

in three of the equatorial CO ligands adopting bridging conformations.

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Crystal structures of $\text{Ln}_5\text{O}_4[\text{AsO}_3]_2\text{Cl}$ ($\text{Ln} = \text{Pr}, \text{Sm}$) and $\text{Ce}_3\text{O}[\text{AsO}_3]_2\text{Cl}$. Hamdi Ben Yahia^a, Ute Ch. Rodewald^a, Rainer Pöttgen^a. ^aInstitut für Anorganische und Analytische Chemie, Universität Münster, Corrensstrasse 30, D-48149 Münster, Germany.

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The title compounds with structures isotopic to the $\text{Nd}_5\text{O}_4[\text{AsO}_3]_2\text{Cl}$ [1] and $\text{Gd}_3\text{O}[\text{AsO}_3]_2\text{Cl}$ [2] types have been synthesised. The crystal structures of $\text{Pr}_5\text{O}_4[\text{AsO}_3]_2\text{Cl}$, $\text{Sm}_5\text{O}_4[\text{AsO}_3]_2\text{Cl}$ and $\text{Ce}_3\text{O}[\text{AsO}_3]_2\text{Cl}$ were determined using single crystal data. The compounds $\text{Ln}_5\text{O}_4[\text{AsO}_3]_2\text{Cl}$ have monoclinic unit cell parameters: $a = 12.493(3)$ Å, $b = 5.5884(13)$ Å, $c = 9.0776(19)$ Å, $\beta = 116.610(14)^\circ$ and $a = 12.2528(26)$ Å, $b = 5.6101(9)$ Å, $c = 8.9059(20)$ Å, $\beta = 115.945(16)^\circ$ for the Pr and Sm compounds, respectively. Both crystallize with space group $C2/m$, $Z = 2$ whereas $\text{Ce}_3\text{O}[\text{AsO}_3]_2\text{Cl}$ crystallises in space group $P4_{2}nm$, $Z = 4$ with unit cell parameters $a = 12.8590(6)$ Å, $c = 5.5627(3)$ Å. The structure of $\text{Ln}_5\text{O}_4[\text{AsO}_3]_2\text{Cl}$ contains three crystallographically independent Ln^{3+} sites with 8, 7 and 8 coordinated Ln^{3+} atoms whereas the $\text{Ce}_3\text{O}[\text{AsO}_3]_2\text{Cl}$ structure contains only two Ce^{3+} sites with 6 and 8 coordinated cerium atoms. The common features to these structures are isolated pyramidal $[\text{AsO}_3]^{3-}$ anions with a stereochemically active non-bonding electron pair.

[1] Kang D.-H., Wontcheu J., Schleid Th., *Solid State Sci.* **2009**, 11, 299. [2] Kang D.-H., Komm Th., Schleid Th., *Z. Kristallogr. Suppl.* **2005**, 22, 157.

Keywords: lanthanide oxide chloride oxoarsenate(III); crystal structure determination; single crystal diffraction