Orbital-exchange and fractional quantum number excitations in an f-electron metal, Yb2Pt2Pb


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Orbital-exchange and fractional quantum number excitations in an f-electron metal, Yb$_2$Pt$_2$Pb


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Materials and Methods

Methods summary

Inelastic neutron scattering measurements on Yb$_2$Pt$_2$Pb were carried out using the Disk Chopper Spectrometer (DCS) [31] at the Center for Neutron Research at the National Institute of Standards and Technology and the Cold Neutron Chopper Spectrometer (CNCS) [32] at the Spallation Neutron Source at Oak Ridge National Laboratory. A 6 g sample of $\approx$ 400 co-aligned Yb$_2$Pt$_2$Pb crystals was used, oriented with (1-10) crystal lattice direction vertical and the $(Q_{HH}, Q_{HH}, Q_L)$ reciprocal lattice plane horizontal. On DCS the sample was mounted in a 10 T vertical field superconducting magnet equipped with the dilution refrigerator, on CNCS a 5T vertical field magnet with 1.5K base temperature was used. The incident neutron energy was $E_i = 3.27$ meV ($\lambda = 5.0$ Å) on DCS and $E_i = 3.316$ meV ($\lambda = 4.97$ Å) on CNCS. For nominal zero field measurements at base temperature 0.07 K a small bias field of about 250 Oe was used to suppress the superconductivity of the aluminum sample holder. A 4 T field along the (1-10) direction fully polarizes the sublattice with magnetic moments parallel to (1-10), allowing us to use measurements at 4 T as a background to isolate the inelastic scattering from the orthogonal sublattice with moments along (110), which is unaffected by the field. The measured intensities were corrected for neutron absorption, and the scattering cross section was put on an absolute scale by comparing the measured Bragg peak intensities with calculated nuclear structure factors. All wave vectors, $(Q_{HH}, Q_{HH}, Q_L)$, are indexed in reciprocal lattice units of $a^* = 2\pi/\alpha = 0.81$ Å$^{-1}$ and $c^* = 2\pi/c = 0.91$ Å$^{-1}$. The susceptibility of XXZ model was calculated using the quantum transfer matrix approach suitable for finite temperature calculations [33, 34] and solving the corresponding non-linear integral equations [35].

Sample preparation

Yb$_2$Pt$_2$Pb single crystals were grown from a Pb flux as described in Ref. 5. The typical crystals have rod-like shape with typical dimensions $\sim 0.5$ mm in width and $\sim 4 - 5$ mm in length, and an approximately square cross-section with flat surfaces normal to the (110) crystal direction and with the (001) lattice $c$—axis along the length. Crystal orientations were checked with X-ray diffraction. The $\sim 6$ g single crystal sample used for inelastic neutron scattering was assembled by co-aligning, using the crystal morphology, between 300 and 400 crystals on 6 aluminum plates. Crystals were held in place by the surface tension from hydrogen free Fomblin oil. These plates were clamped together in an Al box, resulting in the overall crystal mosaic spread perpendicular to the (001) direction of $\sim 2^\circ$. A picture of the sample holder is shown in Fig. S1A. Neutron powder diffraction measurements were performed on a 5 g powder sample prepared from similar single crystals.

Magnetization measurements

For $T = 1.5$ K and $T > 1.8$ K, the bulk magnetization and susceptibility were measured on a single crystal sample using a [Quantum Design] Magnetic Property Measurement System SQUID magnetometer with a gas-flow cryostat. For $T < 1.8$ K,
magnetization and susceptibility were measured on a single crystal using a Hall sensor magnetometer, mounted in the dilution refrigerator insert of the [Quantum Design] Physical Property Measurement System.

Neutron Scattering setups

Figure S1: (A) Single crystal sample holder used for neutron scattering experiments at DCS at NCNR, NIST, and CNCS, SNS. Single crystals of Yb$_2$Pt$_2$Pb were aligned with the (110) crystal direction normal to the surface of the holder plate, and with the (1 − 10) direction parallel to the long side of the rectangular holder. (B) Schematic view of the configuration for both DCS and CNCS neutron scattering setups. Magnetic field B is applied along the vertical, (1 − 10) direction. The sample rotates in the horizontal, (H H L) plane, with access to the vertical (H -H L) scattering plane limited by the acceptance
The distance $D$ from sample to detector and the detector height, $H$, for both DCS and CNCS experiments are described in the text. (C), (D) Contour maps of elastic neutron scattering intensity integrated within the energy range $E = [-0.2,0.2]$ meV, measured in zero field on DCS (C) and CNCS (D). For the DCS data, the detectors are only position sensitive in the horizontal direction, and all scattering in the vertical, $(H-H)$ direction is projected onto the $(H H L)$ scattering plane, (C). For the CNCS data, the detectors have spatial resolution in both horizontal and vertical directions. The data in (D) are integrated only within a narrow region, $Q_{H-H} = [-0.05,0.05]$, in the vertical direction. The diffraction seen in the DCS data (C) at $Q = (001)$ is picked up by the relaxed vertical resolution of that measurement and is absent in the CNCS data (D), as indicated by red arrows.

Most of the data shown in the main manuscript were acquired using the Disk Chopper Spectrometer (DCS) at the National Institute for Standards and Technology (NIST) Center for Neutron Research (NCNR) [31, 36]. Additional measurements of the temperature dependence were performed using the Cold Neutron Chopper Spectrometer (CNCS) at the Spallation Neutron Source at Oak Ridge National Laboratory [32, 37]. The complementary neutron powder diffraction measurements quantifying the order parameter and its detailed temperature dependence were made on the BT-7 double focusing triple-axis spectrometer at NIST NCNR [38] and are reported elsewhere [18].

For the single crystal measurements on DCS and CNCS, the sample holder with the sample was mounted in the $Q_{HH} - Q_L$ scattering plane, with the (1-10) direction vertical. For the DCS experiment, a 10 T cryomagnet with a dilution refrigerator insert with base temperature of 0.1 K was used. A $1.5 \text{ cm} \times 2.5 \text{ cm}$ beam mask was installed in front of the sample to reduce the unwanted background scattering from the sample environment. A small bias field of 250 Oe was used for the nominal zero field measurements at base temperature in order to suppress the superconductivity of the aluminum sample holder. The magnet dark angle was set at $75^\circ$ from the (110) direction, eliminating a substantial source of background scattering. The choppers were set to select neutrons with fixed incident energy of $E_i = 3.27$ meV ($\lambda_i = 5.0$ Å). The vertical acceptance of the detector system to scattered neutrons is limited by the height of the magnet window. Only neutrons scattered from the sample within angle $\pm \theta$ and illuminating a segment of height $H$ on the detector are counted, as illustrated in Fig. S1B. On DCS the distance from sample to detector is $D = 4010$ mm, and only the data of the central detector bank with tubes of height $H = 400$ mm [36] were analyzed, thus resulting in the vertical acceptance of $\pm \theta_{DCS} \approx \pm 5.7^\circ$ in our experiment. As a result, although the scattering plane is (HHL), part of the scattering from the (H-HL) plane within this angular acceptance also contributes to the overall scattering intensity. The corresponding vertical wave vector resolution full width at half maximum (FWHM) is $\approx 0.25$ Å, or about 0.2 reciprocal lattice unit of $a_{110}^* = a^*\sqrt{2} \approx 1.146\hat{\xi}$.

For the CNCS experiment, a 5 T cryomagnet with a He flow cryostat with $T=1.5$ K base temperature was used. The horizontal angular acceptance of the magnet was large and was not significantly restricted by the magnet dark angle. The incident neutron
energy was fixed at 3.316 meV ($\lambda_i = 4.97$ Å), and the high-flux mode of the instrument was used to maximize neutron intensity. The distance from sample to detector was $D = 3500$ mm. Although each 128-pixel detector tube has height $H = 2000$ mm [32, 37], the vertical acceptance of the magnet constrained us to only use the middle 88 pixels in the vertical direction. This gives an effective height of the detector in this experiment of $H = 2000 \times 88/128 = 1375$ mm, and an angular acceptance $\pm \theta_{\text{CNCS}} \approx \pm 11^\circ$, nearly twice the vertical acceptance at DCS. This somewhat increased vertical acceptance angle makes difference for the inelastic neutron scattering measurements by virtue of increasing the overall scattering intensity, but it is of even greater importance for the elastic diffraction measurements. Here, Bragg reflections that are close to, but not exactly in the horizontal scattering plane, may appear unexpectedly if one simply adds all scattered intensity along the vertical dimension of the detectors, thus yielding a misleading result. Fortunately, the CNCS detectors are position sensitive along the vertical direction [32, 37], which provides an access to the out of plane, (H-HL) direction and allows isolating contributions from the (HHL) scattering plane.

By comparing the neutron intensity measured on the DCS with that measured on the CNCS within different vertical wave vector ranges, we conclude that the DCS measurements, which are not position sensitive in the vertical direction, correspond to summing up the intensity over the range of $Q_{H-H} = [-0.1, 0.1]$, in agreement with the FWHM of the vertical resolution of the DCS measurements estimated above. Contour maps of the elastic ($E = [-0.2, 0.2]$ meV) neutron scattering intensity measured at zero field on DCS and CNCS are shown in Fig. S1C and Fig. S1D, respectively. For the DCS data all scattering within the vertical acceptance is projected onto the (HHL) scattering plane. Thus, a diffraction peak is seen at $Q = (001)$ in Fig. S1C, which is actually tails of the satellite magnetic Bragg peaks in the vertical, (H-HL) plane, above and below the (HHL) scattering plane, which are picked up by the vertical resolution of this measurement. For the CNCS data we can limit the integration in the vertical direction to a narrow range, $Q_{H-H} = [-0.05, 0.05]$, thus avoiding the out-of-plane (H-HL) contributions. Consequently, the scattering at $Q = (001)$ is absent, as indicated by the red arrow in Fig. S1D. This will be further discussed later, with reference to Fig. S8A, B.

For powder neutron diffraction measurements on BT-7, a 5 g powder sample was pressed in an Al sample can and mounted in a $^{3}$He cryostat. Neutrons of wavelength $\lambda_i = 2.359$ Å ($E_i = 14.7$ meV) were incident on the sample, and scattered neutrons were measured using the instrument’s position sensitive detector.

**Absolute normalization and extinction effect**

Since our neutron scattering experiments were carried out on two different time of flight spectrometers, DCS and CNCS, in order to compare and combine these two data sets we express both in absolute units. In principle, we could carry out this normalization in several ways: by using the incoherent scattering of the sample, the incoherent scattering of a vanadium standard, or using the structure factor of the diffraction peaks of
Yb$_2$Pt$_2$Pb. The incoherent scattering of Pt and Pb are very small, and the incoherent scattering of the Yb$_2$Pt$_2$Pb sample itself is comparable to scattering from the sample environment, ruling out this approach. Since Yb and Pt have relatively large absorption cross sections $\sigma_{abs}$, combined with the fact that we have a relatively large sample, normalization using a vanadium standard is also problematic, due to this absorption effect. Consequently, we use the structure factors of nuclear and magnetic diffraction peaks in Yb$_2$Pt$_2$Pb to place the experimental intensities on an absolute scale.

<table>
<thead>
<tr>
<th>Nuclear Diffraction</th>
<th>(002)</th>
<th>(110)</th>
</tr>
</thead>
<tbody>
<tr>
<td>$</td>
<td>F_{hkkl}</td>
<td>^2$ (barn-fu$^{-1}$)</td>
</tr>
<tr>
<td>$\int d^3Q \int dEI(Q,E)$ (meV)</td>
<td>0.00165</td>
<td>0.00281</td>
</tr>
</tbody>
</table>

Table S1: The calculated nuclear structure factors and the experimentally measured integrated intensities for the primary diffraction peaks accessed in the (HHL) scattering plane in the CNCS single crystal neutron scattering measurements at 1.5 K in zero field.

<table>
<thead>
<tr>
<th>AF Diffraction</th>
<th>(1.2, 1.2, 0)</th>
<th>(1.8, 1.8, 0)</th>
<th>(0.2, 0.2, 1)</th>
<th>(0.8, 0.8, 1)</th>
</tr>
</thead>
<tbody>
<tr>
<td>$</td>
<td>Q_{hkkl}</td>
<td>$ (Å$^{-1}$)</td>
<td>1.373</td>
<td>2.06</td>
</tr>
<tr>
<td>$</td>
<td>F(Q)</td>
<td>^2$</td>
<td>0.924</td>
<td>0.84</td>
</tr>
<tr>
<td>$</td>
<td>F_{hkkl}</td>
<td>^2$ (barn-fu$^{-1}$)</td>
<td>0.0636</td>
<td>0.1187</td>
</tr>
<tr>
<td>$\int d^3Q \int dEI(Q,E)$ (meV)</td>
<td>0.0001219</td>
<td>0.00023148</td>
<td>0.001805</td>
<td>0.002754</td>
</tr>
</tbody>
</table>

Table S2: List of the calculated antiferromagnetic structure factors, $|F_{hkkl}|^2$, and the experimentally measured integrated intensities for magnetic diffraction peaks accessed in the (HHL) scattering plane in the CNCS single crystal neutron measurements at 1.5 K in zero field ($|F(Q)|^2$ is magnetic form factor [39] squared). The ordered moment at 1.5 K is $M_{order} = 2.5 \mu_B / Yb$, which was extracted from the order parameter measurement shown in Fig. S2A.

