

# Emission of monochromatic sub-THz radiation by IR-active phonons in $KY(MoO_4)_2$

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## Introduction.

Double molybdates  $MRe(MoO_4)_2$  (where  $M^+$  is an alkali metal ion and  $Re^{3+}$  is a rare-earth or  $Y^{3+}$  ion) are optically transparent dielectric compounds with layered orthorhombic structure  $Pbcn$  ( $D_{2h}^{14}$ ) formed by  $[M]^+$  and  $[Re(MoO_4)_2]^-$  layers. Layered structure of  $MRe(MoO_4)_2$  leads to infrared-active shear lattice vibrations with energies below 3.7 meV, which corresponds to the frequencies below 900 GHz [1] where solid state monochromatic radiation sources are rare.

Terahertz time-domain spectroscopy (THz-TDS) is a widely used method to study optical properties of materials based on electromagnetic transients optoelectronically generated by ultrashort, usually femtosecond, laser pulses.

Using THz-TDS setup we demonstrate the re-emission of the sub-THz monochromatic radiation by coherent optical phonons in  $KY(MoO_4)_2$  pumped by a broadband THz excitation pulse.

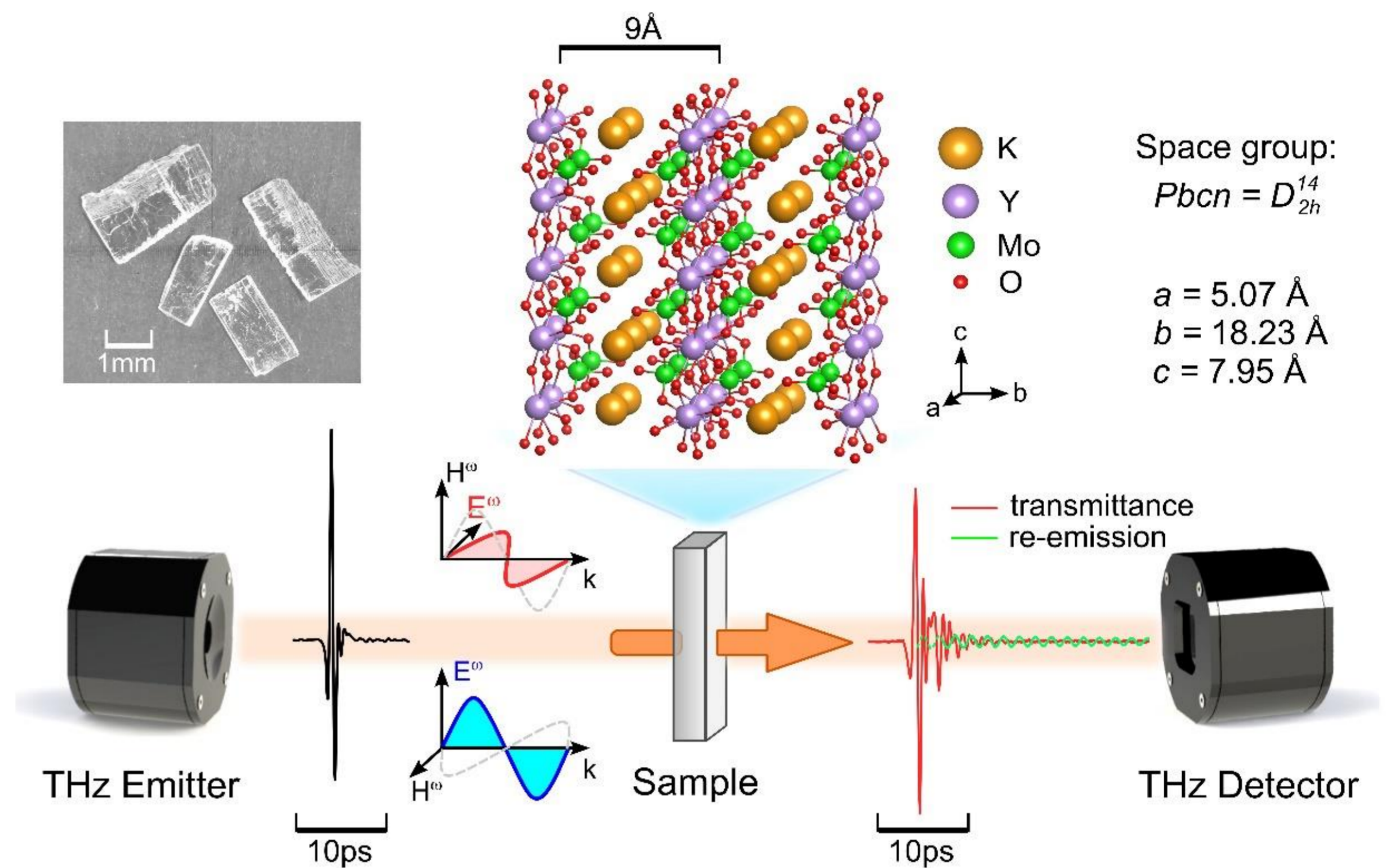


