7.353$ Å, $\alpha = 109.35$, $\beta = 109.88$ and $\gamma = 83.43^\circ$ (Merlino and Perchiazzi, 1985). In addition, in their work on the multi-domains phase called 'guarinite' from Monte Somma, Italy, Bellezza et al. (2012) reported one domain isostructural with hiortdahlite (domain IV). In the first structural investigation of hiortdahlite, Merlino and Perchiazzi (1985) identified three crystallographic sites [M(X7), NaCa (X8), and F3 (W2)] where chemical substitutions occur. The X8 and W2 sites show a mixed occupancy of (Na$_{0.60}$Ca$_{0.40}$) and (F$_{0.60}$O$_{0.40}$), respectively (Fig. 3). The cationic substitutions occurring on the X7 site are complex, with a broad range of cations that may be incorporated (Fe$^{2+}$, Mn$^{2+}$, Ca$^{2+}$ and Zr$^{4+}$).

In their model, Merlino and Perchiazzi (1985), showed that the average charge of the X7 site is +3, and fixed the X7 site population to (Zr$_{0.33}$Ti$_{1.16}$Ca$_{0.10}$Mn$_{0.10}$Fe$_{0.10}$)$_3$F$^3$ according to the chemical data of Brøgger (1890a). Andersen et al. (2010) have reanalysed a crystal fragment of the holotype sample (TYHI0 sample, NRM no. 530976), and gave the formula Na$_{1.55}$Ca$_{4.41}$Mn$_{0.11}$Fe$_{0.10}$REE$_{0.03}$Ce$_{0.01}$Zr$_{1.32}$Ti$_{0.14}$S$_{2.32}$O$_{8.32}$OH$_{0.98}$F$_{2.33}$ (with REE defining lanthanides). New crystallographic and chemical investigations have been performed on samples from the type locality, in order to determine accurately the cationic distribution in the crystal structure. These new data are presented below.

### Table 2. Structural formula for the minerals of the wöhlerite group.

| One topological wall | (X1)$_2$ | (X2)$_2$ | (X3)$_2$ | (X4)$_2$ | (Si$_2$O$_7$)$_2$ | (W1)$_2$ | (W2)$_2$ |
|---------------------|----------|----------|----------|----------|----------------|----------|----------|
| Cuspidine           | Ca       | Ca       | Ca       | Ca       | (Si$_2$O$_7$)$_2$ | F         | F         |
| Baghdadite          | Ca       | Zr       | Ca       |          | (Si$_2$O$_7$)$_2$ | O         | O         |
| Burpalite           | Zr       | Ca       | Na       | Na       | (Si$_2$O$_7$)$_2$ | F         | F         |
| Lävenite            | Zr       | Ca       | Mn       | Na       | (Si$_2$O$_7$)$_2$ | O         | F         |
| Normandite          | Ti       | Ca       | Mn       | Na       | (Si$_2$O$_7$)$_2$ | O         | F         |
| Niocalite           | Ca       | Ca       | Ca       | Na       | Ca             | Ca        | Ca        |
|                     |          |          |          |          | (Si$_2$O$_7$)$_2$ |          | (Si$_2$O$_7$)$_2$ |
|                     |          |          |          |          | (O$_{2.5}$F$_{2.5}$)OH | (O$_{2.5}$F$_{2.5}$)OH | |
| Janhaugite          | Ti       | Na       | Ti       | Mn       | Na             | Mn        | (Si$_2$O$_7$)$_2$ |
|                     |          |          |          |          |          |           | (O$_{2.5}$F$_{2.5}$)OH | (O$_{2.5}$F$_{2.5}$)OH |
| Wöhlerite           | Zr       | Ca       | Na       | Na       | Nb             | Ca        | Ca        |
|                     |          |          |          |          | (Si$_2$O$_7$)$_2$ | O         | (Si$_2$O$_7$)$_2$ |
|                     |          |          |          |          |          |           | (O$_{2.5}$F$_{2.5}$)OH | (O$_{2.5}$F$_{2.5}$)OH |
| Marianosite         | Zr       | Ca       | Na       | Na       | Nb             | Ca        | Ca        |
|                     |          |          |          |          | (Si$_2$O$_7$)$_2$ | O         | (Si$_2$O$_7$)$_2$ |
|                     |          |          |          |          |          |           | (O$_{2.5}$F$_{2.5}$)OH | (O$_{2.5}$F$_{2.5}$)OH |

| Two topological walls* | X1 | X2 | X3 | X4 | X5 | X6 | X7 | X8 | (Si$_2$O$_7$)$_2$ | W1 | W2 | W3 | W4 |
|------------------------|----|----|----|----|----|----|----|----|----------------|----|----|----|----|
| Hiortdahlite           | Zr | Ca | Na | Ca | Ca | Ca | Ca | Ca | Ca             | Ca | Ca | Ca | Ca |
|                       | Zr | Ca | Na | Ca | Ca | Ca | Ca | Ca | Ca             | Ca | Ca | Ca | Ca |
|                       | Zr | Ca | Na | Ca | Ca | Ca | Ca | Ca | Ca             | Ca | Ca | Ca | Ca |

*X1–X4 and X5–X8 sites are part of the first and second wall, respectively. See references from Table 1 for the other minerals.