Using the known crystal structure [5], we have calculated the nuclear structure factors for the diffraction peaks accessed in the (HHL) scattering plane, two of which are listed in Table S1. Magnetic structure factors were calculated based on the powder diffraction experiments on BT-7. Measurements for scattering angles $15^\circ \leq 2\theta \leq 90^\circ$ were performed at different fixed temperatures ($T = 10$ K, 1.5 K, and 0.5 K) above and
below the ordering temperature $T_N = 2.07 \, \text{K}$, and the zero field antiferromagnetic (AF) magnetic structure was solved [18]. Scattered intensity was measured as a function of temperature at $2\theta = 20^\circ$, tracking the AF order parameter at $Q = (0.2, 0.2, 1)$ from 0.48 K up to 3 K, as shown in Fig. S2A. The magnetic transition is clearly seen at $T \approx 2.1 \, \text{K}$, the Néel temperature previously identified in specific heat and magnetic susceptibility measurements [5, 6]. Based on the solution of the magnetic structure [18], and the fitted moment values from the $2\theta$ scans at fixed temperatures, it was possible to extract the ordered moment, $M_{\text{order}} = 2.5 \mu_B/Yb$ at 1.5 K, as indicated by the red solid diamond in Fig. S2A. The calculated magnetic structure factors for diffraction peaks in the (HHL) plane are listed in Table S2.

Figure S2: **Magnetic and nuclear Bragg intensities in Yb$_2$Pt$_2$Pb in zero field.** (A) Ordered moment squared as a function of temperature. Black solid circles are the temperature scan at the (0.2, 0.2, 1) AF peak position at $2\theta = 20^\circ$ [18], which was put on the absolute scale using the ordered moment $M_{\text{order}} = 2.5 \mu_B/Yb$ (red solid diamond) extracted from the refinement of the powder diffraction pattern measured in a $2\theta$ scan at 1.5 K. (B) Comparison of the calculated nuclear and magnetic structure factors and the experimental integrated peak intensities for peaks accessed at 1.5 K at CNCS and listed in Tables S1 and S2. The slope in the limit of small intensity gives the normalization factor for the experimental CNCS data to absolute values, $1/C = \int d^3Q \int dE(Q, E) / |F_{hkl}|^2$ (solid red line). The dashed line is a fit to the semi-empirical secondary extinction factor for the single crystal diffraction intensity, as explained in the text.

By comparing the calculated nuclear and magnetic structure factors, $|F_{hkl}|^2$, with the experimentally measured integrated intensities, $\int d^3Q \int dE(Q, E)$, the normalization factor $C$ can be obtained as a ratio,
However, if we plot the experimentally measured integrated intensity as a function of the calculated structure factor [Fig. S2B], the result does not fall on a straight line. This is due to the secondary extinction effect, which is usually significant for single crystal diffraction experiments on large mosaic crystals. To minimize this extinction effect, we selected the two weakest magnetic diffraction peaks, (1.2, 1.2, 0) and (1.8, 1.8, 0), to extract the normalization factor $C$. As indicated by the red solid line in Fig. S2B, the inverse of the slope gives the normalization factor, $C = 515 \text{ barn} \cdot \text{meV}^{-1}$. On the other hand, an approximate account for the secondary extinction in neutron diffraction can be accomplished by using the following extinction-corrected expression for the measured integrated Bragg intensity as a function of the calculated structure factor,

$$
\int d^3Q \int dE I(Q,E) = C \frac{|F_{hkl}|^2}{1 + a|F_{hkl}|^2}
$$

(S2)

where $\alpha$ is determined by a combination of the sample mosaic and its effective thickness. The dashed line shows fit of our data to Eq. (32) with $C = 515 \text{ barn} \cdot \text{meV}^{-1}$ and $\alpha = 0.44$. The fit provides a good description of the data, and we will use this normalization factor $C$ to express the experimental scattering intensity in absolute units of $\text{barn} \cdot \text{meV}^{-1}$.

In discussing magnetic neutron scattering, it is illuminating to make a direct comparison of the fluctuating magnetic moments with static moments determined from magnetization measurements, so it is convenient to convert the measured cross-section to units of $\mu_B^2 \cdot \text{meV}^{-1}$. The cross-section of (unpolarized) neutron magnetic scattering, which is proportional to the measured intensity, $I(Q,E)$, can be expressed as,

$$
\frac{d^2\sigma(Q,E)}{dEd\Omega} = \frac{k_f}{k_i} \left(\frac{\gamma r_0}{2\mu_B}\right)^2 e^{-2W} \sum_{\alpha} (1 - \tilde{Q}_\alpha^2) M^{\alpha\alpha}(Q,E) = \frac{k_f}{k_i} \cdot C \cdot I(Q,E).
$$

(S3)

[40], where $k_f$ and $k_i$ are the scattered and the incident neutron wave vectors, $r_0 = 2.818 \times 10^{-13} \text{ cm}$ is the classical electron radius, $\gamma = 1.913$ is the magnetic moment of the neutron in nuclear magnetons, $(\gamma r_0)^2 = 2.911 \times 10^{-24} \text{ cm}^2 = 0.291 \text{ barn}$ [11], $\alpha$ denotes the Cartesian coordinate, $x$, $y$, or $z$, which characterizes the polarization of magnetic fluctuations, and $\tilde{Q}_\alpha = Q_\alpha/Q$ are the projections of the unit vector along the wave vector transfer direction on the Cartesian axes. $e^{-2W}$ is the Debye-Waller factor, which at low temperatures and small wave vectors of our measurements is taken to be unity. $M^{\alpha\alpha}(Q,E)$ is the correlation function of magnetic moments, which is related to the spin-spin correlation function, $S^{\alpha\alpha}(Q,E)$, computed for the effective spin-1/2 model through,

$$
M^{\alpha\alpha}(Q,E) = (g_\alpha^{\text{eff}})^2 \mu_B^2 |F(Q)|^2 S^{\alpha\alpha}(Q,E).
$$

(S4)

where $g_\alpha^{\text{eff}}$ is the effective $g$ —factor, and $|F(Q)|$ is the magnetic form factor for Yb $^{3+}$ local moments [39]. Using Eqs. (S2)–(S4), we can express both the measured scattering
intensity, $I(Q,E)$, and the calculated spin-spin correlation function, $S^{aa}(Q,E)$, in units of $\mu_B^2 \cdot meV^{-1}$ [41].

The absolute cross section is based on the structure factor calculated for a crystal unit cell with eight Yb atoms. Thus, in the absence of magnetic field, the scattering must be normalized by a factor of eight on account of 8 Yb sites per unit cell (and also keeping track of the different sublattice polarizations) in order to express it in magnetic units of $\mu_B^2 \cdot meV^{-1}$/Yb. As discussed above, strong Ising anisotropy divides Yb atoms into two orthogonal sublattices, with Ising axes along (110) and (-110) directions, respectively. Magnetic field of 4 T along (110), or (-110), polarizes moments in one of these two sublattices, so that only a single sublattice with four Yb moments per unit cell contributes to the measured magnetic scattering at high fields. Hence, in the 4 T data set, where one sublattice is completely polarized, the remaining inelastic scattering is from the orthogonal 4-Yb sublattice only, and must instead be normalized by a factor of four.

**Absorption Correction**

As mentioned above, Yb and Pt have relatively large absorption cross sections (Table S3), and therefore account for neutron absorption in the sample is important for consistent comparison of the DCS and the CNCS data and for the absolute intensity normalization. Here we explain how absorption has been included in the normalization factor $C$. We estimate that the absorption correction dominates the systematic error, and that the comparisons of the DCS and CNCS data sets are accurate within $\approx 10\%$.

<table>
<thead>
<tr>
<th></th>
<th>$\sigma_{coh}$ (barn)</th>
<th>$\sigma_{inc}$ (barn)</th>
<th>$\sigma_{abs}$ (barn)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Yb</td>
<td>19.42</td>
<td>4</td>
<td>34.8</td>
</tr>
<tr>
<td>Pt</td>
<td>11.58</td>
<td>0.13</td>
<td>10.3</td>
</tr>
<tr>
<td>Pb</td>
<td>11.115</td>
<td>0.003</td>
<td>0.171</td>
</tr>
</tbody>
</table>

Table S3: Neutron scattering and absorption cross sections for Yb, Pt and Pb [42].

Although the dimensions of the aluminum sample holder are $\approx 1.5$ cm wide, 2.5 cm high, and 0.8 cm thick, the overall effective shape of the horizontal cross-section of our multi-crystal Yb-Pt-Pb sample with six co-aligned layers of single crystals can be approximated by a rectangle of width $\approx 0.8$ cm along the crystal $c$ direction and thickness $\approx 0.3$ cm along the crystal (110) direction, which is perpendicular to the sample holder surface.
Figure S3: Effect of absorption in Yb$_2$Pt$_2$Pb. (A), (B) Inelastic neutron intensity in the (HHL) scattering plane integrated within the energy range $E = [0.15, 1.5]$ meV measured at zero field, with the 4 T high field data subtracted, without the absorption correction (A) and with the absorption correction (B). The white dashed lines in (A) indicate the region where the measured intensity has largest suppression due to absorption, which is corrected in (B). The overall intensity is also enhanced upon accounting for absorption in (B). (C), (D) Comparison of the calculated nuclear and magnetic structure factors and the experimental integrated diffraction peak intensities at 0.1 K measured on DCS without (C) and with (D) the absorption correction. The solid and dashed lines are as in Fig. S2, and are explained in the text. (E), (F) Normalized inelastic scattering in the (HHL) scattering plane integrated within the energy range $E = [0.15, 1.5]$ meV measured in 0 T, with the 4 T high field data subtracted, without the absorption correction (E) and with the absorption correction (F). (G), (H) Energy dependence of the scattering averaged over $Q_{HH} = [0, 1]$ and $Q_{00L} = [0, 2]$, as indicated by the dashed rectangles in (E) and (F). The red lines in (C) and (D) are the energy integrated fluctuating moments without the absorption correction (G) and with the absorption correction (H).

For the DCS data measured at 0.1 K, the absorption correction has been carried out using the absorption option in the DCS Mslice program from the DAVE analysis software package [43]. In order to understand the magnitude of systematic errors associated with the absorption, we compare the data with and without the absorption correction in Fig. S3. The absorption effect is evident in the inelastic scattering data of Figure S3A, which shows the raw measured inelastic neutron intensity in the (HHL) scattering plane summed over the energy range $E = [0.15, 1.5]$ meV, without the absorption correction. The white dashed arcs in Fig. S3A indicate the region of suppressed intensity where the absorption is maximum. It occurs when neutron beam is parallel to the crystal $c$-axis, the long dimension of the crystal. This decrease in the intensity is absent once the absorption correction is applied, and the overall intensity is also enhanced [Fig. S3B]. The normalization procedure accounting for the extinction correction described earlier has been performed for the DCS data both without [Fig. S3C] and with [Fig. S3D] the absorption correction. The dashed line in Fig. S3C shows fit of the uncorrected data using Eq. (S2), which yields the normalization factor $C_{no\text{-}absorpt} = 0.926 \text{ barn} \cdot \text{meV}^{-1}$ (solid line) and the extinction coefficient $\alpha = 0.143$. Similar fit of the absorption corrected DCS data [dashed line in Fig. S3D] yields the normalization factor $C_{absorpt} = 0.42 \text{ barn} \cdot \text{meV}^{-1}$ (solid line) and $\alpha = 0.083$. Fig. S3, E and H show the final normalized inelastic scattering cross-section in the (HHL) scattering plane summed within the energy range $E = [0.15, 1.5]$ meV, normalized without and with the absorption correction, respectively. The total scattering intensity averaged over the first Brillouin zone, within $Q_{HH} = [0, 1]$ and $Q_{00L} = [0, 2]$, for data without and with the absorption corrections is plotted in Fig. S3G and Fig. S3H, respectively. As we can see, the final integrated intensity is $6.5 \mu_B^2 \cdot \text{meV}^{-1}/\text{Yb}$ for data without the absorption correction, and $7.5 \mu_B^2 \cdot \text{meV}^{-1}/\text{Yb}$ for data with the absorption correction, with the mean square deviation of only $\approx 10\%$. This is because the diffraction
peaks, which we use for normalization, are subject to the absorption correction on par with the inelastic spectrum that we normalize, so that accounting for absorption changes both the measured inelastic intensities and the value of the normalization factor, $C$. Hence, the absolute values of the normalized intensity remain nearly unaffected by the absorption correction, as they are only sensitive to the difference in absorption between the elastic diffraction peaks used for normalization and the inelastic scattering intensities, and not to the absolute impact of absorption, which is much larger than this difference. This is an advantage of the approach to absolute normalization that we have employed.

