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What we have?

1984 – X-Ray full pumping

1986 - X-Ray diffraction focusing and defocusing in the presence of external excitations

1986- Diffraction of thermal neutrons in piezoelectric single crystals under the influence of external fields.

1987- X-Ray detector based on porous matters

- 1988 Information transfer by means of X-Rays
- 1988 X-Ray diffraction on SAW.
- 1989 Formation of Modulating X-Ray spectroscopy
- 1989 Synthesis of profile single crystals
- 1999 Suppression of X-Ray linear absorption coefficient.
- 2000 Synthesis of new composite porous materials for X-ray detectors
- 2007 X-Ray transparence medium
- 2009 X-Ray acoustic monochromator
- 2010 New X-Ray diffractometer

# Full Pumping of X-ray radiation from transition direction to diffraction direction

**Experimental setup** 

**Experimental results** 



#### Focusing and defocusing of X-ray beams by means of acoustic fields



a) focusing b) defocusing.



What we have obtained during 2012?

Monochromatization of neutron beams Thermal neutrons detector based on new synthesis composite porous materials Formation of periodic structures

# Monochromatization of Diffracted Neutrons by the Acoustic Superlattice

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## Experimental setup



### **Experimental and Theoretical Results**

The integral intensity of thermal neutrons reflection in the presence of acoustic waves.

$$I_h = \frac{I_0}{2} \left( 1 - \frac{\alpha^2 - U_0^2}{\sqrt{(1 + (\alpha + U_0)^2)(1 + (\alpha - U_0)^2)}} \right)$$

where  $\alpha$  –is an immeasurable value and characterizes the deviation from Bragg angle. *Uo* –is the amplitudes of acoustic wave oscillations



# The Coherence of Thermal Neutrons for Full Pumping





#### **High Speed Detector of Thermal Neutrons**

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Figure1 schematic view of the detector based on porous composite materials (2) placed in the special vacuum chamber of stainless steel (5) with a window (1). (3) and (4) electrode contacts for applying electric field .



Figure.2 Registration efficiency  $\eta_{\alpha}$  dependence on the operating voltage U.





Figure.4 Dependence of thermal neutron absorption efficiency with energy  $E_n = 0.025 \text{ eV}$  from the thickness of LiF and LiJ.

Count per Sec



Figure 5. Registration rate dependence from the radiator voltage,  $U_{r_1}$  for the energy  $E_n = 0,025$  eV.

# FORMATION OF PERIODIC STRUCTURES

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#### **The Generator of Micro Particles**



Fig.1 The general view of the generator with the crystallization chamber.

# **Crystallization Chamber**



For micro crystals acquisition, different solutions of watersoluble crystals (NaCl,  $LilO_3$ , KDP, etc.) are used. The crystallization of generated droplets is carried out by their evaporation in the crystallization chamber (fig. 2) shaped as a glass tube with temperature and humidity sensors for crystallization conditions control inside the tube. Thus obtained crystals Fig. 2 subside on a glass substrate.



Figure 3





Figure 4

On fig. 3,4 the micro photoes of the crystals are demonstrated. As it is senn on the photoes, the particles have the same size and form. The measurements show that the average quadratic deviation of particles equals about 5%. On the photoes, it is seen the forms of crystal cells (cubic). Such compact packing is the result of particle self-organization in a viscus medium under external influences (electrical, acoustical fields, temperature gradient, etc.).

While laser illumination of such structures, diffraction patterns are seen on the screen (fig. 5a, b).





Fig.5.a. Particle average size is  $3,4\mu m$ .

Fig.5.b. Particle average size is 5,5  $\mu$ m..



In fig.6.a and 6.b they are presented the photos of micro crystals and their corresponding diffraction patterns for two materials: NaCl and LiIO<sub>3</sub> (for obtaining better photos, it is cut a hole near the central bright maximum in the screen). In the photos of micro crystals, the areas with different distribution of the particles are indicated with circles. On the right, they are presented the diffraction for the both materials. As it is seen in the photos, with more numbers of the particles in the illumination zone, they are more diffraction rings on the screen. At the same time, the diameters of the corresponding rings don't change.

In the presented photos the measurement precision is about 10 % (the size of the particles are  $2,6\pm0,2\mu m$  for NaCl and  $2,3\pm0,2\mu m$  for LiIO<sub>3</sub>).

#### **Development of Periodical Microstructures**

For ordered distribution of the particles on the substrate, masks with different sizes and periodical holes have been used (fig. 9).



Fig.9. The holes size  $40*40 \ \mu\text{m}$ , the distance between the holes  $80 \ \mu\text{m}$ 

What we are expecting in near future?

Diffraction of thermal neutrons in single Quartz crystals in the presence of acoustic fields. Thermal neutrons transparence medium Thermal neutron diffractometer Accumulating systems of X-Rays and thermal neutrons Detectors for elementary particles and gamma quanta with coordinate sensitivity better than 5µm Alternative sources of electron beams Alternative sources of energy Portable X-Ray and thermal neutron diffractometers

# Thank you for attention!