## Nanofibers Pure And Doped With a Transition Metal: BaTiO<sub>3</sub> and LiNbO<sub>3</sub>

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Nowadays one dimensional nanomaterials such as nanofibers, have been synthesized by various processes, e.g. solution method, sol-gel, laser ablation, chemical vapor deposition (CVD), hydrothermal method and mechanochemical activation. In this work, structures ABO<sub>3</sub> such as BaTiO<sub>3</sub> and LiNbO<sub>3</sub> nanofibers were doped with Mn like transition metal and synthesized by electrospinning method, a detailed description of the procedure can be found in the literature [1][2]; this technique has been recognized as an efficient method to make polymeric nanofibers [3], who it is a straightforward way to synthesize nanostructures.

The potentials applications of these materials are focused in one of the extensively studied ferroelectric material with wide range of applications in non-volatile ferroelectric random access memories, as transducers, sensors and actuators, etc [4].

The presence of a pure phase and patterns from BaTiO<sub>3</sub> doped is confirmed by XRD analysis, Fig 1. In the other hand, the presence of a pure phase and patterns from LiNbO<sub>3</sub> doped is confirmed in Fig 2. The metal transition is used to dope the composite taking care the BaTiO<sub>3</sub> and LiNbO<sub>3</sub> stoichiometry. Such as Fig 1. and Fig 2. show the "x" value, that in both cases corresponds to 2.5, 5 and 10%. In BaTiO<sub>3</sub> doped to the 10% change it from tetragonal to hexagonal structure.

Fig 3. and Fig 4. show a SEM micrograph of as-spun fibers, BaTiO<sub>3</sub> and LiNbO<sub>3</sub> respectively. Cylindrical and randomly oriented BaTiO<sub>3</sub> fibers with diameter about 57-453 nm were obtained, compared with diameter about 57-146 nm obtained by LiNbO<sub>3</sub>.

TEM micrographs, Fig 5. and Fig 6, show a isolated and calcined  $BaTi_{0.95}Mn_{0.05}O_3$  and  $LiNb_{0.95}Mn_{0.05}O_3$  nanofibers respectively, both in the same Mn concentration. In Fig 5, it can be observed fibers with few  $\mu m$  in length and an irregular morphology. Fig. 6 shows TEM micrograph, different surface morphology is evident.

## References:

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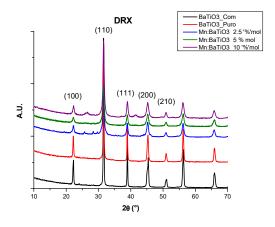


Figure 1. XRD pattern BaTi<sub>1-x</sub>Mn<sub>x</sub>O<sub>3</sub>

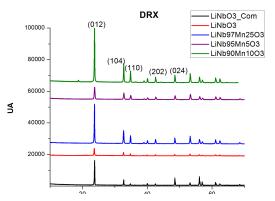
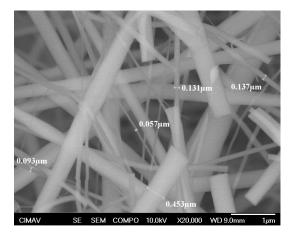
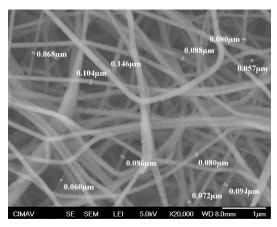


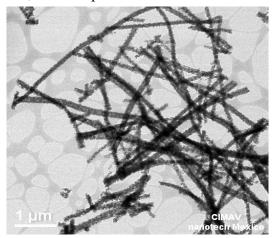
Figure 2. XRD pattern LiNb<sub>1-x</sub>Mn<sub>x</sub>O<sub>3</sub>



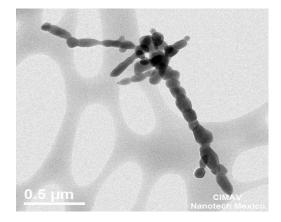
**Figure 3.** SEM images of as-spun Ba(C<sub>2</sub>H<sub>3</sub>O<sub>2</sub>)<sub>2</sub>:Ti[OCH(CH<sub>3</sub>)<sub>2</sub>]<sub>4</sub>:PVP:CH<sub>3</sub>CO OH:H<sub>2</sub>O:C<sub>2</sub>H<sub>5</sub>OH composite.



**Figure 4.** SEM images of as-spun HLiO:Nb(OCH<sub>2</sub>CH<sub>3</sub>)<sub>5</sub>:PVP:CH<sub>3</sub>COOH: C<sub>2</sub>H<sub>5</sub>OH. composite.



**Figure 5.** TEM image of  $BaTi_{0.95}Mn_{0.05}O_3$  nanofibers.



**Figure 6.** TEM image of  $LiNb_{0.95}Mn_{0.05}O_3$  nanofibers.