

Stabilities and coordination modes of α -alaninephosphonic acid in copper(II) heteroligand complexes with ethylenediamine, diethylenetriamine or N,N,N',N',N'' -pentamethyldiethylene triamine in aqueous solution.

Autorzy

Anna Kamecka

Barbara Kurzak

Julia Jezierska

Agnieszka Woźna

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Streszczenie

Solution equilibrium studies on Cu^{2+} –L1–L2 ternary systems have been performed by pH-potentiometry, UV–Vis spectrophotometry and EPR methods (L1 corresponds to polyamines such as ethylenediamine (en), diethylenetriamine (dien), or N,N,N',N',N'' -pentamethyldiethylenetriamine (Me5dien) and L2 represents 1-aminoethylphosphonic acid (α -alaninephosphonic acid)). The obtained results suggest the formation of heteroligand complexes with $[\text{Cu}(\text{L1})(\alpha\text{-Ala}(\text{P}))]$ stoichiometry in all studied systems. Additionally, in the system with en the $[\text{Cu}(\text{en})(\alpha\text{-Ala}(\text{P}))\text{H}-1]^-$ species is formed in basic solution. Our spectroscopic results indicate tetragonal geometry for the $[\text{Cu}(\text{en})(\alpha\text{-Ala}(\text{P}))]$ species, geometry slightly deviated from square pyramidal for the $[\text{Cu}(\text{dien})(\alpha\text{-Ala}(\text{P}))]$ complex and strongly deviated from square pyramidal towards trigonal bipyramidal for the $[\text{Cu}(\text{Me5dien})(\alpha\text{-Ala}(\text{P}))]$ species. The coordination modes in these heteroligand complexes are discussed. Keywords Copper(II) complexes–Heteroligand complexes–Aminophosphonic acids–EPR–Vis spectroscopy–Potentiometry–Equilibria–Stability constants–Polyamines–Five-coordinate complexes

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