

## Europium and terbium pyrrole-2-carboxylates : structures, luminescence, and energy transfer.

### Autorzy

Konstantin P. Zhuravlev  
Łukasz Michnik  
Paula Gawryszewska  
Vera I. Tsaryuk  
Valentina A. Kudryashova

### Rok wydania

2019

### Czasopismo

Inorganica Chimica Acta

### Numer woluminu

492

### Strony

1-7

### DOI

10.1016/j.ica.2019.04.014

### Kolekcja

Naukowa

### Język

Angielski

### Typ publikacji

Artykuł

### Streszczenie

Lanthanide pyrrole-2-carboxylates  $[\{Ln(L)_3(H_2O)_2\} \cdot H_2O]_n$  ( $Ln=Eu, Gd, Tb$ ) were synthesized and investigated using single crystal X-ray diffraction and optical spectroscopy (luminescence, luminescence excitation and vibrational IR spectra). The europium pyrrole-2-carboxylate crystallizes as a 2D network structure in the  $P2_1/c$  space group. The  $Eu^{3+}$  coordination polyhedron formed by five carboxylic groups and two water molecules can be described as a distorted single-capped square antiprism. The luminescence spectra indicate a low symmetry of the charge distribution around the  $Ln^{3+}$  ion that corresponds to the  $C_1$  symmetry of the coordination environment of  $Eu^{3+}$  ion in the  $[\{Eu(L)_3(H_2O)_2\} \cdot H_2O]_n$  crystal lattice. The low-energy ligand–metal charge transfer (LMCT) state was identified in the europium compound. The shortening of the  $^5D_0$  ( $Eu^{3+}$ ) lifetime in the 230–295K temperature region, when the compound is heated from 77 to 295K, and the low luminescence intensity of the europium pyrrole-2-carboxylate are caused by the participation of the LMCT state in quenching processes. The energy of activation determined for the  $^5D_0$  – LMCT crossover is  $3900 \pm 100 \text{ cm}^{-1}$ .

### Słowa kluczowe

$Eu^{3+}$ ,  $Tb^{3+}$ , Pyrrole-2-carboxylate, Luminescence, LMCT, X-ray crystal structure

### Adres publiczny

<http://dx.doi.org/10.1016/j.ica.2019.04.014>

### Strona internetowa wydawcy

<http://www.elsevier.com>