

**1st International Electronic
Conference on Materials**

26 May - 10 June 2014

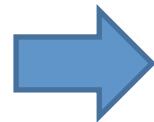
**Sol-gel Synthesis and Antioxidant Properties of Yttrium
Oxide Nanocrystallites Incorporating P-123**

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Arturo López-Marure, Perla Yolanda López-Camacho Angel
de Jesús Morales-Ramírez, Hiram Isaac Beltrán-Conde

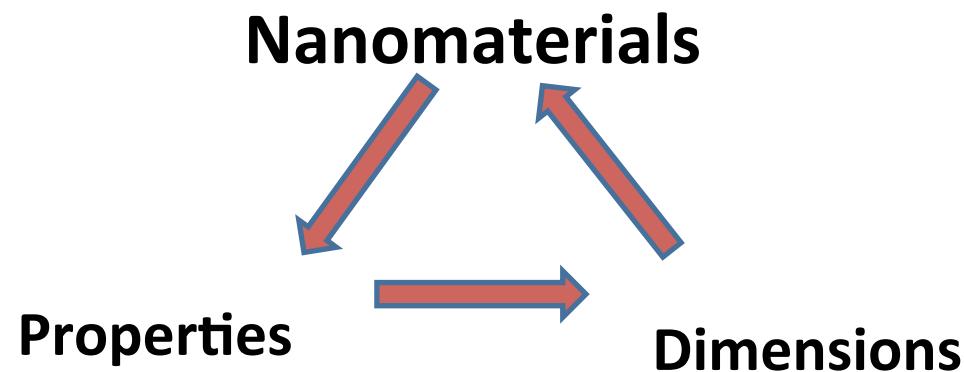
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NANOMATERIALS: Small structures with high potential

**Nanomaterials
Nanoestructurs
Nanoparticles**



**Nanoscience
Nanotechnology**

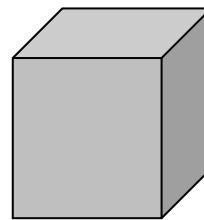


Nanomaterials
1 - 100 nm
one dimension

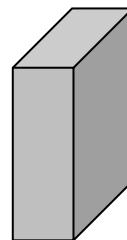
Ceramics
Metals
Semiconductors
Polymers
Combination of them

In nanometer scale the properties are not the same to bulk materials.

Clasification



3D



2D

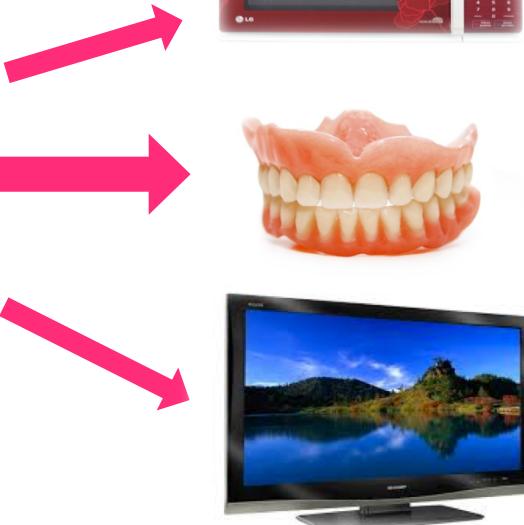


1D

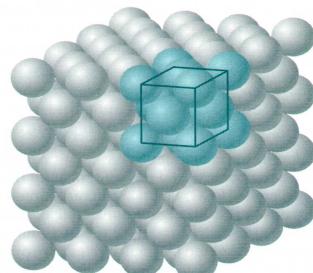
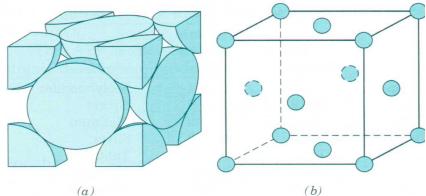


0D

Applications of Y_2O_3



Cubic structure
of Y_2O_3
[PDF cart
201412]