Fig.1. Sketch of the THz-TDS experiment. Black trace shows THz waveform of incident pulse. Red and green curves show the components of the beam passed the sample. At the top of the figure shown the photo of the  $KY(MoO_4)_2$  single-crystals and its crystallographic structure.

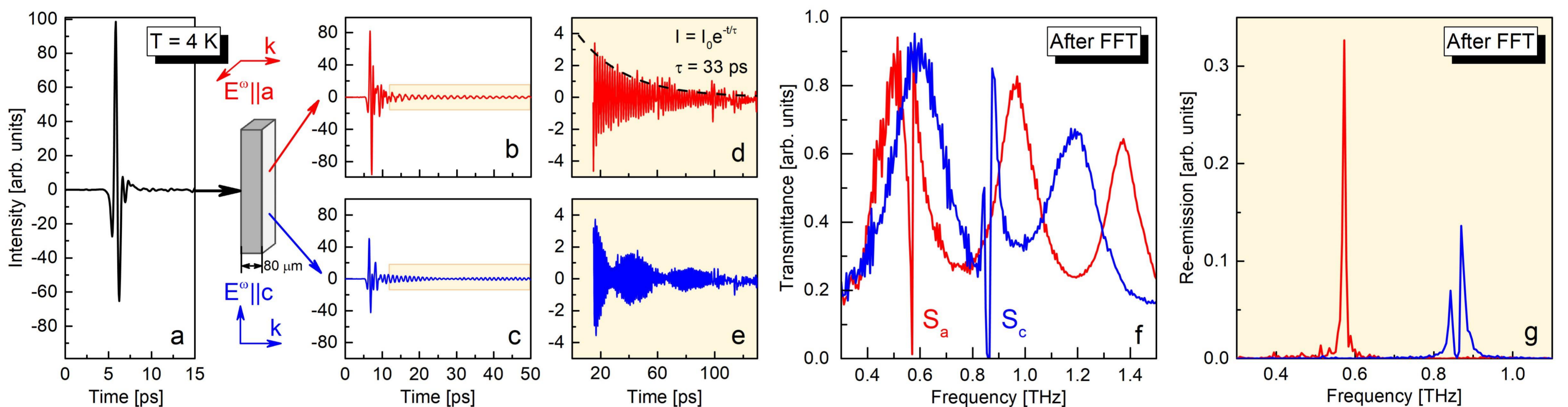


Fig.2. (a) The electric field waveform of the THz pulse before the sample. (b, c) Waveforms of the THz pulse passed the sample (for the case of 80  $\mu\text{m}$  thick sample plate and  $T = 4$  K). (d, e) Zoomed part of the waveforms in the time range 10 – 130 ps (re-emission tail). (f) The transmittance spectra (after FFT post-processing) for  $KY(MoO_4)_2$ . The periodic fringes in the spectra are caused by multiple reflections within the plane-parallel sample (Fabry-Perot type modulation).  $S_a$  and  $S_c$  – phonon modes with energies 568 and 860 GHz respectively. (g) The FFT of the waveforms shown in the Fig.1 (d, e). The spectra obtained at polarizations  $E^\omega \parallel a$  and  $E^\omega \parallel c$  denoted by red and blue colors respectively.