Sum rule

Components of the total fluctuating Yb moment, $M_\alpha$, are related to those of the effective spin-1/2, $S^\alpha$, via the effective $g-$factor,

$$M_\alpha = g_\alpha^{eff} \mu_B S^\alpha, \quad \text{where} \quad (S^\alpha)^2 = 1/4,$$

(S5)

$\alpha = x, y, z$. The zero moment sum rule for the dynamical spin correlation function, $S^{\alpha\alpha}(Q,E)$, in general (for $S \geq 1$) governs the sum of contributions from all three spin polarizations,

$$\sum_\alpha \int \frac{d^3Q}{V^*} \int dE S^{\alpha\alpha}(Q,E) = S(S+1),$$

(S6)

where integrations are over the first Brillouin zone of volume $V^*$ and over all energies [40]. However, for $S = 1/2$, $(S^\alpha)^2 = 1/4$ is a $c-$number, and the sum rule is a combination of three independent sum rules,

$$\int \frac{d^3Q}{V^*} \int dE S^{\alpha\alpha}(Q,E) = (S^\alpha)^2 = 1/4, \quad \alpha = x, y, z,$$

(S7)

one for each polarization. This observation allows us to relate the absolute magnetic scattering intensity measured in Yb$_{2}$Pt$_{2}$Pb with that calculated for the effective spin-1/2 XXZ model Hamiltonian.

Since we have been able to express the scattering intensity in absolute units of magnetic scattering cross-section, as shown in Fig. 4A in the manuscript, it is possible to demonstrate the evolution with temperature of the ordered moment $M^2_{\text{Ordered}}$ (black circles), fluctuating moment $M^2_{\text{Fluctuating}}$ (red circles) and the total moment $M^2_{\text{Total}}$ (blue diamonds). In the paramagnetic state at $T \geq T_N$, there is no static magnetic order, and all magnetic moments are fluctuating. The fluctuating moments are determined by the total inelastic scattering averaged over the first Brillouin zone and integrated in energy, as shown for different temperatures in Fig. 4A of the manuscript. With the onset of magnetic order at $T_N=2.07$ K, magnetic diffraction peaks appear and their intensity increases with the decreasing temperature, while at the same time there is a corresponding drop in the magnitude of the fluctuating moments, although there is a considerable fluctuating moment even at 0.1 K. The total moment, $M^2_{\text{Total}} = M^2_{\text{Ordered}} + M^2_{\text{Fluctuating}}$, is at most weakly temperature dependent, and within the error bar of our neutron measurements is constant and consistent in magnitude with
the value expected for the ground state Yb doublet, $M_{1-10}^2 = 3.95^2 \mu_B^2 / Yb$, which is indicated by the thick gray line in Fig. 4(A) in the manuscript.

**Supplementary Text**

Crystal electric field splitting of the Yb $J = 7/2$ multiplet and the single ion anisotropy

Crystal structure, multiplet splitting, and two Yb sublattices

Yb single ion anisotropy is very important for understanding the magnetic properties and the one dimensional spin dynamics of Yb$_2$Pt$_2$Pb. The electronic configuration of the Yb atom is [Xe]$4f^{14}6s^2$, and therefore Yb$^{3+}$ ions in Yb$_2$Pt$_2$Pb have a single hole in the nearly filled $4f$ ($L = 3$) shell, which renders it a net total spin $S = 1/2$. The strong spin-orbit interaction, one of the strongest in the lanthanide series, $\Delta \sim 380$ meV, mandates that the total angular momentum, $J = L + S$, rather than $S$, or $L$, is a good quantum number, and that the total angular momentum of the ground state multiplet is $J = 7/2$, with the first excited multiplet, $J = 5/2$, about 1.2 eV higher in energy [44]. In the Yb$_2$Pt$_2$Pb layered crystal structure with 2 formula units and, correspondingly, two $a-b$ layers per unit cell, the Yb atoms occupy two inequivalent sites, labeled Yb1 (red) and Yb2 (blue) in Fig. S4A and Fig. S4B, which stagger along the $c-$axis. Both of the Yb sites have the same $C_{2v}$ (m2m) point group symmetry, where each Yb ion is surrounded by six nearest neighbor Pt atoms (three above and three below the Yb-Yb plane), and four nearest neighbor Pb atoms (two above and two below the Yb-Yb plane) [5]. These Pt and Pb atoms make an anisotropic local environment of static charges with markedly uniaxial anisotropy, which determines the magnetic ground state of each of the Yb ions.

The eightfold degeneracy of the $J = 7/2$ multiplet of Yb$^{3+}$ on both Yb1 and Yb2 sites is lifted by the crystal electric field (CEF), which in this point group symmetry results in a manifold of four doublet states. We have calculated the resulting CEF splitting of the $J-$multiplet based on the point charge model, using the McPhase software [45, 46, 47, 48]. Since Yb$_2$Pt$_2$Pb is a good metal, it is hard to assign an accurate point charge to the ions, and so we have simply used charges corresponding to the average valences for Pt and Pb, of Pt$^{2+}$ and Pb$^{4+}$. This assumes that the collectivized electrons are in a good Bloch wave states and are evenly spread throughout the crystal, providing a homogeneous and isotropic negative charge background that does not contribute to the anisotropic crystal electric field (the screening length is larger that the inter-atomic distances), not an unreasonable approximation for a good metal.
Figure S4: Yb crystal environment and magnetic moments in Yb$_2$Pt$_2$Pb. (A), (B) Chemical environment for Yb1 (red) and Yb2 (blue) sites in the $c = 0$ (A), and $c = 1/2$ (B) plane, respectively. The nearest Pt (gray) and Pb (black) neighbors are located below and above the corresponding $c = 0$ and $c = 1/2$ plane [5]. Red and blue arrows indicate magnetic moment directions consistent with the point charge model calculation [45, 46, 47, 48]. (C), (D) Simplified schematics of the two magnetic sublattices, with Yb moments along (-110) direction [sublattice 1, (C)], and along (110) direction [sublattice 2, (D)].

E) T=1.9K

F) T=1.9K
2, (D)], respectively. (E) Experimentally measured magnetization at T = 1.9 K, normalized to the Yb\textsubscript{2}Pt\textsubscript{2}Pb formula unit, for magnetic field B applied along the three principal crystal directions, (110) [blue], (100) [red], and (001) [black]. (F) Magnetization normalized with the account of the two sublattice model illustrated in (A)-(D), as explained in the text. Plotted in blue is $M_{110}^{\text{Nor.}}$ vs. $B_{110}^{\text{Nor.}}$ calculated with Eq. (S11a), in red $M_{110}^{\text{Nor.}}$ vs. $B_{110}^{\text{Nor.}}$ calculated with Eq. (S11b), and in black $M_{001}^{\text{Nor.}}$ vs. $B_{001}^{\text{Nor.}}$ calculated with Eq. (S11c). The saturation moment for field along the diagonal, (110) direction, is $M_{110}^{\text{Nor.}} \approx 3.8\mu_\text{B}/Yb$, which is much larger than the moment for $B = 14T$ along the crystal (001) direction, $M_{001}^{\text{Nor.}} \approx 0.25\mu_\text{B}/Yb$, at 1.9 K. The anisotropy of the magnetic susceptibility at 1.9 K, obtained from the ratio of the two magnetization slopes in the linear region, $B \approx 2T$, indicates the effective g-factor anisotropy, $\frac{g_{\perp}^{\text{eff}}}{g_{\parallel}^{\text{eff}}} = 7.5(4)$, via $\frac{x_{110}^{\text{Nor.}}}{x_{001}^{\text{Nor.}}} = \left(\frac{g_{\perp}^{\text{eff}}}{g_{\parallel}^{\text{eff}}}\right)^2$.

Calculations show that both Yb1 and Yb2 sites have energetically well separated ground state doublets that are almost pure eigenstates of $|m_j| = \pm 7/2$. As shown in Fig. S4A and Fig. S4B, the high symmetry directions for the Yb1 or Yb2 sites do not coincide with the crystalline $a$, $b$, or $c$ axes. For the Yb1 sites in the $c = 0$ plane [red Yb ions in Fig. S4A], the CEF states most nearly diagonalize if the local $J$-quantization $z$-axis is chosen along the $(1, -1, 0)$ direction [Fig. S4C], and the resulting CEF levels are,

\begin{align}
|E_{\pm}\rangle_0 &= 0.992|\pm 7/2\rangle + 0.100|\pm 3/2\rangle + 0.064|\mp 1/2\rangle + 0.032|\mp 5/2\rangle, \\
|E_{\pm}\rangle_1 &= -0.082|\pm 7/2\rangle + 0.300|\pm 3/2\rangle + 0.355|\mp 1/2\rangle + 0.881|\mp 5/2\rangle, \\
|E_{\pm}\rangle_2 &= 0.092|\pm 7/2\rangle + 0.785|\pm 3/2\rangle + 0.420|\mp 1/2\rangle - 0.446|\mp 5/2\rangle, \\
|E_{\pm}\rangle_3 &= -0.153|\pm 7/2\rangle + 0.832|\pm 3/2\rangle - 0.533|\mp 1/2\rangle + 0.005|\mp 5/2\rangle
\end{align}

with the excited levels, $|E_{\pm}\rangle_1$, $|E_{\pm}\rangle_2$, and $|E_{\pm}\rangle_3$, separated from the ground doublet, $|E_{\pm}\rangle_0$, by energies $\Delta_1 = 37$ meV, $\Delta_2 = 64$ meV, and $\Delta_3 = 94$ meV, respectively. This corresponds to the crystal-field Hamiltonian,

$$H = \sum_{i,j|m_i \leq 2l} B^{m}_j O^m_{2l}(J),$$

where $O^m_{2l}(J)$ are the Stevens operators \cite{46}, and the non-zero crystal-field parameters are, $B^0_2 = -2.242230$, $B^0_4 = -1.890630$, $B^0_6 = -0.002223$, $B^0_8 = 0.021004$, $B^4_2 = -0.046010$, $B^6_0 = 0.000008$, $B^6_2 = -0.000015$, $B^6_4 = 0.000124$, and $B^6_6 = -0.000150$.

Consequently, the magnetic properties of Yb\textsubscript{2}Pt\textsubscript{2}Pb at low temperatures must be dominated by the Yb\textsuperscript{3+} ground state doublet, $|E_{\mp}\rangle_0$. Based on the CEF levels of Eqs. (S8), the calculated Yb magnetic moments for the Yb1 (1−10) sublattice in $B = 14T$ magnetic field applied along different directions are,

\begin{align}
M_{110} &\approx 3.95\mu_\text{B}/Yb, \\
M_{110} &\approx 0.2\mu_\text{B}/Yb, \\
M_{001} &\approx 0.3\mu_\text{B}/Yb.
\end{align}
The moment along any direction other than \((1 - 10)\) is very small, indicating strong Ising-like magnetic anisotropy. Although the point charge model for \(\text{Yb}_2\text{Pt}_2\text{Pb}\) may be expected to be not as quantitative as it would be for an insulating magnetic material [49] due to the screening effect of the conduction electrons, the calculated magnetic moments are in a very good quantitative agreement with the \(M(B)\) anisotropy found in magnetization measurements, as we discuss below [Fig. S4, C and D]. Moreover, the calculated CEF splitting of the ground state doublet [Eq. (S8)], \(\Delta_1\), is in good agreement with the crystal field excitation observed in our supplementary neutron scattering measurements.

Since both Yb sites share the same point symmetry and have very similar Yb-Pt and Yb-Pb distances, the CEF levels and the multiplet wave functions, including the ground state doublet, differ very little for the Yb2 sites from those calculated for Yb1 sites, Eqs. (S8). However, the crystal chemical environment of the Yb2 site is rotated by \(90^\circ\) with respect to that of the Yb1 site in the same \((c = 0, \text{ or } c = 1/2)\) plane. Therefore, the local \(J\)-quantization axes and the magnetic moment directions calculated for the Yb2 sites in the \(c = 0\) plane are now along the \((110)\) direction, as indicated by the blue arrows in Fig. S4A. Similarly, for the \(c = 1/2\) plane, the moments are along \((110)\) for the Yb1 sites, and \((1 - 10)\) for the Yb2 sites [Fig. S4B]. Hence, there are two sets of Yb moments in \(\text{Yb}_2\text{Pt}_2\text{Pb}\), forming two sublattices whose moments are orthogonal, directed along \((1 - 10)\) for sublattice 1 [Fig. S4C], and along \((110)\) for sublattice 2 [Fig. S4D].