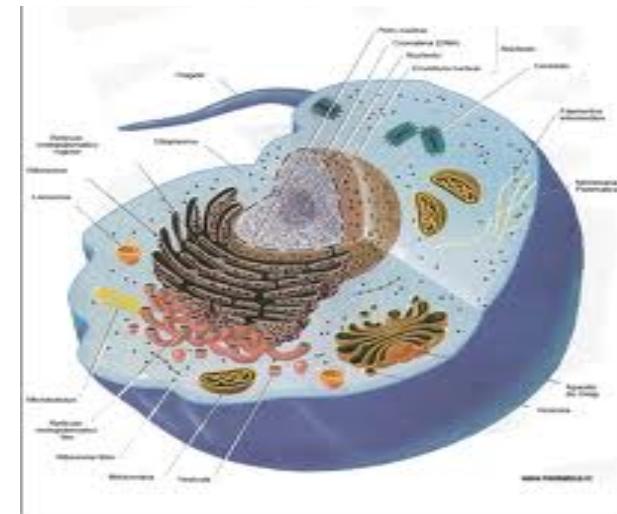


Applications

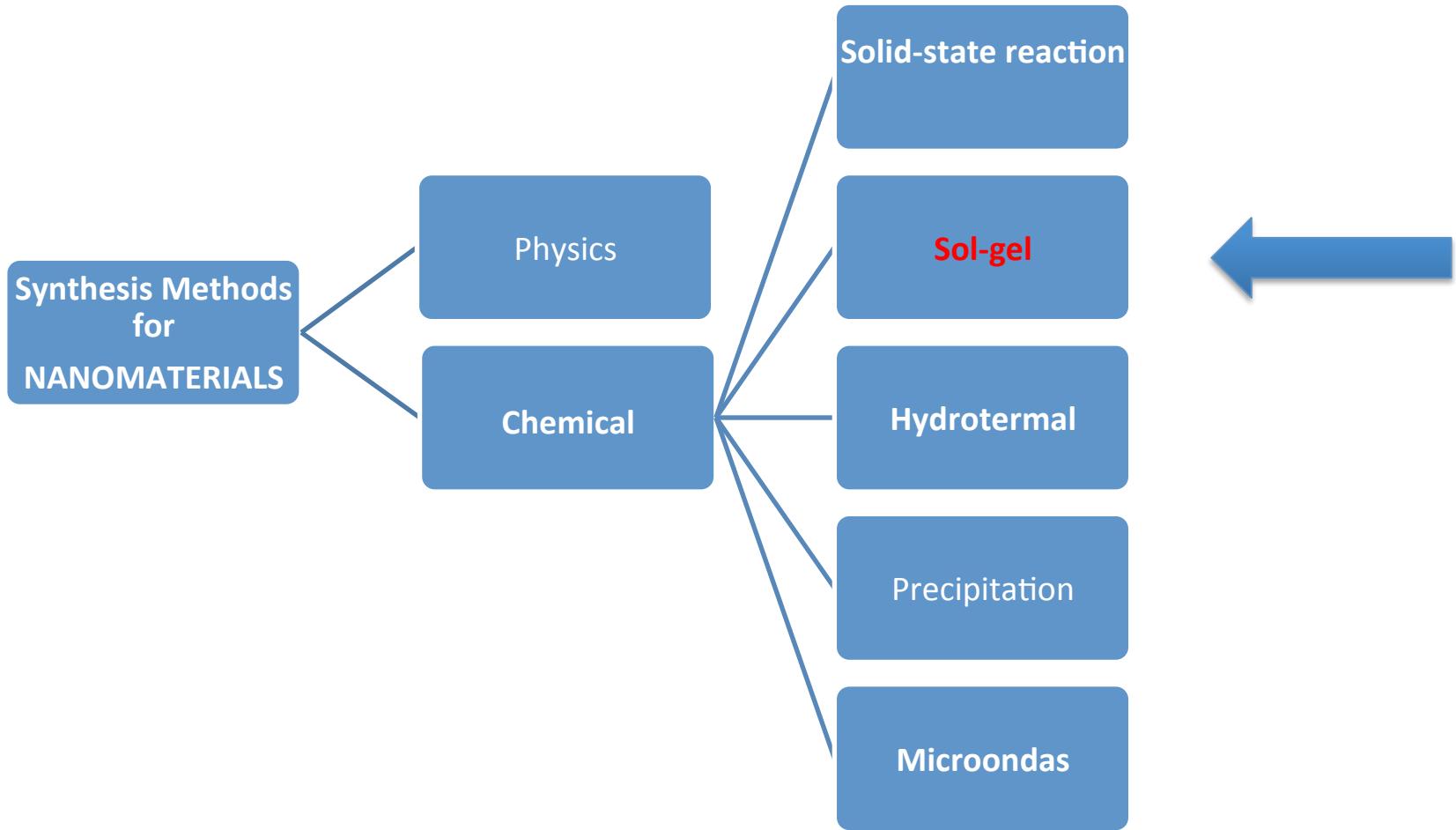
- ✓ Catalyst
- ✓ Medicine
- ✓ Biology
- ✓ Electronic
- ✓ Optique