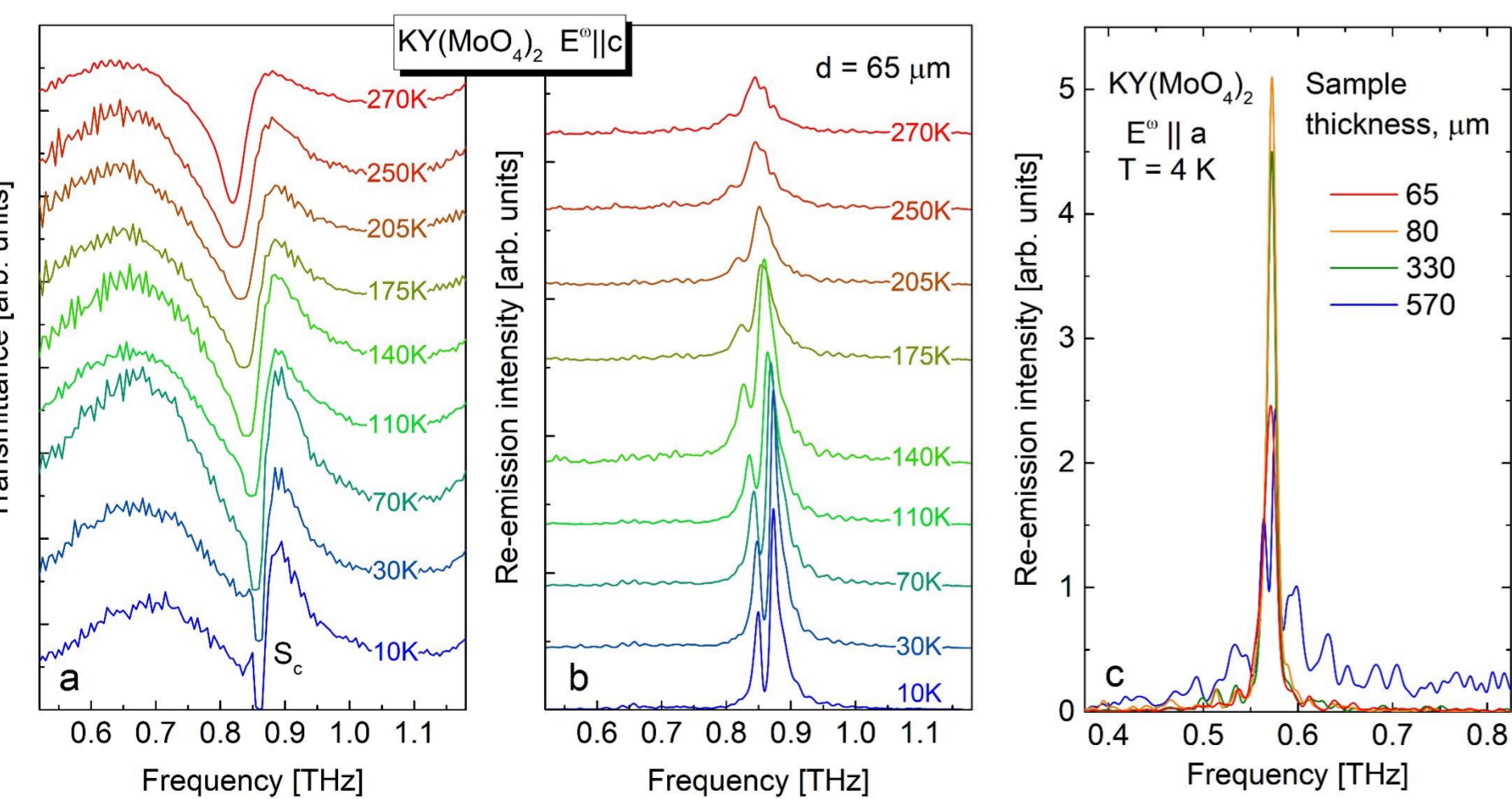


Fig.3. Temperature evolution of the transmittance (a) and re-emission (b) spectra obtained with 65  $\mu\text{m}$  thick sample for  $E^\omega \parallel c$ . The re-emission spectra were obtained in the same manner as in Fig. 2e. (c) The re-emission spectra for  $E^\omega \parallel a$  vs. sample thickness.

## Conclusions.

As the conclusions it should be note the next main results:

- Experimentally observed the re-emission of monochromatic sub-THz electromagnetic radiation by coherent IR-active optical phonons in dielectric material  $KY(MoO_4)_2$  at frequencies 568 and 860 GHz for polarizations  $E^\omega \parallel a$  and  $E^\omega \parallel c$  consequently.
- The revealed re-emission has large decay time ( $> 30\text{ps}$ ), which is exceptionally long for the oscillators with frequencies below 1 THz.
- The splitting of the re-emission peak for polarization  $E^\omega \parallel c$  caused by strong intensity of  $S_c$  phonon and prohibits the propagation of resonance frequency in material, while the lattice vibrations on frequency near by the resonance efficiently re-emit the electromagnetic radiation. Such effect confirmed by temperature evolution of the spectra when increasing the temperature leads to the disappearing of the phonon saturation (Fig.3a,b).
- The sample thickness (80  $\mu\text{m}$  for  $E^\omega \parallel a$ ) provided the maximal magnitude of the re-emission has been found (Fig.3c).
- Significantly large decay time, high intensity of re-emitted radiation and chemical stability of the material make this compound attractive for a variety of electrooptical applications.