



Experiment title: Investigations of phase transitions on Eu compounds by nuclear resonance scattering on $^{151}\text{Eu}$ applying the crossed polarizer technique	Experiment number: HE 286	
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## Report:

After the first successful nuclear resonance scattering experiment at the 21.5 keV resonance of  $^{151}\text{Eu}$  performed by us at the ESRF in October 1995 we report here on nuclear forward scattering (NFS) on Eu compounds undergoing phase transitions. Furtheron we performed the *first inelastic nuclear scattering experiment on the isotope  $^{151}\text{Eu}$* .

We used a highly monochromatized beam with a bandwidth of  $\approx 7\text{meV}$ , the time resolution of the detector system was  $\approx 1.4\text{ns}$ . The samples were  $\text{Eu}_3\text{S}_4^b$  and EuS (both polycrystalline) and a single crystal of  $\text{EuR}_3\text{O}_{14}$ , all with natural abundance of  $^{151}\text{Eu}$ .

EuS which is ferromagnetic below 16.5 K was measured at room temperature (cf. Fig. 1a) and in a liquid helium cryostat at 4.2 K without magnetic field and in a field of 3.67 T applied parallel to the synchrotron beam. The spectra at low temperatures are shown in Fig. 1b, c. The fits of these spectra prove that without external field the magnetic moments in the EuS powder are randomly oriented, while with applied field the magnetization is perfectly aligned along the synchrotron beam.

$\text{Eu}_3\text{S}_4$  exhibits two different valence states ( $\text{Eu}^{2+}$  and  $\text{Eu}^{3+}$ ), coexisting at low temperatures, leading to a pronounced modulation of the time decay of the nuclear excitation (cf. Fig. 1f). At temperatures above  $\approx 150\text{K}$  charge fluctuations set in that drastically influence the time spectra (cf. Fig. 1d, e), since the time scale of the dynamics of these processes fits excellent to the time window accessible in the NFS of synchrotron radiation on  $^{151}\text{Eu}$ . The final "mixed valent" state is reached above room temperature and was not accessible in the present experimental setup. The much faster time decay of the nuclear excitation at higher temperatures indicates the presence of fluctuations. This can also be seen by comparison of Fig. 1d with Fig. 1a where the decay of the EuS single resonance line proceeds much slower than that of  $\text{Eu}_3\text{S}_4$  at the same temperature.

<sup>a</sup> 12 shifts in hybrid mode for adjustment of the setup and 5 shifts out of 15 in 16 bunch mode (shared with two other proposals: HE 289 and HE 175).

<sup>b</sup> We are grateful to Drs. K.-U. Neumann and K.R.A. Ziebeck (Loughborough University, Great Britain) for the loan of the  $\text{Eu}_3\text{S}_4$  sample.

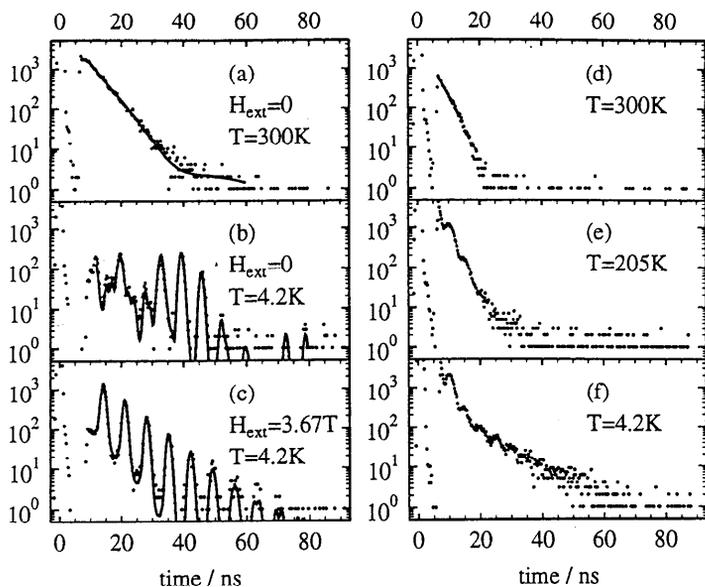


Fig. 1: NFS spectra of  $^{151}\text{Eu}$  in two Eu compounds:

- (a) paramagnetic EuS at room temperature; (b), (c) the ferromagnetic EuS at  $T = 4.2\text{ K}$  (solid lines are fits adopting the dynamical theory of nuclear resonance scattering).  
 (d), (e), (f)  $\text{Eu}_3\text{S}_4$  at various temperatures as indicated.

$\text{EuP}_5\text{O}_{14}$  is of special interest in optics and high pressure studies. Our inelastic nuclear scattering measurement on a single crystal (cf. Fig. 2) is the first experiment of this type at the  $^{151}\text{Eu}$  resonance and proves the feasibility of this new technique at 21.54 keV.

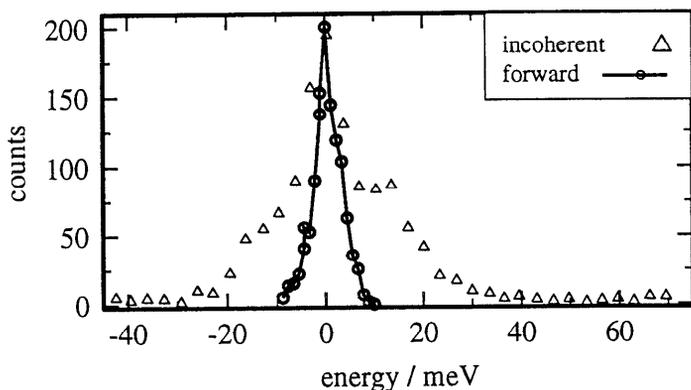


Fig. 2: Inelastic scattering on  $\text{EuP}_5\text{O}_{14}$  after excitation by synchrotron radiation.

$\Delta$  : Intensity of the incoherently reemitted radiation as a function of the energy shift of the incoming radiation relative to the resonance energy in NFS geometry.

$\circ$  : Normalized NFS intensity; the width of the peak is a measure for the energy resolution of the setup (the full line is only to guide the eye).