Moxuanxueite

Moxuanxueite, ideally NaCa$_4$Zr(Si$_2$O$_7$)$_2$OF$_3$, was described recently from the alkaline syenites of the Gejiu intrusion in Honghe Hani and Yi, Autonomous Prefecture, Yunnan Province, China (Qu et al., 2020). The mineral is triclinic, $P\bar{1}$, $a = 10.953$, $b = 10.929$, $c = 7.359$ Å, $\alpha = 109.41$, $\beta = 109.89$ and $\gamma = 83.42^\circ$. Moxuanxueite is isostructural with hiortdahlite, with the X7 and X8 sites fully occupied by Ca (Table 2).

### Chemical compositions of WGM and related species

#### Chemical classification of WGM

The members of the wöhlerite group show a range of compositions with the main cations being Na, Ca, Fe, Mn, Zr, Ti and...
Nb while the main anions are O, F and OH\(^-\). These elements are also the main components of other disilicates with similar optical properties common to alkaline rocks, e.g. rinkite-group minerals of the seidozerite supergroup (Sokolova and Câmara, 2017; Pautov et al. 2019). Most petrological studies of alkaline rocks do not utilise techniques other than chemical data to classify minerals. Therefore, we have explored the feasibility of (1) distinguishing wöhlerite-group minerals from related minerals and (2) classifying WGM down to species level solely based on EMPA data. We have used a total of 908 analyses of WGM and related minerals from our own work and the literature: Aarden and Gittins (1974); Ferguson (1978); Eggleton et al. (1979); Raade and Mlake (1983); Hermezi et al. (1986); Marino and Roeder (1989); Meredith et al. (1990); Keller and Williams (1995); Sharygin et al. (1996a,b); Chao and Gault (1997); Jamtveit et al. (1997); Stoppa et al. (1997); Atencio et al. (1999); Perchiazzi et al. (2000); Roda-Robles et al. (2001); Shiraga et al. (2001); Federico and Peccerillo (2002); Christiansen et al. (2003); Bella et al. (2004a, 2004b); Mitchell and Belton (2004); Casillas et al. (2008); Chakhmouradian et al. (2008); Andersen et al. (2010); Friis et al. (2010); Owens and Kremser (2010); Bella et al. (2012); Biagiioni et al. (2012); Chakrabarty et al. (2013); Chen et al. (2013); Melluso et al. (2014); Ronso et al. (2014); Comin-Chiaramonti et al. (2016); Melluso et al. (2017); Chakrabarty et al. (2018); Sunde et al. (2018) and Guarrino et al. (2019).

Before attempting a classification or discrimination of species based purely on chemistry, all data were recalculated on the basis of 18 anions. We have maintained the identification of each analytical point as given in the respective papers, i.e. we have not changed mineral identifications. The WGM typically have no substitution on the Si sites and the X sites are filled, i.e. there are no, or only limited, vacancies in fresh material. Therefore, only data where 3.9 < Si apfu < 4.1 and 7.8 < ΣX apfu < 8.2 should be treated. We allow for some variation from ideal stoichiometry due to the challenges of analyses some of these minerals. Of the 908 analytical points 258 did not fulfill these criteria and therefore the following is based on the remaining 650 analyses.

Keller and Williams (1995) used three ternary plots to classify WGMs, and we present our data in two of the same diagrams (Fig. 4). Contrary to the paper by Keller and Williams (1995) our data show significant overlaps between species. For example, hainite-(Y), kochite and rosenbuschite overlap in all plots and partly overlap with lävenite. Conversely, wöhlerite forms a distinct group in Fig. 4a,c,e,f. However, Fig. 4a shows that (i) ianjahuate (WGM) overlaps with rinkite-group minerals grennarite and seidozerite and (ii) lävenite (WGM) overlaps with rosenbuschite (rinkite group, seidozerite supergroup). Figure 4b,e show (i) a strong overlap between four minerals: wöhlerite-group minerals wöhlerite and hiortdahlite and rinkite-group minerals hainite-(Y), rinkite-(Ce) and rinkite-(Y).

Chakhmouradian et al. (2008) plotted data for some WGM based only on divalent cations occupying the true octahedral sites. This method suffers from the same issues as Keller and Williams (1995) with large overlaps between different species, especially if data for hiortdahlite is included. Melluso et al. (2014) suggested other graphical methods to classify WGM and related species, but also concluded that the high degree of overlap between species and endmembers of species does not make these plots suitable for species determination.