The existence of these two orthogonal magnetic sublattices provides consistent and very natural explanation of the anisotropic magnetization in \(\text{Yb}_2\text{Pt}_2\text{Pb}\), as has been proposed in [8, 50, 51]. The measured magnetization with magnetic field along the \((100), (110)\) and \((001)\) principal axes [5, 6] are plotted in Fig. S4E, where the normalization to the units of \(\mu_B/\text{Yb}\) was obtained based on the total number of Yb atoms in the sample. However, the large CEF splitting \((\Delta_1 \approx 37\text{ meV})\), which isolates the ground state doublet, means that the magnetic single ion Ising anisotropy is much stronger than the maximum field, \(B = 14\text{ T}\), of the DC magnetization measurements. Hence, the magnetization is weakly sensitive to magnetic field component perpendicular to the local \(J\)-quantization axis, and only half of the Yb moments could be fully polarized when magnetic field is applied along the \((110)\) direction. Thus, for \(B \parallel (110)\), the measured saturation magnetization, \(M_{110}\text{(Meas.)} = 1.9\mu_B/\text{Yb}\), mainly originates from the sublattice 2 with moments along \((110)\), while the other sublattice, with moments along the \((1 - 10)\) direction, contributes very little. As only half of the Yb moments are contributing, \(M_{110}\text{(Meas.)}\) must be corrected by the factor of two [8, 50, 51], leading to the saturation moment \(M_{110}\text{(Nor.)} = 3.8\mu_B/\text{Yb}\), close to the value calculated by McPhase, Eq. (S10a). When magnetic field is applied along the \((100)\) direction, \(45^\circ\) away from the Ising easy-axis direction of either of the two magnetic sublattices, all Yb magnetic moments are polarized along their own Ising axes, but with only a projection of the magnetic field on these axes being effective, so the measured saturation field along the \((100)\) direction, \(B_{100}\text{(Meas.)}\), is \(\sqrt{2}\) times larger than the saturation field when magnetization is measured along the \((110)\) direction, \(B_{110}\text{(Meas.)}\). Similarly, the measured saturation moment, \(M_{100}\text{(Meas.)} = 2.7\mu_B/\text{Yb}\), is the \(45^\circ\) projection of both \(M_{110}\text{(Nor.)}\) and \(M_{110}\text{(Nor.)}\), which are in fact \(\sqrt{2}\) times larger [8, 50, 51]. Magnetic field along the crystal \((001)\) direction is perpendicular to all
Ising axes, and no moments in either sublattice are substantially polarized in fields as low as 14 T. Consequently, the measured magnetization, $M_{001}(\text{Meas.}) = 0.25 \mu_B/Yb$, at 14 T is very small. We thus establish the following relationships between the actual magnetization of Yb ions, which we call normalized magnetization $M(\text{Nor.})$, and the effective magnetic field acting along the Ising axis, $B(\text{Nor.})$, and the experimentally measured magnetization $M(\text{Meas.})$ and the applied magnetic field $B(\text{Meas.})$.

\begin{align}
M_{110 \text{ or } \overline{1}10}(\text{Nor.}) &= M_{110}(\text{Meas.}) \times 2[= 3.8\mu_B/Yb@14T], B_{110}(\text{Nor.}) = B_{110}(\text{Meas.}), \\
M_{110 \text{ or } \overline{1}10}(\text{Nor.}) &= M_{100}(\text{Meas.}) \times \sqrt{2}[= 3.8\mu_B/Yb@14T], B_{110}(\text{Nor.}) = B_{100}(\text{Meas.})/\sqrt{2}, \\
M_{001}(\text{Nor.}) &= M_{001}(\text{Meas.})[= 0.25\mu_B/Yb@14T], B_{001}(\text{Nor.}) = B_{001}(\text{Meas.}).
\end{align}

(S11a)  
(S11b)  
(S11c)

Fig. S4F shows the normalized magnetization against the accordingly normalized field, as defined by Eqs. (S11). We should note that because of the 4-fold macroscopic symmetry of the two Yb sites and the corresponding two Ising axes, there is always a small contribution coming from the Yb magnetization induced by the field component perpendicular to its Ising axis when magnetic field is applied in the $ab$ plane. For example, $M_{110}(\text{Meas.})$ contains small transverse contribution from the $(1 - 10)$ sublattice, and $M_{100}(\text{Meas.})$ has similar transverse contributions from both sublattices, which are neglected in Eqs. (S11a)-(S11c). However, based on the calculation [Eqs. (S10a)-(S10c)] this contribution is even smaller than the magnetization of $B$ along the $(001)$ direction, which justifies our neglect of this component.

It is also impossible to conclude from the DC magnetization measurements alone, which of the two sublattices (i.e. which of the two Yb sites, Yb1, or Yb2, in a given $ab$-layer) has moments along the $(110)$, and which along the $(1 - 10)$ direction. Given the two-sublattice structure, magnetization cannot distinguish whether moments are parallel or perpendicular to the putative Yb-dimer bond direction [5, 8]. Our point charge model CEF calculations indicate, however, that the Yb Ising moments are perpendicular to the nearest neighbor Yb-Yb bond [Fig. S4], which is consistent with the zero field magnetic structure determined from powder neutron diffraction [18]. We note that the magnetic structure that we infer in Yb$_2$Pt$_2$Pb is quite different from the magnetic structure of U$_2$Pt$_2$Sn, and of the SSL system GdB$_4$ [52, 53].

Finally, two-sublattice structure helps to understand the one-dimensionality of Yb$_2$Pt$_2$Pb. The nearest neighbor Yb-Yb pairs in the crystalline $ab$--plane belong to different sublattices [Fig. S4A–D]. That the two sublattices are easily decoupled even by small magnetic fields and are magnetized separately, indicates that the inter-sublattice interactions are very weak. On the other hand, the distance between the nearest neighbor Yb pairs within each sublattice in the $ab$--plane is $a = 7.76\text{Å}$ [Fig. S4A–D]. Therefore, these Yb are only weakly coupled. In contrast, the distance between Yb from the same sublattice along the crystal $c$--axis is only $\approx c/2, \approx 3.5\text{Å}$ [Fig. S4A–D], and therefore their coupling in this direction is stronger. This immediately leads to one dimensional Yb
chains, or ladders, along the crystallographic \( c \) –axis, and thus the one dimensional dispersion with respect to wave vector \( Q_{00L} \), as seen in Fig. S8, C and D.

Figure S5: **Calculated single-ion magnetization based on the CEF model in Yb\(_2\)Pt\(_2\)Pb.** Magnetization for an isolated Yb ion belonging to the Yb1 (1-10) sublattice as a function of magnetic field \( B \) up to 14 T for the three directions of the field at \( T = 2 \) K (A) and at \( T = 10 \) K (B). Solid curves show fits to the Brillouin function for \( S=1/2 \), which determine the effective spin-1/2 \( g \) — factors describing the ground state doublet.

**Magnetization and single ion g-factors**

The point charge CEF model also allows us to determine the single ion effective spin-1/2 \( g \) — factors for the Yb ground state doublet. To this end, we calculate magnetization curves, \( M(B) \), of the Yb CEF multiplet [Eq. S8] for some finite temperature \( T \) for magnetic field along the three principal crystal axes, which we then fit to a Brillouin function with \( S = 1/2 \),

\[
M_{\alpha} = g_{\alpha}^{\text{eff}} \mu_B S B_{S}(x), \quad B_{S}(x) = \frac{25+1}{25} \coth \left( \frac{25+1}{25} x \right) - \frac{1}{25} \coth \left( \frac{1}{25} x \right), \quad x = \frac{g_{\alpha}^{\text{eff}} \mu_B S B}{k_B T}
\]

Here \( \alpha \) denotes the crystallographic direction, \( M_{\alpha} \) is the magnetization per Yb, \( \mu_B \) is Bohr magneton, \( k_B \) is Boltzman constant, and \( g_{\alpha}^{\text{eff}} \) is the effective \( g \) — factor for magnetic field \( B \) in the direction \( \alpha \), which is obtained from the fit. Calculations for the Yb1 site with the Ising axis along the \((1-10)\) direction for \( T = 2 \) K and \( T = 10 \) K (symbols) are shown together with the corresponding fits (lines) in Fig. S5. The obtained \( g \) — factor values,

\[
\begin{align*}
g_{||}^{\text{eff}} &= g_{1-10}^{\text{eff}} = 7.9 \\
g_{\perp,ab}^{\text{eff}} &= g_{110}^{\text{eff}} = 0.6 \\
g_{\perp,c}^{\text{eff}} &= g_{001}^{\text{eff}} = 0.8
\end{align*}
\]
are in good agreement with the experimental results that we report, and in particular with nearly completely longitudinal character of the magnetic excitation spectrum, which is proportional to \((g_{\alpha}^{\text{eff}})^2\).

Figure S6: (A) Magnetization measured on a typical single crystal of Yb\(_2\)Pt\(_2\)Pb in magnetic field \(B\) along the \((1 - 10)\) lattice direction at \(T = 0.09(1)K\) (dark blue) and 1.5(1) K (red), without the two-sublattice correction (1). (B) Magnetization calculated for the effective spin-1/2 XXZ Hamiltonian (20) with \(g_{\alpha}^{\text{eff}} = 7\), \(\Delta = 2.6\) and \(J = 0.205\) meV for \(T = 0.05J \approx 0.1\) K (dark solid line) and \(T = 0.5J \approx 1.2\) K (blue broken line) using the quantum transfer matrix approach suitable for finite temperature calculations, as described in Methods. Red symbols show magnetization measured in Yb\(_2\)Pt\(_2\)Pb at \(T^\approx 0.09\) K, same as in panel A, but corrected by a factor 2 according to Eq. (S11a) and by subtracting a small linear component accounting for the contribution of the perpendicular sublattice.

The magnetization curves measured in Yb\(_2\)Pt\(_2\)Pb for different directions of magnetic field at \(T = 1.9\) K shown in Fig. S4, E and F, allow us to make an experimental estimate of the effective g-factor anisotropy from the anisotropy of magnetic susceptibility at 1.9 K. Using a simple Curie law for a paramagnet, \(\chi_{\parallel}/\chi_{\perp} = (g_{\parallel}^{\text{eff}}/g_{\perp}^{\text{eff}})^2\), we obtain \(g_{\parallel}^{\text{eff}}/g_{\perp}^{\text{eff}} = 7.5(4)\), which agrees remarkably well with the calculations, Eq. (S13), even though susceptibility at 1.9 K might be markedly affected by the antiferromagnetic correlations that are neglected in the single ion calculation (S13). The magnetization curves for \(B \parallel (1 - 10)\) at \(T = 0.09\) K and 1.5 K are shown in Fig. S6A, while in Fig. S6B the measured magnetization is compared with the theoretical magnetization curves for the spin-1/2 XXZ model Eq. S18, obtained using the quantum transfer matrix approach suitable for finite temperature calculations, similar to the T-dependent magnetic susceptibility shown in Fig. 4B of the manuscript. Both the temperature dependence of magnetic susceptibility (it has also been reported in [6]), and the magnetization curves agree quite well with the calculation for the effective nearest neighbor spin-1/2 XXZ.
Hamiltonian Eq. S18 with $g_{\text{eff}} = 7$, $\Delta = 2.6$ and $J = 0.205$ meV. The largest discrepancy in $M(H)$ occurs near the saturation field, where the high energy four spinon contribution that is absent in the approximate nearest neighbor model becomes important (cf Fig. 1 and the discussion in the manuscript).

**Crystal field excitation**

In order to verify directly the hierarchy of the Yb crystal field levels predicted by the point charge model, we have carried out additional measurements on CNCS, using the incident neutron energy of 37 meV, Fig. S7. Inelastic neutron spectra collected at $T = 175$ K reveal a non-dispersing excitation with an energy $E_c = 25.2(1)$ meV [Fig. S7], which is in reasonably good agreement with the splitting, $\Delta_1 = 37$ meV, between the ground state and the first excited doublet predicted by the point charge model. Unlike the phonon intensity that increases with the increasing magnitude of the wave vector transfer, $Q$, the intensity of the observed excitation decreases at large $L$ (large $Q$), which confirms its magnetic origin. This result renders further direct experimental credibility to our CEF calculations, which are already strongly supported by the magnetization and the magnetic susceptibility data. We have to note that in the early reports by some of the present authors [5], a broad contribution to the low-temperature specific heat was found, whose Schottky-peak model analysis indicated the lowest doublet splitting of $\Delta_1 \approx 7$ meV, significantly lower than the excitation observed here by neutron scattering. However, a more recent, closer inspection of that data lead us to conclude that the “Shottky” peak (e.g. in Fig. 6 of Ref. [5]) likely results from a systematic error in subtracting the Debye contribution [18]. Hence, the heat capacity results can only be understood to indicate a lower limit, $\Delta_1 \approx 7$ meV, on the CEF doublet splitting. Moreover, the temperature dependence of the inverse magnetic susceptibility, which was measured in Ref. [5] up to 800 K, indicates a change of slope at around 300 K, which is fully consistent with $E_c = 25.2(1)$ meV observed here.

