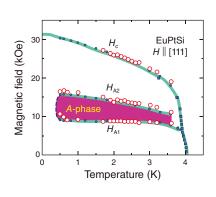
## **5–5** Capturing of a Unique Spin Texture of a 4*f* Electron

- Discovery of a Magnetic Skyrmion Lattice in the 4f Electron Magnet EuPtSi -



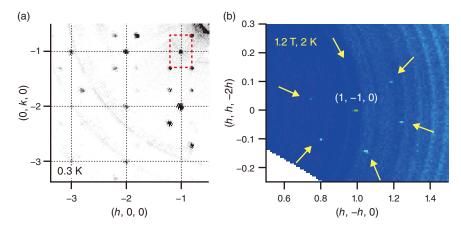


Fig.5-13 Magnetic phase diagram of EuPtSi A field-induced A-phase only exists at

a finite temperature and magnetic field.

**Fig.5-14 Neutron scattering intensity map of EuPtSi (a) at zero field, (b) 1.2 T** The rectangular magnetic scattering pattern found at zero field changes into a characteristic hexagon in the plane normal to the applied field when the system enters the *A* phase.

Magnetic skyrmions are particle-like, topologically non-trivial spin textures that have attracted interest in fundamental aspects as well as applications due to their potential as a low-powerconsumption device. A magnetic skyrmion formation was first identified in a chiral magnet MnSi. This discovery has triggered an intensive search for compounds with similar crystal structures, leading to the discovery of their existence in MnGe, FeGe, and insulator Cu<sub>2</sub>OSeO<sub>3</sub>. A variety of characteristic properties have been unveiled as the number of skyrmion compounds has increased. Unlike this diversity, magnetic ions have been mostly limited to 3*d* transition metal elements.

Recently, high-quality single crystals of the 4f electron compound EuPtSi have been grown to have a similar crystal structure. In EuPtSi, seven 4f electrons carry magnetism. This compound exhibits an antiferromagnetic transition at 4 K at a zero magnetic field. Characteristic features emerged under magnetic fields, including a field-induced phase, i.e., the A phase, which appeared only at a finite temperature and magnetic field, as shown in Fig.5-13. This feature, common to the skyrmion phase of MnSi, implies the existence of a magnetic skyrmion in EuPtSi. To clarify the nature of the A phase, single-crystal neutron diffraction experiments were carried out under a magnetic field.

Neutron diffraction is a powerful probe to determine the spin arrangement under an extreme environment, such as low temperature and a magnetic field. Experiments were performed using the Wide-Angle Neutron Diffractometer (WAND<sup>2</sup>) installed at the High Flux Isotope Reactor (HFIR) in Oak Ridge National Laboratory (ORNL), U.S.A. and SENJU located at BL18 in the Materials Life Science Facility (MLF), J-PARC.

The formation of a magnetic skyrmion lattice in MnSi was evidenced by small-angle neutron scattering via an observation of six-fold magnetic diffraction patterns in the plane normal to the applied field. In EuPtSi, the magnetic structure at a zero field was helical, characterized by a rectangular form as displayed in Fig.5-14. By applying a magnetic field, the diffraction pattern exhibited substantial change when the system enters the *A* phase; the rectangular pattern at the zero field changed into a six-fold pattern in the plane normal to the field. Thus, the pattern common to the skyrmion phase was demonstrated.

This work revealed that magnetic diffraction pattern of EuPtSi has similar characteristics to the archetypical skyrmion lattice magnet MnSi, suggesting a formation of skyrmion lattice in EuPtSi. However, marked differences were seen. The periodicity in MnSi is quite long, requiring small-angle neutron scattering to separate magnetic and nuclear peaks, whereas the standard diffraction is sufficient to reveal magnetic patterns in EuPtSi owing to its shorter periodic length: 18 versus 1.8 nm in MnSi and EuPtSi, respectively. In addition, the *A* phase in EuPtSi was stabilized over a relatively wide temperature range, whereas that of MnSi exists over only a narrow region in close proximity to the transition temperature. Overall, the discovery of a magnetic skyrmion phase in 4*f* electron systems has provided a new direction to explore another compounds and has unveiled more characteristics of skyrmions.

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## Reference

Kaneko, K. et al., Unique Helical Magnetic Order and Field-Induced Phase in Trillium Lattice Antiferromagnet EuPtSi, Journal of the Physical Society of Japan, vol.88, issue 1, 2019, p.013702-1–013702-5.