

SUPPLEMENTAL MATERIAL

Type-I superconductivity in the Dirac semimetal PdTe₂ probed by μ SR H. Leng¹, J.-C. Orain², A. Amato², Y. K. Huang¹, and A. de Visser¹

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1. Powder X-ray diffraction

A powder X-ray diffraction pattern was measured on a crushed piece of the PdTe₂ single crystal, see Fig. S1. The diffraction pattern, taken with Cu-K α radiation, confirms the CdI₂ crystal structure with lattice parameters $a = 4.034 \text{ \AA}$ and $c = 5.132 \text{ \AA}$. The pattern shows single phase homogeneity within the experimental resolution (5 %).

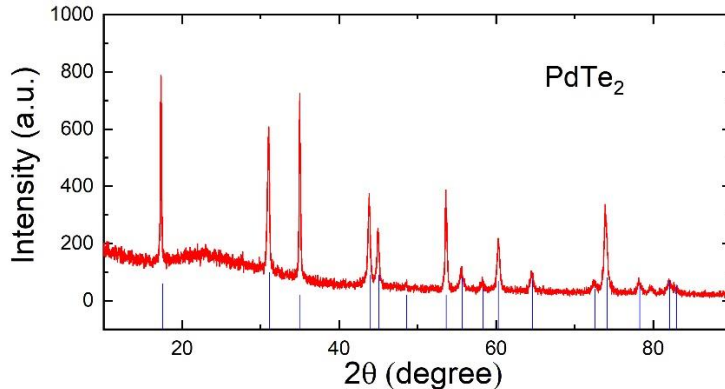


Fig. S1 Powder diffraction pattern of PdTe₂. The blue lines indicate the expected Bragg positions.

2. Energy dispersive X-ray spectroscopy

Scanning electron microscopy (SEM; Hitachi TM3000) and energy dispersive X-ray spectroscopy (EDX; Quantax 70) were carried out on the surface of the PdTe₂ crystal. A typical EDX spectrum is shown in Fig. S2. Surface scans indicate a homogenous distribution of Pd and Te. EDX spectra measured at several positions on the surface show the expected stoichiometry of 1:2 within the experimental resolution of 0.5 %.

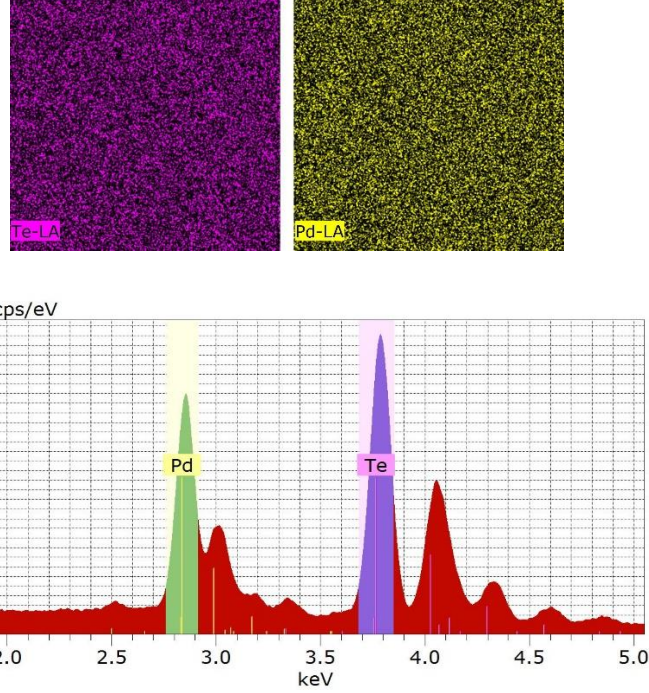


Fig. S2 EDX analysis of the surface of PdTe_2 . The surface scans (upper panels, scanned area $360\mu\text{m} \times 385\mu\text{m}$.) show the distribution of Pd and Te is homogeneous. The EDX spectrum (lower panel) gives a Pd : Te ratio 33.2 : 66.8.