Although in principle it is possible that the Yb1 and Yb2 sites could have slightly different crystal field splitting, we do not resolve any such additional peak structure, or any considerable thermal broadening of this level at 175 K, within the $\approx 3.6$ meV energy resolution of this high energy CNCS measurement, Fig. S7B. Taken together, our experimental data confirm that the first excited CEF level in Yb$_2$Pt$_2$Pb is energetically well separated from the ground state doublet, and provide strong support for our CEF point charge model calculations. Thermal population of the excited doublets is negligible within the temperature range, $T_{100K}$, of the measurements that we report in the manuscript.
Figure S7: **Crystal field excitation in Yb$_2$Pt$_2$Pb.** (A) Contour map of the inelastic scattering intensity measured at T ≈ 175K, with energy dispersion along the wave vector $Q_{00L}$, averaged over the perpendicular wave vector components within (HH) = [−5,5], (H−H) = [−0.15,0.15]. The scattering at high wave vectors and energies less than 20 meV is dominated by the aluminum sample holder and the cryogenic environment. (B) Energy dependent intensity averaged over the wave vector range (HH) = [−5,5], (H−H) = [−0.15,0.15], L = [−5,5]. The inset shows the enlarged plot of the CEF excitation around 25.2 meV.

More precisely, the lowest splitting of the CEF doublet manifold, $\Delta_1 \approx 30$ meV, implies that there is no significant thermal occupation of the first excited CEF level for temperatures T lower than $\approx 300$ K, or magnetic fields less than $B \approx 300$ T. The data we present in both the main text of this paper and the supplement were measured well within this field ($B \leq 14$T) and temperature ($T \leq 100$K) range. Thus, the Ising anisotropy of the doublet ground state is robust, and protected by the magnitude of $\Delta_1$ over our full experimental field and temperature window. Neglecting higher energy states, we can separate the two orthogonally aligned Yb moments into two magnetic sublattices based on the direction of their Ising anisotropy axes. Shown in Fig. S4C is the magnetic sublattice with all the Yb1 sites in the c = 0 plane, and all the Yb2 sites in the c = 1/2 plane, with the magnetic moments pointing along the (−1, 1, 0) direction. All the Yb2 sites in the c = 0 plane and all the Yb1 sites in the c = 1/2 plane form a different, but equivalent magnetic sublattice, with all the magnetic moments pointing along the (1, 1, 0) direction [Fig. S4D]. It is worthwhile to notice that the originally proposed Shastry-Sutherland lattice (SSL) based on the crystal structure in the ab plane [5, 6, 8, 50, 51] is now broken, and instead each sublattice consists of square lattices with well separated ladders, or pairs of chains, along the crystal c−axis. Thus it is not surprising that one dimensional dynamics with spinon excitations are observed along $Q_{00L}$ as shown in the manuscript.

**Supplementary Theory**
The emergence of spinons in $\text{Yb}_2\text{Pt}_2\text{Pb}$ becomes less counterintuitive when one steps back and considers the underlying high-energy $f$-electron Hamiltonian, which can be expressed in the form of a one-dimensional Hubbard model with strong spin-orbit coupling [27],

$$H = \sum_{\langle n,m,j,j' \rangle} t_{nn}^{jj'} f_j^+(n) f_{j'}(m) + \sum_{\langle n,j \rangle} U [f_j^+(n) f_j(n) - N]^2 + \sum_{\langle n,j \rangle} \Delta_j f_j^+(n) f_j(n), \quad \text{(S14)}$$

where $n, m$ number the Yb sites, the last term represents the crystal field, $f_j f_{j'}$ are total angular momentum projections, and $N = 7$ is the occupation number of the $4f$-shell with $J = 7/2$. The hierarchy of interactions is such that $U \gg t_{nm}^{jj'} \gtrsim \Delta_j$. Considering $t_{nm}^{jj'}$ and $\Delta_j$ as perturbations, we assume the axial symmetry of the unperturbed system so that the tunneling is diagonal in the total angular momentum projection. In the second order of perturbation in the tunneling matrix element we obtain the exchange interaction in the form,

$$H_{\text{exchange}} = \frac{t_{nm}^{jj'} \delta_{kk}}{U (2N+1)} f_j^+(n) f_k(n) f_k^+(m) f_j(m) \sum_j f_j^+(n) f_j(n) = N. \quad \text{(S15)}$$

When all tunneling matrix elements $t_{nm}^{jj'}$ are equal, the exchange interaction is just a permutation operator, $P_{nm}$, acting on $(2J+1) \times (2J+1)$-dimensional space of sites $n$ and $m$,

$$H_{\text{exchange}} = \frac{t^2}{2 (2N+1) U} P_{nm}. \quad \text{(S16)}$$

This interaction has a high, $SU(2J+1)$ symmetry, and in this form it has been known for a long time in the theory of low dimensional magnetism [23, 24, 25], and is also frequently discussed in the theory of cold atoms (see, for example, [26]). The most striking feature of (S15) or its simplified form (S16) is that, unlike the Heisenberg-Dirac bilinear exchange $\sim \mathbf{J}_n \cdot \mathbf{J}_m$, it contains matrix elements between all angular momentum projections, in the latter case even with equal weight. Thus, it can be represented as a polynomial of order $(2J+1)$ in $J^+, J^-$ raising and lowering operators of the total angular momentum. The next step towards obtaining the effective low energy exchange Hamiltonian is to take into account the crystal field. If, as in our case, it is stronger than the exchange integral $\sim t^2 / U$, then one can project (S15) on the lowest-energy multiplet (in our case it is the $j = \pm 7/2$ Kramers doublet) to obtain,

$$H_{\text{eff}} = \sum_{j,j' = \pm 7/2} \frac{t_{nn}^{jj'} \delta_{kk}}{U (2N+1)} f_j^+(n) f_{j'}(n) f_{j'}^+(m) f_j(m) \sum_{j=\pm 7/2} f_{j}^+(n) f_{j}(n) = 1. \quad \text{(S17)}$$

The operators $S^\alpha = \sum f_j^+ \sigma^\alpha_{j,j'} f_{j'}$ ( $\sigma^\alpha$ being Pauli matrices, $\alpha = x, y, z$), subject to the occupation constraint, represent the components of a spin-1/2 operator. Hence, (S17) is equivalent to the $S = 1/2$ Heisenberg-Dirac XXZ Hamiltonian for pseudospins $S^\alpha$,

$$H = J \sum_n \left( S_n^x S_{n+1}^x + S_n^y S_{n+1}^y + \Delta S_n^z S_{n+1}^z \right). \quad \text{(S18)}$$

[Eq. (1) in the main text], replacing the $f$-electron Hamiltonian (S14) at low energies. The Hamiltonian (S17) accounts for the exchange of the two electrons between the $j = \pm 7/2$ states ($l = \pm 3$ states of the $f$-orbital) on the neighbor sites, Fig. 1D in the main
text. For these states, strong spin-orbit coupling effectively quenches the electronic spin as an independent degree of freedom, simply dictating that it is co-aligned with the orbital angular momentum, $L$, which is the main contributor to the total angular momentum, $J$. Therefore, $U(2)$ pseudospin $S$ essentially describes the orbital degree of freedom [23] of the Yb f-electrons, and the effective spin-1/2 XXZ Hamiltonian (S18) represents the orbital-exchange Hamiltonian (S17). Although it has the familiar Heisenberg-Dirac-like bilinear form for the effective spin-1/2, in the general case it is anisotropic, and can also have non-diagonal elements which couple different spin components, $\sim S_n^a S_m^b$. Most importantly, it contains matrix elements between the states that the bilinear exchange interaction of the form $\sim J_n J_m$ does not have.

**Spinon dispersion in spin-1/2 XXZ model.**

In the Heisenberg-Ising case the anisotropy parameter in the XXZ Hamiltonian, Eq. (S18), is $\Delta > 1$, and the spinon dispersion, $\varepsilon_s(q)$, acquires a finite gap, $\Delta_s$. Using the conventional parameterization of the algebraic Bethe ansatz solution [12, 13, 14],

$$\varepsilon_s(q) = I \sqrt{1 - k^2 \cos^2(qa)} = \sqrt{I^2 \sin^2(qa) + \Delta_s^2 \cos^2(qa)},$$  \hspace{1cm} (S19)

where $q \in [0, \pi]$ is the spinon’s one-dimensional wave vector, $\alpha$ is the lattice spacing in the chain, and the spinon gap, $\Delta_s = I \sqrt{1 - k^2} = I k^l$, and the dispersion bandwidth parameter,

$$I = J \frac{K(k)}{\pi} \sqrt{\Delta^2 - 1},$$  \hspace{1cm} (S20)

are both encoded by the integral equation,

$$\frac{K(k')}{K(k)} = \frac{K(\sqrt{1 - k'^2})}{K(\sqrt{1 - k^2})} = \frac{1}{\pi} \ln(\Delta - \sqrt{\Delta^2 - 1}).$$  \hspace{1cm} (S21)

Here $K(k)$ is the complete elliptic integral of the first kind with elliptic modulus $k$, and $k'$ is the complementary elliptic modulus. For small $k'$, $K(k')$ can be expressed as a power series,

$$K(k') = \frac{\pi}{2} \left( 1 + \left( \frac{1}{2} \right)^2 k'^2 + \left( \frac{3}{8} \right)^2 k'^4 + O(k'^6) \right),$$  \hspace{1cm} (S22)

while for sufficiently large $\Delta$ the logarithm in Eq. (S21) can be Taylor-expanded,

$$\frac{K(k)}{K(\sqrt{1 - k^2})} = \frac{1}{\pi} \ln(2\Delta) - \frac{1}{4\Delta^2} + O(\frac{1}{\Delta^4}).$$  \hspace{1cm} (S23)

Equivalently, the integral equations relating $I$ and $\Delta$ with $k$, $k'$ are [12, 13, 14],

$$\Delta = \cosh \left( \pi \frac{K(k')}{K(k)} \right), \hspace{0.5cm} I = J \frac{K(k)}{\pi} \sinh \left( \pi \frac{K(k')}{K(k)} \right) = J K(k') \sqrt{\frac{\Delta^2 - 1}{-\ln(\Delta - \sqrt{\Delta^2 - 1})}}.$$  \hspace{1cm} (S24)

Using the experimentally determined values $I = 0.322$ meV and $\Delta_s = 0.095$ meV, we obtain $k' = 0.295$, $k = \sqrt{1 - k'^2} = 0.955$, and $K(k') \approx 1.606$, $K(k) \approx 2.638$, $K(k')/K(k) \approx 0.609$. Substituting these values in Eq. (S24), we obtain $\Delta \approx 3.46$, and $J \approx 0.116$ meV, so that $I \approx 2.78J$ instead of $\frac{\pi}{2}J$ in the isotropic Heisenberg case where $k' = 0$.