Antioxidants

Biological affinity



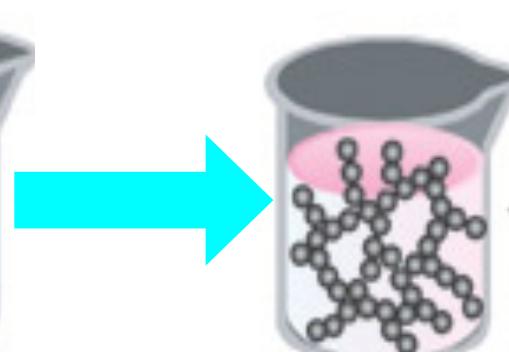
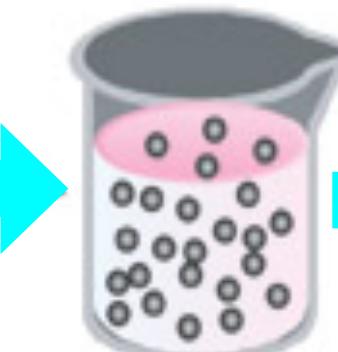
Synthesis Methods of Y₂O₃



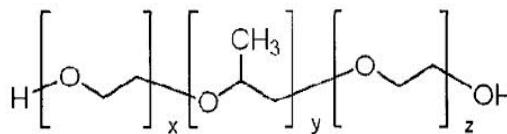
SYNTHESIS PROCEDURE OF Y_2O_3

Precursors:

- ✓ $[\text{Y}(\text{NO}_3)_3]$
- ✓ $[\text{YCl}_3]$
- ✓ $[\text{CH}_3\text{OH}]$
Acetylacetone
- ✓ Poloxámero P-123



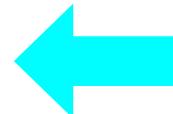
XEROGEL / 24 h a 90 °C



Densify and cristallize:

Heat treatmeat at 270 °C (2 h), 500 °C, 700 °C, 800 °C y 900 °C for 1 h

Chemical,
estructural and
antioxidant
characterization
of Y_2O_3
nanocrystallites



SYNTHESIS PROCEDURE OF Y_2O_3

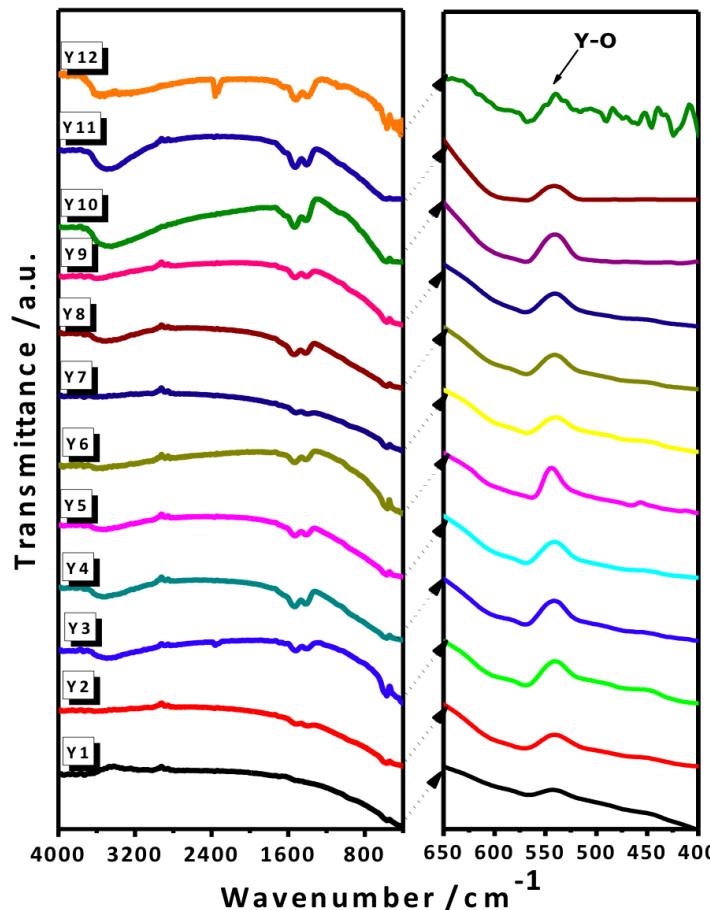
Table 1. Y_2O_3 systems prepared by sol-gel method, key words and general description.