The previously proposed methods for classification do not enable a satisfactory determination at species level from only chemical data. At first glance, the plots by Keller and Williams (1995) do seem to create some distinct groups and it may be possible to separate some species based on them. However, the data presented by Keller and Williams (1995) has in a sense already been filtered as they only presented WGM data. Therefore, these plots may help identify some WGM, but only when it is already known that the chemical data is actually from a WGM. Conversely, they fail when the mineral has not been identified, at least, to a group level. On the basis of the available chemical data it is not possible to make a graphic interpretation to identify WGM or distinguish them from related minerals solely based on chemical data. However, the plots may work if additional methods are applied, for example X-ray diffraction (XRD), to determine if the mineral is a member of the wöhlerite group or another chemically related group e.g. the seidozerite supergroup.

As the graphical methods do not enable distinction of WGM form related mineral groups, let alone identification at species level we have used the same chemical data to establish a workflow for treating chemical data. It must be stressed that the flow below requires the sequential removal of data so that the next step in the flow are criteria to be applied on the remaining data after samples have been removed by the previous step (Fig. 5).

1. Remove poor quality data.
2. Remove data with REE+Y > 0.5 apfu, which are REE-bearing rinkite-group minerals.
3. Remove data with Ti+Zr+Hf > 2.5 apfu. This removes seidozerite and grennartite.
4. Remove data with Fe+Mn > 3 apfu, which will be janhaugite.
5. Remove data where Ca > 5.9 apfu. This step removes cuspidine–baghdadite–niocalite.
6. Remove samples with Ca < 2.9 apfu. This step removes lävenite, normandite and burpalite, but also some kochite.
7. Remove data with Nb+Ta > 0.5 apfu, which corresponds to wöhlerite.
8. Remove data with Ti/(Zr+Hf) < 0.2, which will separate hiortdahlite, moxuanxueite and one data point given as rosenbuschite in the literature.

After these steps a total of 122 data points remain corresponding to götzenite, hainite-(Y), kochite and rosenbuschite. In addition, one data point given as hiortdahlite and the Zr–Ti-cuspidine of Sharygin et al. (1996a) remains. The method above provides a good separation of WGM from seidozerite-group minerals.

**Solid solution in WGM**

The literature often refers to the WGM having a flexible structure resulting in large degree of solid solution (e.g. Perchiazzi et al., 2000; Mitchell and Belton, 2004; Chakhmouradian et al., 2008). Regardless of the diverse composition of the WGM, solid solutions are not as extensive as the chemical data may suggest. There seems to be a high degree of solid solution between cuspidine and niocalite, as well as between cuspidine and baghdadite, however, the solid solution between niocalite and baghdadite is limited (Fig. 4f). In baghdadite, the two Zr sites are edge sharing, resulting in a highly distorted site. If niobium completely replaces Zr, two Nb polyhedra would be edge sharing, which is highly unlikely to happen. Furthermore, a complete replacement of Zr by Nb is not possible as the additional two charges cannot be balanced because all anions in baghdadite are already oxygen. A full solid solution would be the coupled substitution 2Zr\(^{4+} +\)
Fig. 4. (a–c) Ternary plots of 650 chemical analyses of WGM and related species from our data and the references listed in the text and (d–f) only WGM. Abbreviations: Baghdadite (Bgd); burpalite (Brp); cuspidine (Csp); götzenite (Göz); grenmarite (Grn); hainite-(Y) (Hai); hiortdahlite (Hio); janhaugite (Jhg); kochite (Koh); låvenite (Låv); mxouanxueite (Mox); niocalite (Nio); normandite (Nmd); nacareniobsite-(Ce) (Nns); rosenbuschite (Rbs); rinkite-(Ce) (Rin); seidozerite (Sdz) and wöhlerite (Wöh). In the figure the abbreviations are in italic for WGM.
O$_2^-$ ↔ Ca$_2^+$ + Nb$_5^+$ + F$^-$, but this is not a simple substitution as baghdadite and niocalite are not isostructural, and it would lead to a major structural change (Fig. 2). However, it is likely that there is a limited solid solution between the two species as indicated by Casillas et al. (2008). Keller and Williams (1995) suggested a degree of solid solution between niocalite and wöhlerite, but most of these data have more than 8.2 apfu X site cations and were removed by the above data processing. Wöhlerite and niocalite are not isostructural, therefore solid solution between the minerals not only requires substitution between Ca, Na, Zr and F, but also a change of the position of Nb in the structure between X5 and X6 (Tables 1, 2; Fig. 3).