**The boundaries of 2- and 4-spinon continua.**

In the case of non-interacting quasi-particles, the energy of a multi-particle state is simply given by a sum of energies of the constituent particles, while the total wave vector
is a sum of their wave vectors by the law of quasi-momentum conservation. Such multi-
particle states then form a continuum, whose upper and lower boundaries are determined
by the maximum and the minimum values of the total energy for a given total momentum
and with respect to all possible relative momenta of the constituent particles. Thus, for
the case of the 2-particle spinon continuum we have,
\[ \epsilon_{2s}(q) = \min, \max\{\epsilon_s(q_1) + \epsilon_s(q_2)\}, \]  
(S25)
where the extremum is sought for the fixed total wave vector of the two-particle state,
\( q = q_1 + q_2 \), of two spinons with wave vectors \( q_1 \) and \( q_2 \). The condition can be
eliminated by solving the conservation law for the wave vector of one of the particles,
leading to,
\[ \epsilon_{2s}(q) = \min, \max\{\epsilon_s(q_1) + \epsilon_s(q - q_1)\} = \min, \max\{\epsilon_s(q/2 + q') + \epsilon_s(q/2 - q')\}, \]  
(S26)
where \( q' = (q_1 - q_2)/2 \) is half the spinons relative momentum, \( q_{1,2} = q/2 \pm q' \). This
results in the following minimum/maximum condition,
\[ \epsilon'_s(q/2 + q') = \epsilon'_s(q/2 - q'), \text{ or } \epsilon'_s(q_1) = \epsilon'_s(q_2). \]  
(S27)
which essentially requires that spinons at the lower (upper) boundary have equal velocity,
or that momentum of one of the spinons is at the boundary of the 1-spinon Brillouin zone
(BZ), i.e. \( q_1 = q/2 - q' = \pi \), and \( q_2 = q/2 + q' = q - \pi \equiv q \), or vice versa. For the
spinon dispersion given by Eq. (S19), the equal-velocity condition (S27) within the \([0,\pi]\)
spinon BZ is only achieved for \( q' = 0 \), and therefore the upper and the lower boundaries
of the 2-spinon continuum are defined by the following condition [13, 14, 16],
\[ \epsilon'_{2s}^{\text{up,low}}(q) = \min, \max\{2\epsilon_s(q/2), \Delta_s + \epsilon_s(q)\}, \]  
(S28)
where the minimum and the maximum values of the two expressions, \( 2\epsilon_s(q/2) \) and
\( \Delta_s + \epsilon_s(q) \), determine the lower and the upper boundary of the 2-spinon continuum,
respectively. Near \( q = \pi \) the lower boundary is given by \( \Delta_s + \epsilon_s(q) \), while near \( q = 0 \)
by \( 2\epsilon_s(q/2) \). The situation is opposite for the upper boundary. These boundaries are
shown by solid white lines in Figure 2 of the manuscript.

The boundaries of the 4-spinon continuum are obtained using a similar extremum
search procedure and applying the method of Lagrange multipliers, which leads to the
same requirement of equal velocities. The upper boundaries, which in a range near
\( q = \pi/2 \) are given by \( \max\{4\epsilon_s(q/4), 4\epsilon_s((q + 2\pi)/4)\} \), [14], are shown by broken
white lines in Figure 2 of the manuscript.

**Supplementary neutron scattering data**

**Scattering in zero and high magnetic field and two magnetic sublattices.**

As discussed above, strong Ising anisotropy divides Yb atoms in Yb\(_2\)Pt\(_2\)Pb into
two orthogonal sublattices, with Ising axes along (110) and (1-10) directions,
respectively. Magnetic field \( B \geq B_c \approx 2.3 \text{ T} \) applied along (110), or (1-10) direction, can
separately polarize moments in one of these two sublattices, so that only a single
sublattice contributes to the measured inelastic neutron magnetic scattering at high fields.
Here, we present the neutron scattering data at zero and 4 Tesla, which support this
conclusion.
Figure S8: Magnetic neutron scattering from the two sublattices in Yb$_2$Pt$_2$Pb in 0 T and 4 T magnetic field. (A), (B) Contour maps of the elastic diffraction in the (HH, H–H, 1) scattering plane within $E = [-0.2, 0.2]$ meV, and $Q_{00L} = [0.9, 1.1]$ measured on CNCS at 1.5 K in 0 T (A) and 4 T field along (1-10) (B). (C), (D) Contour maps of the inelastic scattering intensity as a function of energy transfer $E$ and wave vector $Q_{00L}$, within $Q_{HH} = [0,1]$ and $Q_{H–H} = [-0.1,0.1]$, measured on DCS at 0.1 K in zero magnetic
field (C), and in field $B = 4\, \text{T}$ along $(1 - 10)$ (D). The solid lines, dashed lines, and data points are explained in the main text. (E) Energy dependent scattering intensity in magnetic fields 0 T and 4 T, as indicated. The plot is obtained by averaging the neutron scattering intensity over $Q_{HH} = [-0.1,0.1]$, and $Q_{00L} = [0.2]$. (F) Energy integrated scattering intensity in 0 T and 4 T. The total scattering intensity in 0 T (black symbols) is nearly twice the intensity at 4 T (red symbols), as indicated by the open red circles.

First, our neutron scattering measurements demonstrate that two magnetic sublattices with the orthogonal moments order independently below the Neél temperature $T_N = 2.07$ K, and that magnetic order of one of the two sublattices can be tuned separately by a magnetic field along the (1–10) direction. Shown in Fig. S8, A and B are the contour maps of the elastic neutron scattering intensity in the (HH, H–H,1) scattering plane measured on CNCS at 1.5 K, in 0 T and 4 T field, respectively. This plane is perpendicular to the horizontal (HHL) scattering plane [Fig. S1, C and D], and the $Q_{H–H}$ dependence is only resolved by using the vertical position sensitivity of the CNCS detectors. In zero field [Fig. S8A], two sets of satellite peaks, with $Q_1 = (\pm 0.2, \pm 0.2, 1)$ and $Q_2 = (\pm 0.2, \mp 0.2, 1)$, sit symmetrically around the (001) position. This indicates a $5 \times 5$ magnetic superstructure in the $ab$ plane in real space [18], and the propagation vectors $Q_1$ and $Q_2$ correspond to the respective AF ordering of the two sublattices. Application of a 4 T magnetic field along the (1–10) direction polarizes the sublattice with magnetic moments along this direction, while the other, perpendicular sublattice appears essentially unaffected. As sublattice parallel to the field is magnetized, the diffraction intensity of the corresponding antiferromagnetic satellites at $Q_1 = (\pm 0.2, \pm 0.2, 1)$ decreases, and is transferred to the lattice Bragg positions by the arising ferromagnetic component. We indeed observe an increase of the (002) lattice Bragg reflection proportional to the square of magnetization. At 4 T, peaks at $Q_1$ are fully suppressed, while the other set of satellite peaks, at $Q_2$, remain nearly unchanged [Fig. S8B].

The two-sublattice behavior is further revealed by the inelastic neutron scattering data. Shown in Fig. S8C is the contour map of the inelastic scattering intensity as a function of energy transfer and the wave vector $Q_{00L}$, averaged over $Q_{HH} = [0.1]$ and $Q_{H–H} \approx [-0.1,0.1]$, measured on DCS at 0.1 K in nominal magnetic field $B = 0$ T. The solid and dashed lines indicate the lower and upper boundaries of the two-spinon and four-spinon excitations, as explained in the main text. The experiment was repeated with $B = 4$ T, as shown in Fig. 8D, and the energy and wave vector dependencies of the scattered intensity are very similar to those measured in zero field. However, the absolute magnitude of the scattered intensity in 4 T is significantly lower than in zero field (the intensities in Fig. 8 C and D are normalized to units of $\mu_B^2 \cdot \text{meV}^{-1} \cdot \text{per crystal unit cell}$ with 8 Yb). Further quantitative comparison of the 0 T and the 4 T data is presented in Fig. S8E, which displays magnetic scattering intensity averaged over $Q_{00L} = [0.2]$, $Q_{H–H} \approx [-0.1,0.1]$, and $Q_{HH}$ near zero, $Q_{HH} = [-0.1,0.1]$, as a function of energy transfer, $E$. A broad, asymmetric peak is observed in both 0 T and 4 T data at energies from about 0.15 meV to 1.5 meV. This is the spinon continuum, and it is considerably broader than the instrument resolution, which is about 0.1 meV, as estimated from the width of the elastic diffraction peaks. The shape of the broad peak at 4 T is similar to that at 0 T, but with about twice lower intensity. As we integrate this broad continuum with
respect to energy up to an energy $E$ [Fig. S8F], we find that the integrated intensity at 0 T is almost exactly twice the integrated intensity at 4 T. The filled black symbols in Fig. S8F represent the 0 T integrated scattering intensity, the open red circles are twice the integrated scattering intensity at 4 T, and they overlap very closely with the 0 T data within the entire energy range covered, from 0.15 meV up to 1.75 meV.

Another remarkable observation is the notable absence of the coherent spin wave excitation of the fully polarized ferromagnetic sublattice at $B = 4$ T in Fig. S8D. Such spin wave is naturally expected in the fully saturated ferromagnetic state of a spin-1/2 Heisenberg, or XXZ chain at high field, and has been experimentally observed, e. g. in copper sulphate [15]. In Yb$_2$Pt$_2$Pb, however, such excitation, while theoretically present, and contributing to the system’s energy and entropy in the effective spin-1/2 Hamiltonian, is not visible in experiment. The reason is, that it involves *transverse* fluctuations of the effective spins-1/2, and such fluctuations do not lead to any measurable magnetization fluctuations. The latter are almost entirely longitudinal by virtue of the strong anisotropy of the effective $g$-factor. Excitation of a transverse spin wave corresponds to lowering (raising) of the angular momentum of the Yb magnetic moment by $\Delta m_j = \pm 1$ via magnetic dipole interaction with a neutron, or a photon. However, such processes are ruled out by the nearly pure $|m_j = \pm 7/2|$ nature of the Yb ground state doublet, which is protected by a large, $\gtrsim 25$ meV gap to the nearest excited state. Only orbital-exchange processes with $\Delta m_j = 0$ are allowed, as illustrated in Fig. 1 of the main text. This invisible transverse spin wave excitation of the ferromagnetically polarized sublattice at high-field, a “dark magnon” quasi-particle, perhaps offers a condensed matter analogue of the weakly interacting massive particles (WIMPs) of the cosmological dark matter.

The above observations indicate that, while in the absence of magnetic field both sublattices contribute equally to magnetic scattering, when magnetic field is applied along one of the orthogonal Ising directions [here $B \parallel (1 - 10)$] and polarizes magnetic moments in the corresponding sublattice, only fluctuations of the other sublattice, with moments perpendicular to the field direction and essentially unaffected by the magnetic field, contribute to the magnetic scattering intensity. Hence, the difference between the 0 T and the 4 T data yields the net contribution in zero field of the sublattice with moments along the (vertical) field direction, $(1 - 10)$. We use the crystal unit cell with eight Yb atoms, and thus there are four Yb moments in each sublattice. We account for this when normalizing the measured magnetic scattering intensity in units of $\mu_B^2 \cdot \text{meV}^{-1}/\text{Yb}$, as discussed above and shown in figures in the main text and in most of the figures here.

*Polarization factor in the $Q_{HH}$ wave vector dependence of magnetic scattering.*

The energy dependent intensity shown in Fig. S8 E and F was only integrated over $Q_{HH} = [-0.1,0.1]$, and there is a good reason for that. The total intensity integrated over the full Brillouine zone, $Q_{HH} = [0,1]$, in 4 T is much less than a half of the total intensity in 0 T integrated over the same $Q$-range. This results from the different polarization factors for the two sublattices, which clearly reveal the longitudinal (with
respect to the local Yb Ising axis) character of the measured neutron spectra, as we discuss in this section.

CNCS, $E=\left[0.15, 1.5\right]$ meV, $(H-H)=\left[-0.1, 0.1\right]$

A 1.5 K, 0-4.75 T

B 5 K, 0-4.75 T

C 1.5 K, 4.75 T

D 5 K, 4.75 T

CNCS, $E=\left[0.8, 1.0\right]$ meV, $L=\left[0.8, 1.2\right]$, $(H-H)=\left[-0.1, 0.1\right]$

E 1.5 K

F 5 K

Figure 9: Polarization factor and magnetic intensity dependence on $Q_{HH}$. (A), (B) Contour maps of the inelastic scattering intensity ($E=\left[0.15, 1.5\right]$ meV, $Q_{HH}=\left[-0.1, 0.1\right]$) in the (HHL) scattering plane measured on CNCS at $T=1.5$ K (A)
and at $T = 5$ K (B), with the 4.75 T high field data subtracted. (C), (D) Contour maps of the inelastic scattering intensity ($E = [0.15, 1.5]$ meV, $Q_{H-H} = [-0.1, 1.0]$) in the (HHL) scattering plane measured on CNCS in 4.75 T at $T = 1.5$ K (C) and at $T = 5$ K (D). (E), (F) Inelastic scattering intensity integrated within $E = [0.8, 1]$ meV and averaged over $Q_{000} = [-0.8, -1.2]$, $Q_{H-H} = [-0.1, 0.1]$, as a function of wave vector $Q_{HH}$, at temperature 1.5 K (E) and 5 K (F), for different magnetic fields as indicated in the figure. The white dashed lines in (A), (B), (C) and (D) indicate the region corresponding to the line cuts shown in (E) and (F). The black and red solid lines indicate the different $Q_{HH}$ dependent polarization factors of the two sublattices, as explained in the text. The 4.75 T data in (F) (solid red circles) give the open red circles when a constant is subtracted, as explained in the text.