3. Estimate of the demagnetization factors N_{\perp} and N_{\parallel}

The single crystal used for the μSR measurements had a disk-like shape with the c -axis perpendicular to the plane of the disk. Its thickness and diameter were 0.65 mm and 10 mm, respectively. A small piece was removed from the disk along the a -axis for other measurements. This reduced the size in the perpendicular a^* -direction to 6.8 mm and hampered the precise calculation of N . For the configuration with the field applied along the c -axis (N_{\perp}) we made two approximations. Assuming the crystal is a short cylinder with diameter 10 mm we calculate $N_{\perp} = 0.88$ (Ref. 1). Assuming the crystal is a rectangular bar with size $10 \times 6.8 \times 0.65\text{ mm}^3$ we calculate $N_{\perp} = 0.86$ (Ref. 2). In the manuscript we use the average value $N_{\perp} = 0.87 \pm 0.02$, where the error is partly due to the uncertainty in the dimensions of the sample. Under the assumption that $N = N_{\perp} + 2N_{\parallel} = 1$, we calculate for the field in the plane of the disk $N_{\parallel} = 0.07$. On the other hand, the calculated value of the radial demagnetization factor of the short cylinder is $N_{\parallel} = 0.09$ (Ref. 3). Again, in the manuscript we use the average value $N_{\parallel} = 0.08 \pm 0.02$. We remark these values are obtained for a complete diamagnetic screening, $\chi = -1$.

4. Temperature variation of the μ SR damping rate

The field and temperature variations of the different relaxation rates derived from fitting the μ SR spectra to Eqs. 1-3 are shown in Fig. S3. In the left diagram $\sigma(H)$ is traced for the configuration with the applied field perpendicular to the plane of the disk and in the right panel $\sigma(T)$ for the field in the plane of the disk. Note the volume fraction of the background contribution is always relatively small, $\sim 10\%$.

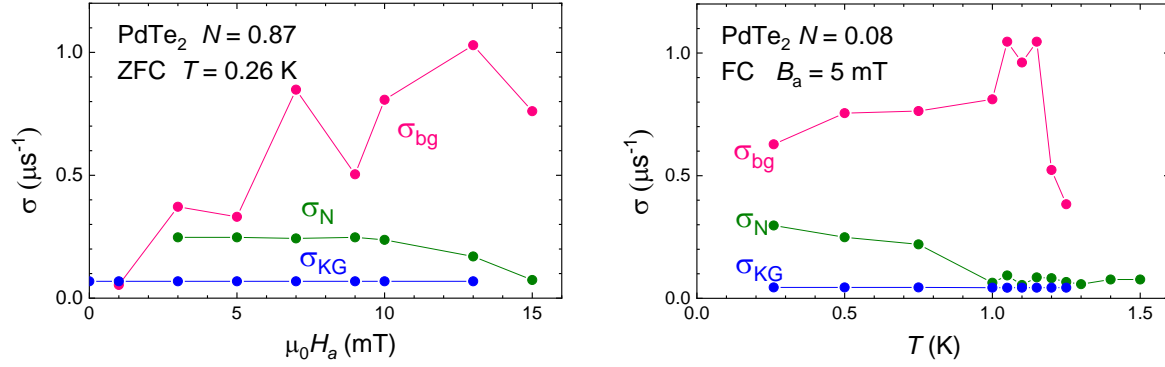


Fig. S3 Field (left panel) and temperature (right panel) variation of the μ SR relaxation rate for the Meissner phase, σ_{KG} , the intermediate or normal phase, σ_{N} , and the background contribution, σ_{bg} .

References

1. D.-X. Chen, J.A. Brug and R.B. Goldfarb, IEEE Trans. Magn. **27**, 3601 (1991).
2. E. Pardo, D.-X. Chen and A. Sanchez, J. Appl. Phys. **96**, 5365 (2004).
3. D.-X. Chen, E. Pardo and A. Sanchez, IEEE Trans. Magn. **37**, 3877 (2001).