Sharygin et al. (1996a) investigated a series of cuspidine and götzenite minerals from Pian di Celle in Italy and suggested a partial solid solution between cuspidine and an end-member with the composition NaCa$_4$Zr(Si$_2$O$_7$)$_2$OF$_3$. From powder XRD and Raman spectroscopy Sharygin et al. (1996a) showed this phase to be structurally more closely related to hiortdahlite than cuspidine, and in fact this composition corresponds to the recently approved mineral moxuanxuieite (Qu et al., 2020).

Extended solid solution has been documented between lăvenite and normandite (Perchiazi et al., 2000) as Zr in lăvenite and Ti in normandite both occupy the X1 site and hence can replace each other with no additional structural modifications. Similarly, the Mn/Fe ratio differs in lăvenite and as both these cations occupy the X3 site, it is likely that Fe-equivalent species may be found of both minerals.

In summary, solid solution in the WGM is controlled by the crystal structure and occurs where no major structural modifications are required, e.g. between lăvenite and normandite, or

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**Fig. 5.** Schematic workflow for discriminating chemical data into different disilicate groups.
between cuspidine, baghdadite and niocalite. Therefore, it is not uncommon to find several different WGM in the same rocks as small chemical changes favour the formation of multiple species rather than creating solid-solution series.

Nomenclature for wöhlerite-group minerals

Classification

The general formula for WGM can be expressed as $X_8(SiO_2)_{12}W_4$, without any further distinction among the X and W sites. As shown in Fig. 2, a specific chemical element will have a different preferential X site in different WGM. For instance, high-field-strength elements (HFSE) such as Ti, Zr and Nb, are hosted on crystallographic sites located in different columns of the walls and on different cationic X sites. Consequently, there is no crystal-chemical feature to assign elements to specific X sites. As a rule, the topological representation of the cationic walls has been made by drawing the projection of the wall along the crystallographic c axis. The labelling of the sites always starts with X1, which is the smallest site belonging to outer columns of the wall. The site sitting in the outer column and connected to X1 though edge-sharing is labelled X2. Afterwards the sites are labelled in succession, going from the outer to the inner columns of the wall, and according to the symmetry of the mineral. The crystal structure of hiortdahlite is characterised by two cationic walls, where wall I contains the HFSE-dominated site located in the outer column (Fig. 3). The anionic W sites correspond to the ligands that are not bonded to the disilicate groups. W1 and W2 are the anionic sites bonded to X1 and X2, respectively. Depending of the symmetry of the mineral, W3 and W4 are either belonging to the same outer column as W1 and W2, or to the second outer column of the wall. The W anionic sites occurring between the two inner columns are the symmetrical equivalents of the W sites defined previously.

The classification of WGM is based on the occupancy of the structural X and W sites, and the different combination of chemical elements on these sites are regarded as different mineral species. Considering only chemical data is in some cases insufficient to correctly identify WGM at a species level. Consequently, a structural refinement is needed for a complete characterisation. The sequential application of the dominant-valency and the structural walls is not a valid criterion to define a new species, as it would lead to a proliferation of the number of species. A mineral phase showing the same chemical composition as previously described species but with a different cationic ordering must be considered as an analogue to that species. Polytypism is likely to occur among WGM (Merlino and Perchiazzi, 1988), and polytypes are not considered as different mineral species (Nickel and Grice, 1998). The polytypes must be described according to the nomenclature proposed in the IMA–CNMNC guidelines (Guinier et al., 1984; Nickel, 1993; Nickel and Grice, 1998).

Name, prefixes and suffixes

All the wöhlerite-group minerals have a distinct name, with no prefix or suffix. Due to the relatively large number of structural sites that are able to host the same cation (e.g. Ca$^{2+}$), we strongly discourage the use of a compositional suffix (e.g. calcio-, ferro-), as this is unclear to which structural site these prefixes will be related. For the same reason, we also discourage the use of prefix referring to the composition of anionic W sites. In the case of a species characterised by one crystallographic site dominated by Y or REE, we recommend using a new rootname as well as the Levinson suffix (Bayliss and Levinson, 1988).

Hiortdahlite end-member formula

In order to define the correct end-member formula of hiortdahlite new chemical data was collected and a crystal structure refinement was performed on a sample from the type locality, Langodden, Langesundsfjord, Norway (samples located in the NHM Oslo collections, catalogue number KNR 24099). The chemical data were acquired using the Cameca SX100 electron microprobe housed at the Department of Geosciences, University of Oslo. The instrument was operated with a beam current of 15 nA and an acceleration voltage of 15kV, creating a 10 μm spot. The following natural and synthetic standards were used: albite (Na), zircon (Zr), wollastonite (Ca and Si), pyrophanite (Ti and Mn), REE orthophosphate (Y, La, Ce and Nd; Jarosewich and Botnearer, 1991), MgO (Mg), Fe metal (Fe), Nb metal (Nb) and fluorite (F). The intensity data were corrected for inter-element overlaps and matrix effects using the PAP routine (Pouchou and Pichoir 1984). The chemical data are compared with those published by Andersen et al. (2010) on type-locality material (Table 3).