Since neutrons only scatter from magnetic moments that are perpendicular to the wave vector $Q$, the scattering from the two sublattices will have different polarization factors in the (HHL) scattering plane due to their different moment orientations. The moments in the field-dependent (1-10) sublattice directed along magnetic field are vertical and perpendicular to the wave vectors in the scattering plane, which leads to the polarization factor of 1, independent of $Q$. Moments of the perpendicular, (110) sublattice, lie in the scattering plane, and therefore their contribution depends strongly on the alignment of $Q$ with respect to the (110), or, equivalently, (001) direction. Shown in Fig. S9A is the contour map of the inelastic scattering intensity ($E = [0.15, 1.5]$ meV, $Q_{H-H} = [-0.1, 1.0]$) in the (HHL) scattering plane, measured at $T = 1.5$ K, with the 4.75 T high field data directly subtracted. This is the net inelastic scattering that originates from the sublattice with moments along the (1-10) direction, perpendicular to the (HHL) scattering plane. We observe stripes of intensity that are symmetrically positioned at $Q_{000} = \pm 1$ and are nearly independent of $Q_{HH}$, consistent with the $Q_{HH}$-independent polarization factor. The inelastic scattering intensity measured at 1.5 K in 4.75 T is rather different, as shown in Fig. S9C. This intensity originates almost entirely from the sublattice with magnetic moments along the (110) direction, which lies in the (HHL) scattering plane. We observe that scattering intensity in 4.75 T decreases continuously with the increasing wave vector $Q_{HH}$.

In Fig. S9E, the inelastic scattering intensity at 1.5 K, integrated within $Q_{000} = [-0.8, -1.2]$, $Q_{H-H} = [-0.1, 0.1]$ [white dashed region in Fig. S9, A and C] and $E = [0.8, 1]$ meV is plotted for the two sublattices as a function of wave vector $Q_{HH}$. The intensity at $B = 4.75$ T (red circles) decreases quickly with the increasing $Q_{HH}$, and is well described by the polarization factor for magnetic moment fluctuations along the (110) direction (red solid line),

$$P_{(110)} = 1 - \frac{Q_{110}^2}{Q_{H-H}^2} = 1 - \frac{(a_{110}H)^2 + (cL)^2}{(a_{110}H)^2 + (cL)^2}.$$

(S29)

In contrast, for the zero field minus 4.75 T data, where the fluctuating magnetic moments are along (1-10), i.e. perpendicular to the (HHL) plane, the polarization factor is 1. Consistent with this, apart from a small dip around $Q_{HH} = 1$ that we explained above is a consequence of absorption, the integrated intensity of the $I(0T) - I(4.75T)$ data is almost constant along $Q_{HH}$, as indicated by the solid black line. Data in Fig. S9 B, D, and F show the scattering measured in the paramagnetic state at $T = 5$ K. Both the difference, $I(0 T) - I(4.75 T)$ [Fig. S9B], and the 4.75 T data [Fig. S9D], at 5 K demonstrate very
similar \( Q_{HH} \) dependencies to those measured at 1.5 K [Fig. S9, A and C]. The nearly \( Q_{HH} \) independent stripe of scattering in Fig. S9B and the black solid symbols in Fig. S9F are consistent with the idea that the \( I(0\ T) - I(4.75\ T) \) data represent scattering from the sublattice with moments along the field, (1-10) direction, perpendicular to the (HHL) scattering plane, whose polarization factor is 1, independent of \( Q_{HH} \). The \( Q_{HH} \) dependence of the 4.75 T data [solid red symbols in Fig. 9F] is similar to that at 1.5 K, up to a \( Q_{HH} \)-independent constant. This additional constant contribution is not unexpected, and can be explained by the increased importance of thermal fluctuations at 5 K. As a result, \( B = 4.75 T \) field appears insufficient to fully polarize the moments of the (1-10) sublattice, and it can still contribute to the inelastic scattering intensity. We believe that this is the origin of the extra \( Q_{HH} \) independent scattering at 4.75 T, which we therefore subtract in order to isolate the net contribution from the (110) sublattice [open red circles in Fig. 9F]. As indicated by the red solid line, the resulting \( Q_{HH} \) dependence is well described by the same polarization factor of Eq. (S29), as for the 1.5 K data. These observations show that strong Ising anisotropy imposed by the crystal field constrains the moments to fluctuate only along their own Ising axes both in the ordered state, where, as a result, magnetic fluctuations are parallel to the ordered moments, and in the paramagnetic state above the ordering temperature \( T_N = 2.07 \) K, where fluctuations remain uniaxial.

That the dependence of magnetic scattering intensity on the wave vector transfer in the \( ab \) plane can be explained entirely by the magnetic polarization factor and has no \( Q_{HH} \) dependent structure factors, such as would be expected for a dimer, or a ladder, is the most direct evidence that Yb magnetic moments form nearly uncoupled 1D chains, which fluctuate independently.

The temperature dependence of magnetic scattering.

In this section, we show how the fluctuating moments change as a function of temperature, below and above the AF ordering temperature \( T_N = 2.07 \) K. Since the experimental temperature range for the DCS experiment is below \( T_N \), we will mostly focus on the CNCS data from 1.5 K up to 100 K. Shown in Fig. S10A is the contour map of the spinon dispersion along \( Q_{00L} \) with \( Q_{HH} = [0,1], Q_{H-H} = [0,1] \), measured at 1.5 K, with the high field 4.75 T data subtracted. The overall fan shape along the wave vector \( Q_{00L} \) is similar that observed at 0.1 K, but the dispersion appears smeared due to thermal fluctuations. The energy dependence of the scattered intensity at 1.5 K summed over \( Q_{00L} = [0, \pm 2], Q_{HH} = [0,1] \) and \( Q_{H-H} = [0,1] \) is plotted in Fig. S10B. The spinon continuum centered around 0.5 meV is much broader than the instrumental resolution, which is indicated by the gray area. Integrating the scattered intensity over the first Brillouin zone, we find the fluctuating moment squared of about \((8.4 \pm 1.2) \mu_B^2/Yb\) which is slightly larger than what is found at 0.1 K. The error bars here are dominated by the systematic error, which we estimate based on the difference of the integrated intensity over the wave vector ranges of \( Q_{00L} = [0, -2] \) (black circles and line) and \( Q_{00L} = [0,2] \) (red circles and line) [Fig. S10B].
Figure S10: **Temperature dependence of the inelastic magnetic scattering in Yb$_2$Pt$_2$Pb.** (A) Contour map of the energy dependence of the neutron scattering intensity along $Q_{00L}$ with $Q_{HH} = [0,1], Q_{HH} = [-0.1,0.1]$ at 1.5 K, with the 4.75 T high field data subtracted. (B) The energy dependence of the $I(0 \text{T}) - I(4.75 \text{T})$ scattering measured at 1.5 K, summed over $Q_{00L} = [0, \pm 2], Q_{HH} = [0,1]$ and $Q_{HH} = [-0.1,0.1]$. (C) Contour map of the spinon dispersion along $Q_{00L}$ with $Q_{HH} = [0,1], Q_{HH} = [-0.1,0.1]$ at 5 K, with the 4.75 T high field data subtracted. (D) The energy dependence of the $I(0 \text{T}) - I(4.75 \text{T})$ scattering measured at 5 K summed over
\[ Q_{\text{OO}L} = [0, \pm 2], Q_{\text{HH}} = [0,1] \text{ and } Q_{H-H} = [-0.1,0.1]. \] 
\( E \) Contour map of the spinon dispersion along \( Q_{\text{OO}L} \) with \( Q_{\text{HH}} = [0,1], Q_{H-H} = [-0.1,0.1] \), at 100 K in zero field. (F) Energy dependence of the 0 T scattering measured at 100 K, summed over \( Q_{\text{OO}L} = [0, \pm 2], Q_{\text{HH}} = [0,1] \) and \( Q_{H-H} = [-0.1,0.1] \). The gray area in (B), (D), (F) indicates the energy range of elastic diffraction, mainly determined by the instrumental resolution, which is \( \approx 0.1 \) meV in this experiment.

Once the static order disappears above \( T_N \), all the Yb moments are fluctuating. Figure S10C is the contour map of the spinon dispersion measured at 5 K, summed over the same range of wave vectors as at 1.5 K. The overall dispersion along \( Q_{\text{OO}L} \) is now considerably smeared by thermal fluctuations, and the net fluctuating moment obtained by summing the intensity at 5 K over the first Brillouin zone [Fig. S10D] is \((16.4 \pm 1.8) \mu_B^2/Yb\), almost twice the value at 1.5 K. Roughly speaking, about half of the total Yb moments are ordered in the AF state, while the other half remain fluctuating even at the lowest temperatures. This is a very different situation from that found in most rare earth based classical magnets, where spin wave excitations exist only in the presence of magnetic order, and only slightly suppress the magnitude of the ordered moments. The spinon dispersion at 100 K is shown in Fig. S10E. At this temperature, the magnetic field required to fully polarize the sublattice with moments in the field direction is much larger than 4.75 T, which is the largest field available in this experiment. Consequently, we report only zero field measurements. As expected, thermal fluctuations substantially broaden the spectrum for both neutron energy gain and neutron energy loss. The wave vector integrated intensity at 100 K is plotted as a function of energy in Fig. S10F, where the black and red lines are again obtained by integrating the intensity within \( Q_{\text{OO}L} = [0, \pm 2] \). This procedure yields a total fluctuating moment of \((17.3 \pm 1.5) \mu_B^2/Yb\), which is, within our experimental accuracy, identical to the one we found at 5 K. This result is explained by pointing out that little variation in the fluctuating moment is expected as long as the experimental temperature is small compared to the energy splitting, \( \Delta_1 \sim 30 \) meV, of the CEF ground state and the first excited state.
References and Notes


6. See supplementary materials on Science Online for details.


37. More information about the instrument configuration is on the CNCS website: http://neutrons.ornl.gov/cncs/.


Fig. S1.

(A) Single crystal sample holder used for neutron scattering experiments at DCS at NCNR, NIST, and CNCS, SNS. Single crystals of Yb$_2$Pt$_2$Pb were aligned with the (110) crystal direction normal to the surface of the holder plate, and with the (1 1 0) direction parallel to the long side of the rectangular holder. (B) Schematic view of the configuration for both DCS and CNCS neutron scattering setups. Magnetic field B is applied along the vertical, (H 1 0) direction. The sample rotates in the horizontal, (H H L) plane, with access to the vertical (H -H L) scattering plane limited by the acceptance angle, ±θ. The distance D from sample to detector and the detector height, H, for both DCS and CNCS experiments are described in the text. (C), (D) Contour maps of elastic neutron scattering intensity integrated within the energy range $E = [-0.2,0.2]$ meV, measured in zero field on DCS (C) and CNCS (D). For the DCS data, the detectors are only position sensitive in the horizontal direction, and all scattering in the vertical, (H -H) direction is projected onto the (H H L) scattering plane, (C). For the CNCS data, the
detectors have spatial resolution in both horizontal and vertical directions. The data in (D) are integrated only within a narrow region, \(Q_{H-H} = [-0.05,0.05]\), in the vertical direction. The diffraction seen in the DCS data (C) at \(Q = (001)\) is picked up by the relaxed vertical resolution of that measurement and is absent in the CNCS data (D), as indicated by red arrows.
Fig. S2

Magnetic and nuclear Bragg intensities in Yb$_2$Pt$_2$Pb in zero field. (A) Ordered moment squared as a function of temperature. Black solid circles are the temperature scan at the (0.2, 0.2, 1) AF peak position at $2\theta = 20^\circ$ [18], which was put on the absolute scale using the ordered moment $M_{\text{order}} = 2.5\mu_\text{B}/\text{Yb}$ (red solid diamond) extracted from the refinement of the powder diffraction pattern measured in a $\theta-2\theta$ scan at 1.5 K. (B) Comparison of the calculated nuclear and magnetic structure factors and the experimental integrated peak intensities for peaks accessed at 1.5 K at CNCS and listed in Tables S1 and S2. The slope in the limit of small intensity gives the normalization factor for the experimental CNCS data to absolute values, $1/C = \int d^3Q \int dE I(Q,E)/|F_{hkl}|^2$ (solid red line). The dashed line is a fit to the semi-empirical secondary extinction factor for the single crystal diffraction intensity, as explained in the text.
Fig. S3