Keywords	Precursor	Matrix	Poloxamer	Y:P123 Molar Rat.	T / °C	Crystalite size / (nm)
Y1					700	26
Y2	$\text{Y}(\text{NO}_3)_3$		-	-	800	27
Y3					900	26
Y4					700	32
Y5	YCl_3	Y_2O_3	-	-	800	29
Y6					900	29
Y9			P-123	1:1	900	21
Y10	$\text{Y}(\text{NO}_3)_3$			2:1	900	28
Y11			P-123	1:1	900	29
Y12	YCl_3			2:1	900	29

RESULTS

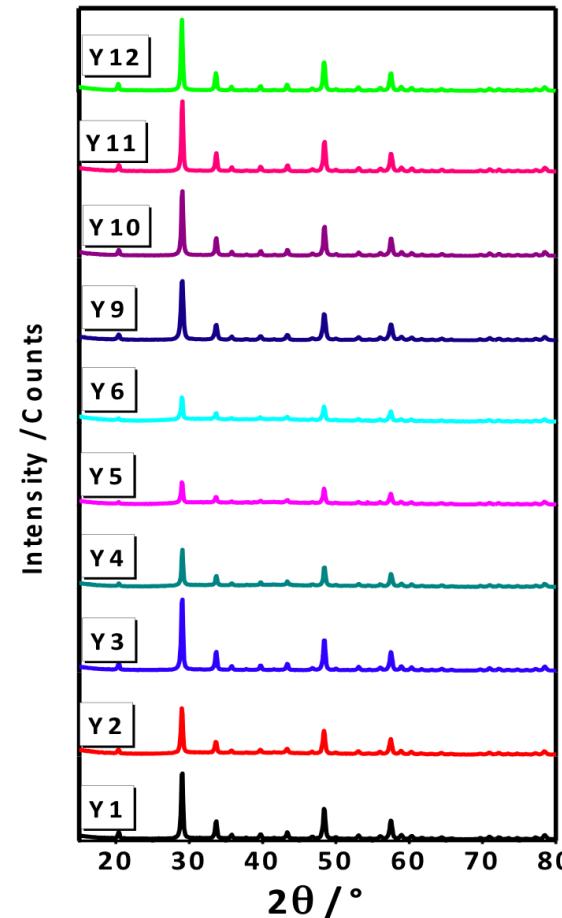
FTIR – DRX of Y_2O_3

Figure 1. IR spectra of Y_2O_3 systems heat treated at 700 and 900 °C.



In the samples Y1 and Y2 is only observed the absorption band of oxygen-metal in around 500 and 600 cm^{-1}

Figure 2. XRD patterns of Y_2O_3 systems heat treated at 700 and 900 °C.



No secondary phases are found. It can be noticed that the cubic structure of Y_2O_3 is formed at 700 °C and its remained stable until 900 °C [PDF cart 201412].

DRX of Y_2O_3

Figure 3. Y_2O_3 systems prepared from yttrium nitrate and yttrium chloride at different temperatures.

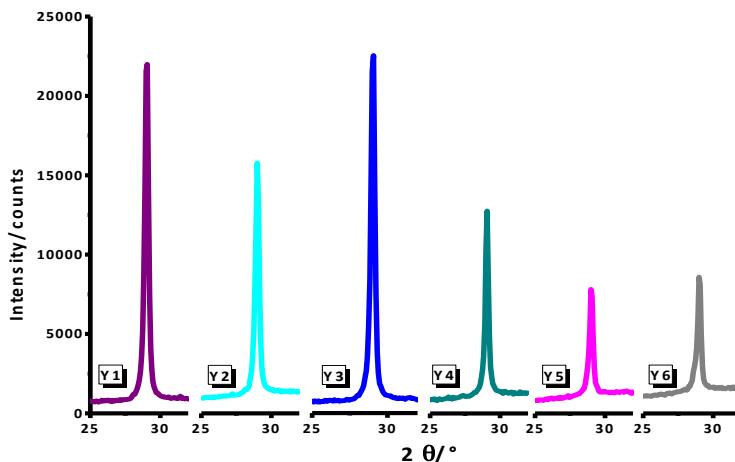


Figure 4. Y_2O_3 powder synthetized from yttrium nitrate with and without P-123 poloxamer heat treated at $900\text{ }{}^\circ\text{C}$ for 1 hour.