Single-crystal X-ray data were collected at room temperature with monochromated MoKα radiation ($\lambda = 0.71703 \text{ Å} - 50 \text{ kV}$ and 1 mA) on a Rigaku Synergy-S diffractometer equipped with a HyPix-6000He detector housed at NHM Oslo. The instrument has Kappa geometry and both data collection and

| Constituent | Average (wt.%) | S.D. (2σ) | Range | afu | Average (wt.%) | afu |
|-------------|----------------|-----------|-------|-----|----------------|-----|
| Na$_2$O     | 6.62           | 0.30      | 6.39-6.87 | 1.66 | 6.84           | 1.74 |
| CaO         | 32.90          | 0.69      | 32.32-33.62 | 4.55 | 32.77          | 4.60 |
| Y$_2$O$_3$  | 0.56           | 0.08      | 0.48-0.63 | 0.04 | 0.48           | 0.03 |
| La$_2$O$_3$ | 0.09           | 0.05      | 0.03-0.16 | 0.01 | 0.08           | 0.01 |
| Ce$_2$O$_3$ | 0.26           | 0.05      | 0.21-0.31 | 0.01 | 0.30           | 0.01 |
| Nd$_2$O$_3$ | 0.12           | 0.14      | 0.01-0.24 | 0.01 | 0.08           | 0.01 |
| MgO         | 0.04           | 0.02      | 0.02-0.06 | 0.01 | -              | -   |
| MnO         | 0.92           | 0.12      | 0.80-1.06 | 0.10 | 0.92           | 0.10 |
| FeO         | 1.17           | 0.18      | 1.00-1.40 | 0.13 | 0.87           | 0.10 |
| TiO$_2$     | 0.98           | 0.12      | 0.83-1.13 | 0.10 | 0.87           | 0.09 |
| ZrO$_2$     | 17.12          | 0.81      | 16.32-17.81 | 1.08 | 18.21          | 1.16 |
| Nb$_2$O$_5$ | 1.90           | 0.42      | 1.43-2.28 | 0.11 | 1.81           | 0.11 |
| SiO$_2$     | 30.76          | 0.49      | 30.23-31.45 | 3.97 | 30.52          | 4.00 |
| F            | 8.41           | 0.56      | 3.33-3.77 | 3.44 | 8.21           | 3.40 |
| O = F        | 3.54           |           |         | 3.46 |               |     |
| Total       | 98.31          |           |         | 98.96 |               |     |

S.D. = standard deviation
subsequent data reduction, together with face-based absorption corrections were carried out using the Rigaku CrystalisPro software. The details of the data collection and refinement are provided in Table 3. The initial structure solution in space group P1 was determined by the charge flipping method using the Superflip algorithm (Palatinus and Chapuis, 2007), and the structural model was subsequently refined on the basis of $F^2$ with the Jana2006 software (Petřiček et al., 2014). All atoms were refined with anisotropic thermal parameters. The details of the refinement are provided in Table 4 and the atoms coordinates, anisotropic thermal parameters and detailed bond distances are provided in the Supplementary tables S1, S2 and S3. Free refinement of the site-scattering factors showed that the occupancy on the $X$7 site is in the central columns. Note that both walls in the second site is in the outer columns, while in the second $X$5Ca$\gamma$ site is in the central columns.

The empirical formula is recalculated on the basis of $\Sigma (O+F) = 18$ apfu. The site-scattering factors and the established cationic distribution in the crystal structure are provided in Table 5 and the bond-valence table is provided in Table 6. The crystallographic information file has been deposited with the Principal Editor of Mineralogical Magazine and is available as Supplementary material.

As shown by Merlino and Perchiazzi (1985), the main chemical substitutions occur on the $X$7 and $X$8 sites and the bond-valence analysis indicate a mixed $O^2$ + $F^\prime$ occupancy on the anionic W1 site (Table 7). The refinement of the site-scattering for the Ca1 ($X$2) and Ca3 ($X$5) sites shows a deficit and an excess of electronic density, respectively. Therefore, considering the average bond distances around these sites, Ca + Na and Ca + Y + REE have also been refined on these sites, respectively (Table 5). All Nb, Ti and Hf has been attributed to the $Zr$ site (X1), which was then filled up with Zr. The remaining Zr was attributed to the $X$7 sites, along with all the Mn and Fe content. The $X$7 site was then filled with Ca. Refinement of the site scattering in the $X$8 position indicates an excess of electron density and therefore Ca was assumed to replace Na. The cationic distribution proposed is in excellent agreement with the different structural parameters and the bond-valence analysis (Tables 5, 6). Bond-valence sums show that the W2, W3 and W4 sites are populated by F and that a substitution $O^2$ ↔ $F^\prime$ is occurring on the W1 site (Table 6). Taking into account previous work and the new data presented here, the end-member formula of hiortdahlite is Na$_2$Ca$_4$(Ca$_{0.5}$Zr$_{0.5}$)Zr(Si$_2$O$_7$)$_2$OF$_3$, with a constrained mixed occupancy of Ca$_{0.5}$Zr$_{0.5}$ (M$_{2+}$), in order to obtain a charge-balanced formula.

