## Electrodeposition of luminescent composite metal coatings

## containing rare-earth phosphor particles

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**Electronic supplementary information (ESI)** 

Figure S1. Cyclic voltammogram of an electrolyte with composition 0.593:0.391:0.016 mole fractions of acetamide-DMSO<sub>2</sub>-NiCl<sub>2</sub> on a Pt electrode at 130 °C.



Figure S2. Cyclic voltammogram of an electrolyte with composition 0.617:0.366:0.017 mole fractions of acetamide-DMSO<sub>2</sub>-CoCl<sub>2</sub> on a Pt electrode at 130 °C.



Figure S3. SEM pictures of Ni- $Y_2O_3$  coatings, electrodeposited from electrolyte solutions with different concentration of  $Y_2O_3$  (For experimental conditions: see Table 1)



Figure S4. SEM pictures of Ni- $Y_2O_3$  coatings, electrodeposited from electrolyte solutions with different concentration of  $Y_2O_3$  (For experimental conditions: see Table 1). The coatings are the same as those shown in Figure S3, but at a different magnification.



Table S1. Effect of the deposition potential on the loading of  $Y_2O_3$  particles (at 130 °C).

Potential vs. Ni	atomic % of O	atomic % of Y	atomic % of Ni	vol. % of Y <sub>2</sub> O <sub>3</sub>
(V)				
-0.2	14.5	9.1	76.4	29
-0.4	19.4	15.6	65.0	45
-0.8	19.4	15.3	65.3	45

Table S2. Influence of temperature on Y<sub>2</sub>O<sub>3</sub> codeposition in nickel.<sup>[a]</sup>

temperature (°C)	atomic % of O	atomic % of Y	atomic % of Ni	vol. % of $Y_2O_3$
130	19.5	15.5	65.0	45
150	15.3	11.3	73.4	35
[a] Deposits obtaine	d at _0 / V vs Ni	with stirring (250 r	nm) in an electroly	te bath with 100 a

<sup>[a]</sup> Deposits obtained at -0.4 V vs. Ni, with stirring (250 rpm), in an electrolyte bath with 100 g  $L^{-1}$  of Y<sub>2</sub>O<sub>3</sub>.

Figure S5. Effect of the deposition potential on the morphology of  $Ni-Y_2O_3$  coatings obtained at 130 °C (SEM pictures at two different magnifications, left and right).



Figure S6. Influence of temperature on  $Y_2O_3$  codeposition in nickel. SEM pictures (left) and EDX spectra (right).



Figure S7. SEM pictures of Co- $Y_2O_3$  coatings, electrodeposited from acetamide:DMSO<sub>2</sub>:CoCl<sub>2</sub> electrolyte solutions with different concentration of  $Y_2O_3$ , and different deposition potentials.



Figure S8. EDX spectrum of a Co-Y<sub>2</sub>O<sub>3</sub> coating.



Figure S9. SEM pictures of: (a) Ni coatings with  $Y_2O_3$ :Eu<sup>3+</sup> phosphor particles, and (c) Ni coatings with  $Gd_2O_2S$ :Tb<sup>3+</sup> phosphor particles, deposited at -0.4 V vs. Ni at the temperature of 130 °C. The electrolyte was stirred at 250 rpm. (b) and (d) are the EDX spectra of (a) and (c), respectively.



Figure S10. SEM pictures of: (a) Ni coatings with  $BaMg_2Al_{16}O_{27}$ :Eu<sup>2+</sup> phosphor particles, and (c) Ni coatings with  $Y_3Al_5O_{12}$ :Ce<sup>3+</sup> phosphor particles, deposited at -0.4 V vs. Ni at the temperature of 130 °C. The electrolyte was stirred at 250 rpm. (b) and (d) are the EDX spectra of (a) and (c), respectively.







Figure S12. Excitation spectra of  $Gd_2O_2S:Tb^{3+}$  powder ( $\lambda_{em} = 545$  nm) as a function of the thickness of the powder layer.



Figure S13. Pictures of a nickel coating with  $Y_3Al_5O_{12}$ :Ce<sup>3+</sup> phosphor particles, illuminated (a) by daylight and (b) with UV radiation of 340 nm.



Figure S14. Room temperature luminescence spectra of  $Y_3Al_5O_{12}$ :Ce<sup>3+</sup> excited at different wavelengths.



Figure S15. Room temperature emission spectra under excitation at 254 nm of the mixtures of red ( $Y_2O_3:Eu^{3+}$ ) and green ( $Gd_2O_2S:Tb^{3+}$ ) phosphors in (a) 90:10, (b) 75:25, (c) 50:50, (d) 25:75 and (e) 10:90 wt.% ratio, respectively.



Table S3. Luminescence lifetimes ( $\tau_{obs}$ , ms) of Eu( ${}^{5}D_{0}$ ) and Tb( ${}^{5}D_{4}$ ) levels for red (Y<sub>2</sub>O<sub>3</sub>:Eu<sup>3+</sup>) and green (Gd<sub>2</sub>O<sub>2</sub>S:Tb<sup>3+</sup>) phosphors, the mixtures of them and the corresponding composite Ni coatings.

Ratio Eu:Tb / wt. %	State	$Eu(^{5}D_{0})$	$\text{Tb}(^{5}\text{D}_{4})$
100:0	powder	1.04(1)	-
	Ni coating	1.04(1)	-
90:10	powder	1.03(1)	0.57(1)
75:25	powder	1.02(1)	0.56(1)
	Ni coating	1.00(1)	0.56(1)
50:50	powder	1.03(1)	0.56(1)
	Ni coating	1.02(1)	0.58(1)
25:75	powder	1.02(1)	0.58(1)
	Ni coating	1.00(1)	0.57(1)
10:90	powder	1.02(1)	0.57(1)
0:100	powder	-	0.55(1)
	Ni coating	-	0.55(1)

Table S4. CIE coordinates of the red  $(Y_2O_3:Eu^{3+})$  and green  $(Gd_2O_2S:Tb^{3+})$  phosphors, the mixtures of them and the corresponding composite Ni coatings.

Ratio Eu:Tb / wt. %	State	( <i>x</i> ; <i>y</i> )
100:0	powder	0.652; 0.345
	Ni coating	0.638; 0.346
75:25	powder	0.563; 0.408
	Ni coating	0.578; 0.401
50:50	powder	0.515; 0.464
	Ni coating	0.489; 0.471
25:75	powder	0.414; 0.522
	Ni coating	0.410: 0.523
0:100	powder	0.411; 0.537
	Ni coating	0.373; 0.568

Figure S16. Room-temperature emission spectra of (a) the 50:50 wt.% mixture of blue and yellow phosphors and (b) the corresponding Ni coating. Black lines:  $\lambda_{ex} = 254$  nm, green lines:  $\lambda_{ex} = 340$  nm.



Figure S17. (Left) Photograph of the powder of 50:50 wt.% mixture of yellow (YAG: Ce<sup>3+</sup>) and blue (BAM: Eu<sup>2+</sup>) phosphors and the corresponding composite Ni coating under 254 nm excitation and (right) CIE trichromatic coordinates (circles: powder, triangles: Ni coating; black:  $\lambda_{ex} = 254$  nm, red:  $\lambda_{ex} = 340$  nm).



Figure S18. SEM pictures (two different magnifications) and EDX spectrum of binary mixtures of the red  $(Y_2O_3:Eu^{3+})$  and green  $(Gd_2O_2S:Tb^{3+})$  phosphors, co-deposited with Ni at -0.4 V vs. Ni.