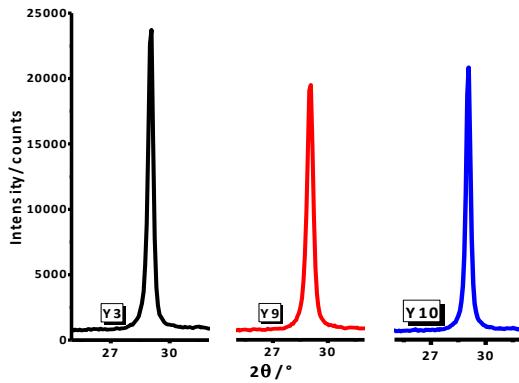
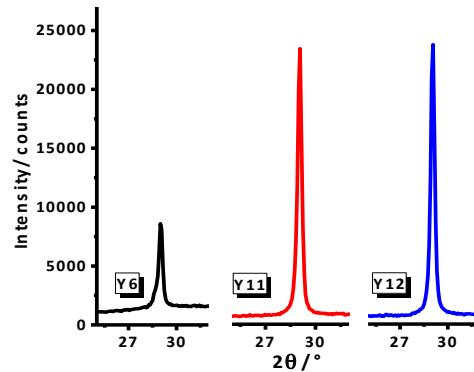


Figure 5. Y_2O_3 powder synthetized from yttrium chloride with and without P-123 poloxamer heat treated at $900\text{ }{}^\circ\text{C}$ for 1 hour.

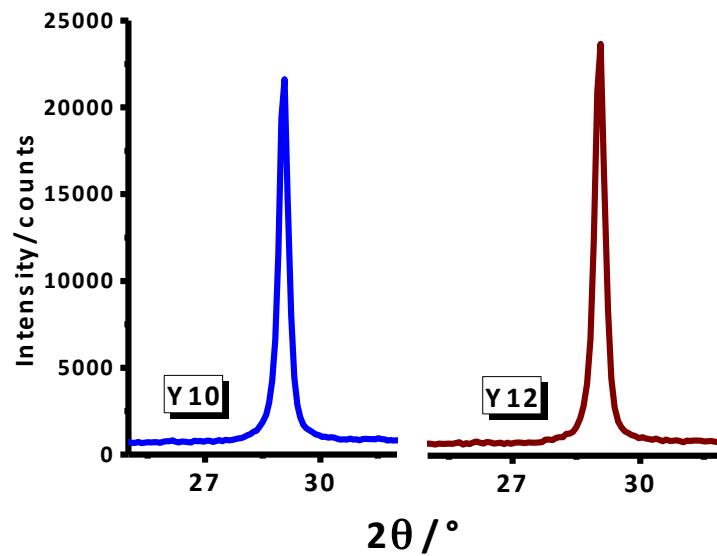


The samples synthesized from yttrium nitrate (Y1-Y3) showed a greater degree of crystallization compared with synthesized from yttrium chloride (Y4-Y6).

The crystallization degree comparison of the system prepared from yttrium nitrate and yttrium chloride with and without P-123 poloxamer.

DRX of Y_2O_3

Figure 6. XRD pattern of Y_2O_3 powder synthesized from yttrium nitrate and yttrium chloride using P-123 poloxamer in a molar ratio of P-123:Y, 2:1.



The XRD results revealed that the yttrium oxide systems embedded in P-123 poloxamer in a molar ratio of P-123:Y; 2:1 presented better crystallization degree.

ANTIOXIDANT ASSAYS of Y_2O_3

Figure 7. Time dependent of DPPH without nanocrystal.

