

Relationships between structural and spectroscopic properties of complexes of o-phenanthroline and 2,2'-bipyridine with lanthanide (Ln = Sm, Eu) thiocyanates.

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Rok wydania

2020

Czasopismo

Polyhedron

Numer woluminu

190

Strony

114770/1-114770/6

DOI

10.1016/j.poly.2020.114770

Kolekcja

Naukowa

Język

Angielski

Typ publikacji

Artykuł

Streszczenie

Three new complexes, [(NCS)₃(phenanthroline)₂(H₂O)Eu] and [(NCS)₃(2,2'-bipyridine)₂(H₂O)Ln] (Ln = Sm, Eu) have been synthesized as potential candidates for optical materials. The absorption spectra of the [(NCS)₃(2,2'-bipyridine)₂(H₂O)Sm] crystal were measured at 293 K and 8 K. For all of these complexes emission and excitation spectra as well as luminescent decay curves at room temperature and 77 K were recorded. Additionally, the quantum yields of luminescence were determined. The crystal structure of [(NCS)₃(phen)₂(H₂O)Eu] complex was determined by X-ray diffraction analysis. The [(NCS)₃(phen)₂(H₂O)Eu] compound crystallizes in a monoclinic system with the space group *P*2₁/*c*. XRPD patterns of [(NCS)₃(2,2'-bipyridine)₂(H₂O)Ln] (Ln = Sm, Eu) reveals that they are isostructural. The inner sphere coordination of the lanthanide ions is composed of three isothiocyanate nitrogen atoms, four nitrogen atoms from the organic ligand (1,10-phenanthroline or 2,2'-bipyridine) and a water molecule. The luminescent studies of complexes discussed in relation to their crystal structures. The intensities of the f–f transitions in [(NCS)₃(2,2'-bipyridine)₂(H₂O)Sm] crystal were determined and examined in accordance to the Judd–Ofelt theory. The modification of ratio β and Sinha's parameter δ and their dependence on the nature of lanthanide-ligand bond is discussed for the samarium complex.

Słowa kluczowe

O-phenanthroline, molecule, 2'- bipyridine, Isothiocyanate, UV-Vis spectroscopy, Lanthanides

Adres publiczny

<http://dx.doi.org/10.1016/j.poly.2020.114770>

Strona internetowa wydawcy

<http://www.elsevier.com>

Plik został wygenerowany dnia 2026-05-13 11:50:41

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