The refinement provided in this work is slightly different than the one proposed by Biagioni et al. (2012) on a mineral phase structurally related to hiortdahlite (Table 7). The incorporation of Ti$^{4+}$ on the larger $X$7 site (<$X$7–$O$> = 2.233Å) instead of the smaller $X$1 site (<$X$1–$O$> = 2.082Å) is unlikely considering the ideal bond distance for octahedrally coordinated Ti (2.005Å, Shannon (1976); Table 5, S3). The partitioning of Y and REE between the $X$5 and Na sites ($X$3) is not clear, however, we have not detected a refined site-scattering factor higher than 11 eπ pm for the $X$3 site in our investigations.

Hiortdahlite and moxuanxueite are the only approved WGM with a structure containing two topologically independent octahedral walls. The ideal compositions of the walls are given by wall-I$^\prime$:X$_1$Zr$^2$X$_2$Ca$^2$ X$_3$Na$^2$Ca$^2$ and wall-II: X$_5$Ca$^2$X$_6$X$_7$(Zr$_{0.5}$Ca$_{0.5}$)$^2$X$_8$Na. The chemical compositions of the walls are similar, although in the first wall the Zr site is in the outer columns, while in the second wall the X7 site is in the central columns. Note that both walls in hiortdahlite are topologically and chemically unique among the WGM (Table 2).

### Table 4. Data collection and structure refinement details for hiortdahlite from Langodden, Langesundsfjord, Norway.

| Parameter | Value |
|-----------|-------|
| Temperature (K) | 293 |
| Space group | P1 |
| Lattice parameters | |
| $a$ (Å) | 10.9517(1) |
| $b$ (Å) | 10.9251(1) |
| $c$ (Å) | 7.3555(1) |
| $\alpha$ (°) | 109.369(1) |
| $\beta$ (°) | 109.180(1) |
| $\gamma$ (°) | 83.873(1) |
| $V$ ($Å^3$) | 784.17(2) |
| $Z$ | 2 |

### Table 5. Site population assignment and structural parameters for the crystal structure of hiortdahlite.

| Site | RSS | Site-population (apfu) | CSS | ABL | CBL |
|------|-----|------------------------|-----|-----|-----|
| X1   | 38.4 | Zr$_{0.79}$Nb$_{0.11}$ | 38.3 | 2.084 | 2.100 |
| X2   | 18.8 | Ca$_{0.80}$Nb$_{0.15}$ | 18.7 | 2.515 | 2.549 |
| X3   | 11.0 | Na$_{0.00}$ | 11.0 | 2.444 | 2.420 |
| X4   | 20.0 | Ca$_{0.00}$ | 20.0 | 2.407 | 2.460 |
| X5   | 21.0 | Ca$_{0.30}$Zr$_{0.24}$ | 21.9 | 2.340 | 2.396 |
| X6   | 20.0 | Ca$_{0.00}$ | 20.0 | 2.378 | 2.400 |
| X7   | 26.6 | Ca$_{0.80}$Zr$_{0.25}$ | 27.1 | 2.231 | 2.233 |
| X8   | 14.4 | Na$_{0.00}$Ca$_{0.40}$ | 14.6 | 2.567 | 2.576 |

RSS: Refined site scattering (epfu); CSS: calculated site scattering (epfu); ABL: average observed bond-distances (Å); CBL: calculated averaged bond-distances (Å), using the ionic radii of Shannon (1976).