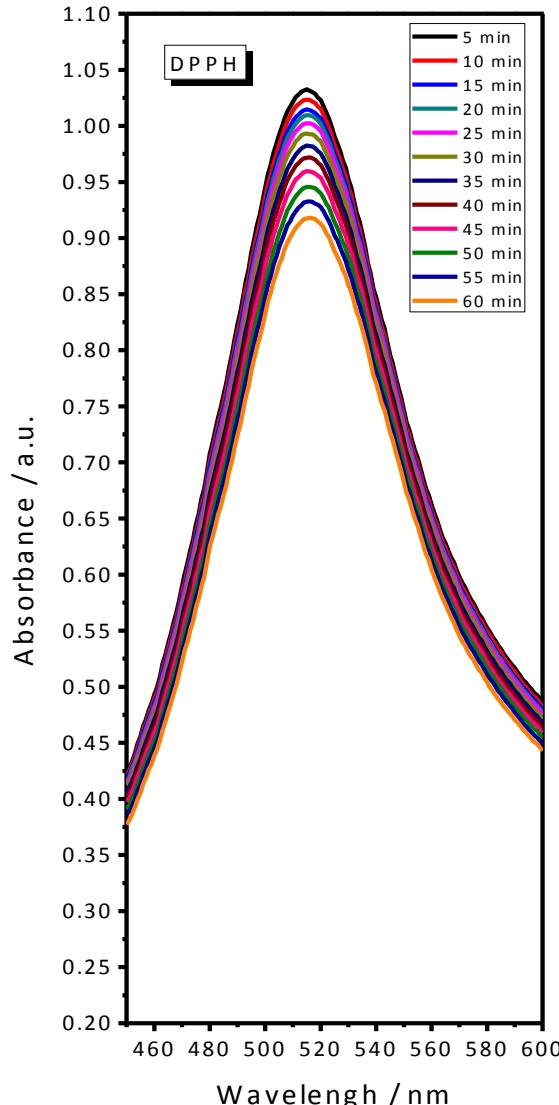
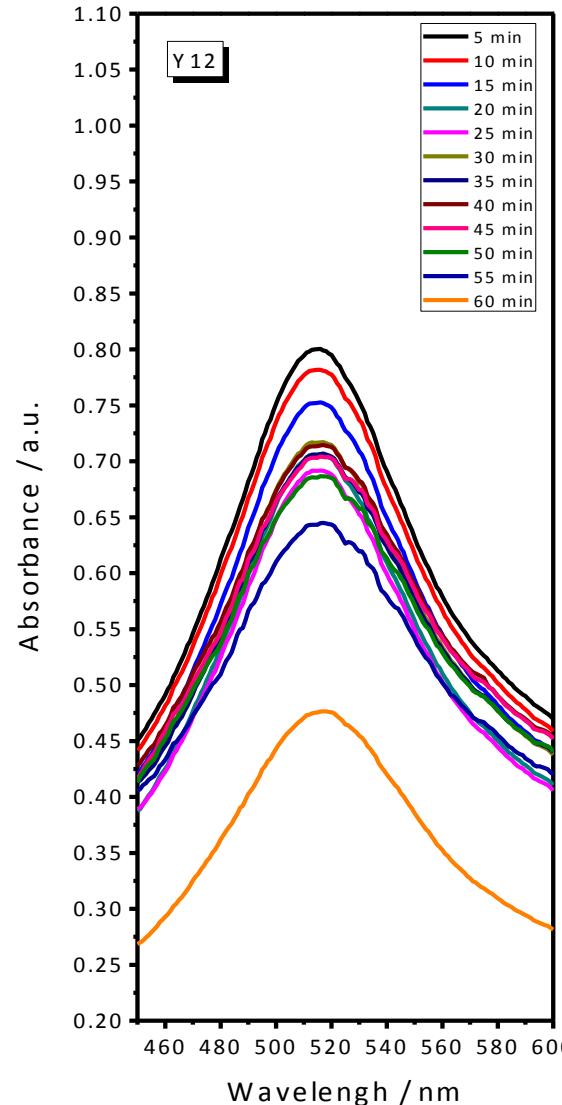


Figure 8. Time dependent of DPPH scavenging by yttrium oxide nanocrystallites (sample Y12).



The DPPH without nanoparticles does not reveal changes in absorption characteristic peak.

the diminishing of the DPPH begin from first 5 min and it is evident after 60 min. It is known that the antioxidant property may be due to the neutralization of free radical character of DPPH which is by transfer a electron between the reactant.

CONCLUSION

- ✓ Yttrium oxide nanocrystallites were successfully synthesized by sol-gel method from yttrium nitrate and yttrium chloride as precursor.
- ✓ The yttrium oxide nanostructured powders elaborated from yttrium chloride and embedded in P-123 poloxamer in a molar ratio of P-123:Y 2:1 presented better physicochemical properties (crystallinity and purity) than systems prepared from yttrium nitrate precursor.
- ✓ Yttrium oxide powder presented crystallites size in the range of 21 to 32 nanometers.
- ✓ The DPPH studies are reported for first time for yttrium oxide synthesized by sol-gel method due to a directly comparison cannot be made.
- ✓ Yttrium oxide nanocrystallites show enhanced antioxidant potency which leading a new promising material in biological system.

Acknowledgments

The authors acknowledge to the financial support of
SEP-ConacyT 178817, PROMEP 47310345 and
UAM-C-CA-23 projects.

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THANK YOU!

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