**Effect of absorption in Yb\textsubscript{2}Pt\textsubscript{2}Pb.** (A), (B) Inelastic neutron intensity in the (HHL) scattering plane integrated within the energy range $E = [0.15, 1.5]$ meV measured at zero field, with the 4 T high field data subtracted, without the absorption correction (A) and with the absorption correction (B). The white dashed lines in (A) indicate the region where the measured intensity has largest suppression due to absorption, which is corrected in (B). The overall intensity is also enhanced upon accounting for absorption in (B). (C), (D) Comparison of the calculated nuclear and magnetic structure factors and the experimental integrated diffraction peak intensities at 0.1 K measured on DCS without (C) and with (D) the absorption correction. The solid and dashed lines are as in Fig. S2, and are explained in the text. (E), (F) Normalized inelastic scattering in the (HHL) scattering plane integrated within the energy range $E = [0.15, 1.5]$ meV measured in 0 T, with the 4 T high field data subtracted, without the absorption correction (E) and with the absorption correction (F). (G), (H) Energy dependence of the scattering averaged over $Q_{HH} = [0, 1]$ and $Q_{002} = [0, 2]$, as indicated by the dashed rectangles in (E) and (F). The red lines in (C) and (D) are the energy integrated fluctuating moments without the absorption correction (G) and with the absorption correction (H).
**Fig. S4**

**Yb crystal environment and magnetic moments in Yb$_2$Pt$_2$Pb.** (A), (B) Chemical environment for Yb1 (red) and Yb2 (blue) sites in the $c = 0$ (A), and $c = 1/2$ (B) plane, respectively. The nearest Pt (gray) and Pb (black) neighbors are located below and above the corresponding $c = 0$ and $c = 1/2$ plane [5]. Red and blue arrows indicate magnetic moment directions consistent with the point charge model calculation [45, 46, 47, 48]. (C), (D) Simplified schematics of the two magnetic sublattices, with Yb moments along
(-110) direction [sublattice 1, (C)], and along (110) direction [sublattice 2, (D)], respectively. (E) Experimentally measured magnetization at \( T = 1.9 \) K, normalized to the \( \text{Yb}_2\text{Pt}_2\text{Pb} \) formula unit, for magnetic field \( B \) applied along the three principal crystal directions, (110) [blue], (100) [red], and (001) [black]. (F) Magnetization normalized with the account of the two sublattice model illustrated in (A)-(D), as explained in the text. Plotted in blue is \( M_{110} \, (\text{Nor}.) \) vs. \( B_{110} \, (\text{Nor}.) \) calculated with Eq. (S11a), in red \( M_{110} \, (\text{Nor}.) \) vs. \( B_{110} \, (\text{Nor}.) \) calculated with Eq. (S11b), and in black \( M_{001} \, (\text{Nor}.) \) vs. \( B_{001} \, (\text{Nor}.) \) calculated with Eq. (S11c). The saturation moment for field along the diagonal, (110) direction, is \( M_{110} \, (\text{Nor}.) \approx 3.8\mu_B/\text{Yb} \), which is much larger than the moment for \( B = 14T \) along the crystal (001) direction, \( M_{001} \, (\text{Nor}.) \approx 0.25\mu_B/\text{Yb} \), at 1.9 K. The anisotropy of the magnetic susceptibility at 1.9 K, obtained from the ratio of the two magnetization slopes in the linear region, \( B \lesssim 2T \), indicates the effective g-factor anisotropy, \( g_{||}^{\text{eff}}/g_{\perp}^{\text{eff}} = 7.5(4) \), via \( \frac{\chi_{110} \, (\text{Nor}.)}{\chi_{001} \, (\text{Nor}.)} = \left( \frac{g_{||}^{\text{eff}}}{g_{\perp}^{\text{eff}}} \right)^2 \).
Fig. S5

Calculated single-ion magnetization based on the CEF model in Yb$_2$Pt$_2$Pb.

Magnetization for an isolated Yb ion belonging to the Yb1 (1-10) sublattice as a function of magnetic field B up to 14 T for the three directions of the field at T = 2 K (A) and at T = 10 K (B). Solid curves show fits to the Brillouin function for S=1/2, which determine the effective spin-1/2 $g$ -- factors describing the ground state doublet.
Fig. S6

(A) Magnetization measured on a typical single crystal of Yb$_2$Pt$_2$Pb in magnetic field B along the (1 - 10) lattice direction at T = 0.09(1) K (dark blue) and 1.5(1) K (red), without the two-sublattice correction [1]. (B) Magnetization calculated for the effective spin-1/2 XXZ Hamiltonian (20) with $g^{eff} = 7$, $\Delta = 2.6$ and $J = 0.205$ meV for $T = 0.05J \approx 0.1$ K (dark solid line) and $T = 0.5J \approx 1.2$ K (blue broken line) using the quantum transfer matrix approach suitable for finite temperature calculations, as described in Methods. Red symbols show magnetization measured in Yb$_2$Pt$_2$Pb at $T \approx 0.09$ K, same as in panel A, but corrected by a factor 2 according to Eq. (S11a) and by subtracting a small linear component accounting for the contribution of the perpendicular sublattice.
Fig. S7

Crystal field excitation in Yb$_2$Pt$_2$Pb. (A) Contour map of the inelastic scattering intensity measured at $T \approx 175$K, with energy dispersion along the wave vector $Q_{00L}$, averaged over the perpendicular wave vector components within $(HH) = [-5,5]$, $(H-H) = [-0.15,0.15]$. The scattering at high wave vectors and energies less than 20 meV is dominated by the aluminum sample holder and the cryogenic environment. (B) Energy dependent intensity averaged over the wave vector range $(HH) = [-5,5]$, $(H-H) = [-0.15,0.15]$, $L = [-5,5]$. The inset shows the enlarged plot of the CEF excitation around 25.2 meV.
Magnetic neutron scattering from the two sublattices in Yb$_2$Pt$_2$Pb in 0 T and 4 T magnetic field. (A), (B) Contour maps of the elastic diffraction in the (HH, H–H, 1) scattering plane within $E = [-0.2, 0.2]$ meV, and $Q_{00L} = [0.9,1.1]$ measured on CNCS at 1.5 K in 0 T (A) and 4 T field along (1-10) (B). (C), (D) Contour maps of the inelastic scattering intensity as a function of energy transfer $E$ and wave vector $Q_{00L}$, within...
$Q_{HH} = [0,1]$ and $Q_{H-H} \approx [-0.1,0.1]$, measured on DCS at 0.1 K in zero magnetic field (C), and in field $B = 4\, T$ along $(1-10)$ (D). The solid lines, dashed lines, and data points are explained in the main text. (E) Energy dependent scattering intensity in magnetic fields 0 T and 4 T, as indicated. The plot is obtained by averaging the neutron scattering intensity over $Q_{HH} = [-0.1,0.1]$, and $Q_{00L} = [0,2]$. (F) Energy integrated scattering intensity in 0 T and 4 T. The total scattering intensity in 0 T (black symbols) is nearly twice the intensity at 4 T (red symbols), as indicated by the open red circles.
CNCS, $E=[0.15, 1.5]\, \text{meV}$, $(H-H)=[-0.1, 0.1]\]

(A) 1.5 K, 0-4.75 T

(B) 5 K, 0-4.75 T

(C) 1.5 K, 4.75 T

(D) 5 K, 4.75 T

CNCS, $E=[0.8, 1.0]\, \text{meV}$, $L=[0.8, 1.2]$, $(H-H)=[-0.1, 0.1]\]

(E) 1.5 K

(F) 5 K

**Fig. S9**

**Polarization factor and magnetic intensity dependence on $Q_{HH}$**

(A), (B) Contour maps of the inelastic scattering intensity $(E = [0.15,1.5]\, \text{meV}$, $Q_{HH} = [-0.1,0.1]$) in the (HHL) scattering plane measured on CNCS at $T = 1.5$ K (A) and at $T = 5$ K (B), with the 4.75 T high field data subtracted. (C), (D) Contour maps of the inelastic scattering intensity $(E = [0.15,1.5]\, \text{meV}$, $Q_{HH} = [-0.1,0.1]$) in the (HHL) scattering plane.
measured on CNCS in 4.75 T at T = 1.5 K (C) and at T = 5 K (D). (E), (F) Inelastic scattering intensity integrated within $E = [0.8,1]$ meV and averaged over $Q_{00L} = [-0.8,-1.2], Q_{H-H} = [-0.1,0.1]$, as a function of wave vector $Q_{HH}$, at temperature 1.5 K (E) and 5 K (F), for different magnetic fields as indicated in the figure. The white dashed lines in (A), (B), (C) and (D) indicate the region corresponding to the line cuts shown in (E) and (F). The black and red solid lines indicate the different $Q_{HH}$ dependent polarization factors of the two sublattices, as explained in the text. The 4.75 T data in (F) (solid red circles) give the open red circles when a constant is subtracted, as explained in the text.
Fig. S10
Temperature dependence of the inelastic magnetic scattering in Yb$_2$Pt$_2$Pb. (A) Contour map of the energy dependence of the neutron scattering intensity along $Q_{00L}$ with $Q_{HH} = [0, 1]$, $Q_{H-H} = [-0.1, 0.1]$ at 1.5 K, with the 4.75 T high field data subtracted. (B) The energy dependence of the I(0 T) - I(4.75 T) scattering measured at 1.5 K, summed over $L=\lfloor-2, 0\rfloor$, $L=\lfloor0, 2\rfloor$ and $Q_{HH} = [0, 1]$, $Q_{H-H} = [-0.1, 0.1]$. (C) Contour map of the spinon dispersion along $Q_{00L}$ with $Q_{HH} = [0, 1]$, $Q_{H-H} = [-0.1, 0.1]$ at 5 K, with the 4.75 T high field data subtracted. (D) The energy dependence of the I(0 T) - I(4.75 T)
scattering measured at 5 K summed over \( Q_{00L} = [0, \pm 2], Q_{HH} = [0,1] \) and \( Q_{H-H} = [-0.1,0.1] \). (E) Contour map of the spinon dispersion along \( Q_{00L} \) with \( Q_{HH} = [0,1], Q_{H-H} = [-0.1,0.1] \) at 100 K in zero field. (F) Energy dependence of the 0 T scattering measured at 100 K, summed over \( Q_{00L} = [0, \pm 2], Q_{HH} = [0,1] \) and \( Q_{H-H} = [-0.1,0.1] \). The gray area in (B), (D), (F) indicates the energy range of elastic diffraction, mainly determined by the instrumental resolution, which is \( \approx 0.1 \) meV in this experiment.
**Table S1.**
The calculated nuclear structure factors and the experimentally measured integrated intensities for the primary diffraction peaks accessed in the (HHL) scattering plane in the CNCS single crystal neutron scattering measurements at 1.5 K in zero field.

<table>
<thead>
<tr>
<th>Nuclear Diffraction</th>
<th>(002)</th>
<th>(110)</th>
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<tr>
<td>$</td>
<td>F_{hkl}</td>
<td>^2$ (barn·fu $^{-3}$)</td>
</tr>
<tr>
<td>$\int d^3Q \int dEI(Q,E)$(meV)</td>
<td>0.00165</td>
<td>0.00281</td>
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Table S2.
List of the calculated antiferromagnetic structure factors, $|F_{hk1}|^2$, and the experimentally measured integrated intensities for magnetic diffraction peaks accessed in the (HHL) scattering plane in the CNCS single crystal neutron measurements at 1.5 K in zero field ($|F(Q)|^2$ is magnetic form factor [39] squared). The ordered moment at 1.5 K is $M_{\text{order}} = 2.5 \mu_B/Yb$, which was extracted from the order parameter measurement shown in Fig. S2A.

<table>
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<tr>
<th>AF Diffraction</th>
<th>(1.2, 1.2, 0)</th>
<th>(1.8, 1.8, 0)</th>
<th>(0.2, 0.2, 1)</th>
<th>(0.8, 0.8, 1)</th>
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<tr>
<td>$</td>
<td>Q_{hk1}</td>
<td>$ (Å$^{-1}$)</td>
<td>1.373</td>
<td>2.06</td>
</tr>
<tr>
<td>$</td>
<td>F(Q)</td>
<td>^2$</td>
<td>0.924</td>
<td>0.84</td>
</tr>
<tr>
<td>$</td>
<td>F_{hk1}</td>
<td>^2$ (barn-fu$^{-1}$)</td>
<td>0.0636</td>
<td>0.1187</td>
</tr>
<tr>
<td>$\int d^3Q \int dE I(Q,E)$ (meV)</td>
<td>0.0001219</td>
<td>0.00023148</td>
<td>0.001805</td>
<td>0.002754</td>
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### Table S3.
Neutron scattering and absorption cross sections for Yb, Pt and Pb [42].

<table>
<thead>
<tr>
<th></th>
<th>$\sigma_{coh}$ (barn)</th>
<th>$\sigma_{inc}$ (barn)</th>
<th>$\sigma_{abs}$ (barn)</th>
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<tr>
<td>Yb</td>
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<td>Pt</td>
<td>11.58</td>
<td>0.13</td>
<td>10.3</td>
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<tr>
<td>Pb</td>
<td>11.115</td>
<td>0.003</td>
<td>0.171</td>
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