### Discreditation of marianoite

Marianoite was discovered from the silicocarbonate Prairie Lake complex, Ontario, Canada (Chakhmouradian et al., 2008), and was considered the Nb-analogue of wöhlerite. Its simplified formula is Na$_2$Ca$_4$(Nb$_{0.33}$Zr$_{0.25}$)(Si$_{2.00}$O$_{7.33}$)(O,F)$_3$. Marianoite was described as monoclinic, $P2_1$, with $a = 10.846$, $b = 10.226$, $c = 7.273$ Å and $\beta = 109.33^\circ$. The highest Nb content reported by Chakhmouradian et al. (2008) for marianoite is 1.019 apfu, which is roughly 0.3 apfu more than in wöhlerite from the Langesundsfjord (Sunde...
et al., 2018). The approval of marianoite as a valid mineral species was based on the assumption that both Zr and Nb are disordered on the two smallest octahedral sites [average bond lengths: 2.031 (X6) and 2.080 Å (X1)]. As a result of the similar X-ray and neutron scattering characteristics of Zr and Nb, it is not possible to solve the ordering issue between these two chemical elements by using standard diffraction methods. Following the description of marianoite, Merlino and Mellini (2009) published a discussion arguing that in wöhlerite and marianoite there is an ordering of Zr and Nb, with Nb preferentially occupying the smallest site (X6) and Zr the second smallest octahedra (X1) of the structure. The same authors have proposed to solve this question through anomalous scattering using synchrotron radiation sources that will allow Zr and Nb to be distinguished. Bellezza et al. (2012) and Biagioni et al. (2012) have used an ordered approach in their refinements. Readers are referred to Merlino and Mellini (2009) and Chakhmouradian and Mitchell (2009) for more information on that discussion.

Following the classification system proposed herein for WGM, wöhlerite and marianoite are equivalent. If one considers a complete cationic ordering between Zr4+ and Nb5+, the resulting end-member formula for both wöhlerite and marianoite is Na2Ca4(X1(Zr)X6(Nb)(Si2O7)2O3F. The maximum Nb content reported for marianoite is 1.02 apfu (associated with 0.85 Zr pfu) (Chakhmouradian et al., 2008), which is not enough to achieve Nb > Zr on both X1 and X6 sites and then define a

| Site       | X1   | X2   | X3   | X4   | X5   | X6   | X7   | X8   | Si1  | Si2  | Si3  | Si4  | Σ     |
|------------|------|------|------|------|------|------|------|------|------|------|------|------|-------|
| Hiortdahlite | 11.0149(9) | 10.9517(1) | 10.991(7) | 10.970(2) | 3.85 | 1.71 | 0.94 | 2.00 | 2.12 | 1.92 | 2.40 | 1.21 | 4.03 | 4.04 | 4.11 |
| Langodden [2]** | 10.9499(9) | 10.9251(1) | 10.930(3) | 10.9431(2) | 1.07 |
| Los Archipelago [3] | 7.3534(3) | 7.3555(1) | 7.3660(2) | 7.365(3) | 1.00 |
| Monte Somma [4] | 83.4344(4) | 83.873(3) | 83.95(3) | 83.39(1) | 1.00 |

*Not hiortdahlite s.s. **Langodden (TL) - Structural formula: Na2(4)Ca4.66Fe0.13Mn0.10Y0.04Ce0.03Zr1.08Nb0.11Ti0.10(Si2O7)2(O0.6F0.4)F3, Chemical formula: Na1.66Ca4.55Fe0.13Mn0.10Y0.04Ce0.03Zr1.08Nb0.11Ti0.10(Si1.98O7)2(O0.57F0.43)F3 (Table 3) [1] Merlino and Perchiazzi (1983); [2] this work; [3] Biagioni et al. (2012); [4] Domain IV, Bellezza et al. (2012).
Nb-dominant end-member. In the second case, if one considers that Zr$^{4+}$ and Nb$^{5+}$ are disordered, the X1 + X6 sites would have a total charge of +9, to keep the charge balance of the formula Na$_2$Ca$_4$(X1)(X6)(Si$_2$O$_7$)$_2$O$_3$F. The possible charge arrangements of the dominant cations R occupying the X1 and X6 sites are (i) X1(R$^{4+}$) + X6(R$^{5+}$) and (ii) X1(R$^{4.5+}$) + X6(R$^{4.5+}$). The charge arrangement (ii) is not valid, because it implies a double occupancy (Zr$^{4+}$Nb$^{5+}$) on two sites. The charge arrangement (i) leads to the end-member formula Na$_2$Ca$_4$(X1)(R$^{4+}$)X6(R$^{5+}$)(Si$_2$O$_7$)$_2$O$_3$F, and to the atomic arrangement Na$_2$Ca$_4$(X1)(Zr$^{4+}$)X6(Nb$^{5+}$)(Si$_2$O$_7$)$_2$O$_3$F. The atomic arrangement and the end-member formula are identical to those of wöhlerite. Consequently, marianoite must be considered equivalent to wöhlerite and is discredited.

