ELECTRONIC STRUCTURE AND HYDRO-OXIDATION OF LaNiO_{3-δ} THIN FILMS

S. Mickevičius^{1*}, S. Grebinskij¹, V. Bondarenka¹, H. Tvardauskas¹, M. Senulis¹, V. Lisauskas¹, K. Sliužienė¹, and <u>B.A. Orlowski²</u>

> ¹ Semiconductor Physics Institute, A. Goštauto 11, 00-00 Vilnius, Lithuania ² Institute of Physics, PAS, Al. Lotnikow 32/46, 02-668 Warsaw, Poland

Keywords: synchrotron radiation, X-ray photoelectron spectroscopy, LaNiO₃

*) e-mail: orbro@ifpan.edu.pl

LaNiO₃ is one of the few conductive oxides with a crystal structure suitable for integration in epitaxial heterostructures with perovskites of enormous potential technological colossal such as magnetoresistance materials, high-temperature superconductors and ferroelectrics.

It is known that electronic structure of LaNiO_{3- δ} strongly depends on the degree of stoichiometry δ and metal-dielectric transition take place at $\delta \approx 0.25$ [1]. Moreover, the considerable segregation of elements takes place in chemically synthesized LaNiO_{3-z} samples i.e. the surface concentrations of various species differ from the volume one. Another factor to be considered is the tendency of rare earth and nickel oxides to absorb water vapor.

In the previous paper [2] it was shown that the interaction with ambient water lead to the formation of lanthanum and nickel hydroxide containing phase at LaNiO_{3- δ} surface. The thickness of hydroxide enriched ~ 2 nm layer was estimated by means of Tunable High-Energy X-ray photoelectron spectroscopy using synchrotron radiation

In this paper, in addition to resistivity measurements, we utilize the surface sensitivity of XPS to study the influence of preparation conditions and heat treatment in various atmospheres on the properties of LaNiO_{3- δ} film.

It was found that after annealing in vacuum at 700°C the relative concentration of hydroxide species increases and electrical conductivity becomes dielectric in nature.

The LaNiO_{3- δ} films remain metallic after hightemperature (750°C) annealing in an oxygen atmosphere, while concentration of the hydroxide species slightly decreases.

Unfortunately, the ESCA sensitivity is insufficient to determine surfaced chemical composition with an accuracy enough to distinguish are the oxygen vacancies or La/Ni hydroxide species responsible for metaldielectric transition after high-temperature annealing in vacuum. Acknowledgements: This work was done in part within the research projects 72/E-67/SPB/DESY/P-03/DWM 68/2004-20061 and EC program G1MA-CT-2002-4017 (Center of Excellence CEPHEUS) and P03B 053 26.

References

- [1] M. Abbate, G. Zampieri, F. Prado, A. Caneiro, J.M. Gonzalez-Calbet, M. Vallet-Regi, "Electronic structure and metal-insulator transition in LaNiO_{3-δ}", *Phys. Rev. B* 65 (2002) 155101-155106.
- [2] S. Mickevicius; S. Grebinskij, V. Bondarenka, V. Lisauskas, K. Sliuziene, H. Tvardauskas, B. Vengalis, B.A. Orlowski, V. Osinniy, W. Drube, "The surface hydro-oxidation of LaNiO_{3-x} thin films", *Acta Phys. Polon. A* **112** (2007) 113-120

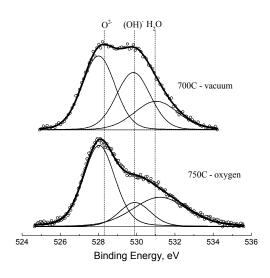


Figure 1. O 1s XPS spectra of LaNiO_{3- δ} films annealed in vacuum and in oxygen.