## Structural and optical Properties of Anodized p-InP(100) in 1M HF

U. Schlierf<sup>1</sup>, P. Schmuki<sup>1</sup>, D. J. Lockwood<sup>2</sup>, and M. J. Graham<sup>2</sup>

<sup>1</sup> Department of Material Science, LKO, University of Erlangen – Nuremberg, Martensstr. 7, D-91058 Erlangen, Germany

<sup>2</sup> Institute for Microstructural Sciences, National Research Council of Canada, Ottawa, Ontario, Canada K1A 0R6

In recent years, the localized dissolution of semiconductors has generated a great deal of renewed interest as a result of light emitting porous silicon and its large number of potential applications (see e.g., Refs. in [1-4]). Also for other semiconductor materials such as III-V compounds, research on localized dissolution has increased in the last few years (see e.g., Refs. in [3]).

The anodization of p-type semiconductors in acidic solutions, leads normally to electropolishing of the surface. It is well known only for p-type silicon that in HF containing solutions porosification takes place [5]. In this work it was found that by a potentiostatic anodic polarization of p-type InP in 1M HF instead of electropolishing a modifycation of the surface occurs. The surface exhibited blue, red and yellow colours. In the SEM images a porous layer with different structures is obtained (Fig.1). Polarization experiments were done for different times and at different potentials. All curves showed a linear current density decrease. The slope of the current density drop and the current density itself increases with increasing potentials. As a result of previous work on the porosification of n-type InP, where different pore morphologies appeared by using different halogenic acids [6], polarization was also carried out in HCl. Here, however, only electropolishing occured. Surface analyses of the HF-samples show that the different regions have also different compositions of indium- and phosphorus-oxides. The direction of growth of these oxides is from the air/electrolyte interface downwards to the edge of the immersed sample (Fig. 2). The spatial expansion and the growth rate are determined by the polarization time and the applied potential. The structure on the p-InP surface under the oxide shows crystallographic ordered etching trenches (Fig. 3). The formation of these oxides starts only at the air/electrolyte interface, because by covering this interface with a varnish, no surface modification occured. PL measurements show a bright visible photoluminescence from these porous layers (Fig. 4).

- [1] R.L. Smith, S.D. Collins, J. Appl. Phys., 71, R1 (1992)
- [2] S.M. Prokes, Interface, 3, 41 (1994)
- [3] A.G. Cullis, L.T. Canham, P.D.J. Calcott, J. Appl. Phys., 82, 909 (1997)
- [4] D.J. Lockwood, Editor Light Emission in Silicon: From Physics to Devices. Academic Press, San Diego, (1997)
- [5] V. Lehmann, S. Roennebeck, J. Electrochem. Soc., 146, (8), 2968 (1999)
- [6] P. Schmuki, L. Santinacci, T. Djenizian, D.J. Lockwood, *Phys. Stat. Sol. (a)*, **182**, 51 (2000)



Fig. 1: Porous oxide formed on p-InP in 1M HF after polarization at 3000mV for 180s



Fig. 2: Schematic drawing of the immersed sample and the growth direction of the oxides



Fig. 3: Crystallographic ordered etching trenches on the p-InP surface after the lift-off of the porous layer



Fig. 4: PL-spectra of p-InP of different coloured regions after polarization at 3000mV for 180s in 1M HF