**Comment on the phase ‘hiortdahlite II’**

‘Hiortdahlite II’ is not an approved mineral species, although Merlino and Perchiazzi (1987) stated that the name ‘hiortdahlite II’ was approved by the IMA Commission on New Minerals and Mineral Names (merged with CNMNC in 2006), a subsequent new mineral proposal was never submitted. Hiortdahlite II was described by Aarden and Gittins (1974) in samples from the Kipawa River, Kipawa alkaline complex, Quebec, Canada. Roda Robles et al. (2001) reported hiortdahlite II from the Ililmaussaq alkaline complex, Greenland, Tamazeght complex, Morocco, and Iles de Los, Guinea, based on chemical analyses and powder X-ray diffraction. We have analysed material from the same locality in the Ililmaussaq complex, and all of the supposed hiortdahlite II crystals have a unit-cell setting and a crystal structure identical to those of hiortdahlite. Therefore, it may be questionable that hiortdahlite II exists at these localities. This further emphasises the need for full crystal structure refinement to correctly identify WGM at a species level.

Chemical analysis on the ‘type’ material gave the formula (Na$_{1.70}$Ca$_{4.02}$Mn$_{0.04}$Fe$_{0.02}$Mg$_{0.02}$Al$_{0.02}$Y$_{0.24}$REE$_{0.08}$Zr$_{1.16}$Nb$_{0.04}$Ti$_{0.02}$)$_{16}$(Si$_{2.05}$O$_7$)$_2$O$_0.82$OH$_{0.36}$F$_{2.68}$, and therefore hiortdahlite II was interpreted as a cationic-deficient analogue of hiortdahlite (Aarden and Gittins, 1974). However, recent chemical analyses performed on the type material of hiortdahlite (Andersen et al., 2010) indicate roughly the same amount of Zr per unit formula than in the material described by Aarden and Gittins (1974). Note that hiortdahlite II contains up 0.24 Y apfu, while in hiortdahlite from Langodden, the Y content is below 0.05 apfu. The total REE content is also slightly larger in hiortdahlite II than in hiortdahlite.

Hiortdahlite II is reported as triclinic, PI, with a = 10.95, b = 10.31, c = 7.29 Å, α = 90.19, β = 109.02 and γ = 90.05° (Aarden and Gittins, 1974). The crystal structure refinement was performed on samples from Kipawa, and gave a structural model based on two independent topological octahedral walls (Merlino and Perchiazzi, 1987). The wall-I in hiortdahlite II has the same chemical composition as the wall-I in hiortdahlite, though the cationic distribution is not strictly equivalent (Fig. 3). The main difference between species is observed in the topology of wall-II, with a composition of [X$_5$Ca$_6$X$_6$Ca$^{4+}$Zr$_{0.5}$Ca$_{0.5}$]X$_8$Na] and [X$_5$Ca$_6$X$_6$Ca$^{4+}$Zr$_{0.5}$Ca$_{0.5}$] in hiortdahlite and hiortdahlite II, respectively. However, the difference between the crystal chemical formula provided by Merlino and Perchiazzi (1987) and the chemical data reported by Aarden and Gittins (1974) is significant, for instance 1.76 Y apfu is reported in the structure while the chemical data indicate 0.32 Y + REE apfu.

Consequently, the presence of a Y-dominant site in the structure of hiortdahlite II is questionable, and new investigations must be performed on hiortdahlite II material to explore if it is a poly-type of hiortdahlite or a distinct and valid mineral species. A new mineral proposal would still be required to be submitted to the IMA–CNMNC.

**Conclusions**

The general formula of the wöhlerite-group minerals is given by $X_6(Si,O,F)_2W_{α}$, where W denotes the cations occurring in the polyhedra building the four-column wall, and where $W$ denotes the anionic sites that are not bonded to the disilicate groups. The crystal structure of WGM is characterised by ‘octahedral walls’ made of four columns of edge-sharing X sites. The symmetry of the different species can vary from monoclinic to triclinic, according to the cationic ordering on the X sites and the relative position of the disilicate groups. Distinction between the mineral species is made based on the dominant elements at the X and W sites, and different combinations of X and W constituents should be regarded as separate mineral species.

In addition to the classification scheme, the following changes have been approved by the IMA–CNMNC: (i) the end-member formula of hiortdahlite has changed to Na$_2$Ca$_4$(Ca$_{0.5}$Zr$_{0.5}$)Zr (Si$_2$O$_7$)$_2$O$_3$F$_{2.5}$, with a valency-imposed double-site occupancy of (Zr$^{4+}$Zr$^{5+}$) on the X7 site, and (ii) marianoite is discredited, as it is structurally and chemically equivalent to wöhlerite. The chemical variation in WGM results in the formation of individual species rather than solid-solution series. The reason being that despite similar compositions between many of the members they are not isostructural, therefore, heterovalent substitutions typically require a complete reordering of the structure. Such reorders appear energetically unfavourable compared to the formation of another species, commonly resulting in rocks containing several WGM or even seidozerite supergroup minerals. Furthermore, the co-existence of different WGM and seidozerite-supergroup minerals in the same rock makes it a challenge for petrologists to identify minerals on a species level. We proposed a discrimination flow-chart for separating various WGM from chemically related species.

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