Accepted Manuscript

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PII:	S0167-577X(18)30340-9
DOI:	https://doi.org/10.1016/j.matlet.2018.02.126
Reference:	MLBLUE 23952
To appear in:	Materials Letters
Received Date:	15 January 2018
Revised Date:	19 February 2018
Accepted Date:	26 February 2018



Please cite this article as: S. Stojadinović, N. Tadić, R. Vasilić, Down-conversion photoluminescence of ZrO₂:Er³⁺ coatings formed by plasma electrolytic oxidation, *Materials Letters* (2018), doi: https://doi.org/10.1016/j.matlet.2018.02.126

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Down-conversion photoluminescence of ZrO₂:Er³⁺ coatings formed by plasma

electrolytic oxidation

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Abstract

Plasma electrolytic oxidation of zirconium in alkaline solution containing Er_2O_3 powder was used for preparation of $ZrO_2:Er^{3+}$ coatings. Photoluminescence (PL) emission spectra of $ZrO_2:Er^{3+}$ excited by ultraviolet irradiation are composed of broad PL band associated with ZrO_2 host and sharp bands corresponding to *f*–*f* transitions of Er^{3+} . The strongest green PL emission band of Er^{3+} in the range from 540nm to 580nm is assigned to ${}^4S_{3/2} \rightarrow {}^4I_{15/2}$ transition. The PL excitation spectra of $ZrO_2:Er^{3+}$ characterize broad band from 250nm to 350nm associated with charge transfer state of Er^{3+} and the series of peaks in the range from 350nm to 530nm which are associated with 4*f* transitions of the Er^{3+} from ground state ${}^4I_{15/2}$ to higher levels. Obtained results allowed the identification of down-conversion PL mechanism.

Keywords: Plasma electrolytic oxidation; ZrO₂; Er³⁺; Luminescence; Phosphors.

1. Introduction

 ZrO_2 has been widely used as a highly efficient host matrix for trivalent rare-earth ions for the fabrication of photoluminescent materials because of its low phonon frequency (about $470cm^{-1}$) as well as excellent chemical, photo-chemical and photo-thermal stability, high refractive index, wide optical band gap, high transparency in the visible and near infrared region, etc. [1,2]. Trivalent rare-earth ions are characterized by a partially filled 4*f* shell that is Download English Version:

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