

Darstellung, Kristallstruktur, Schwingungsspektren und Normalkoordinatenanalyse von *trans*-[OsO₂(N₂H₂C₂O₂)₂]²⁻

Synthesis, Crystal Structure, Vibrational Spectra, and Normal Coordinate Analysis of *trans*-[OsO₂(N₂H₂C₂O₂)₂]²⁻

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Z. Naturforsch. **53 b**, 1338–1342 (1998); eingegangen am 12. August 1998

trans-Dioxo-bis(oxamido)osmate(VI), Synthesis, Crystal Structure, Vibrational Spectra, Normal Coordinate Analysis

The treatment of K₂[OsO₂(OH)₄] with oxamide in aqueous solution yields [OsO₂(N₂H₂C₂O₂)₂]²⁻. The crystal structure of *trans*-(Ph₄P)₂[OsO₂(N₂H₂C₂O₂)₂]·CH₂Cl₂ (triclinic, space group P $\bar{1}$, $a = 10.447(1)$, $b = 14.102(4)$, $c = 16.962(2)$ Å, $\alpha = 90.037(1)$, $\beta = 90.633(7)$, $\gamma = 98.812(2)^\circ$, $Z = 2$) has been determined by single crystal X-ray diffraction analysis. The IR and Raman spectra were measured at room temperature. Based on the molecular parameters of the X-ray determination a normal coordinate analysis has been performed and the vibrations are assigned. The valence force constants are $f_d(\text{Os}=\text{O}) = 6.7$, $f_d(\text{Os}-\text{N}) = 2.4$, $f_d(\text{C}-\text{N}) = 4.9$, $f_d(\text{C}=\text{O}) = 11.15$ and $f_d(\text{C}-\text{C}) = 4.7$ mdyn/Å.

* Sonderdruckanforderungen an Prof. Dr. W. Preetz.