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* In Kazan University the Electron Paramagnetic Resonance (EPR) was discovered by Zavoisky E.K. in 1944.
Magnetic properties of powders LiTbF$_4$ and TbF$_3$\textsuperscript{†}

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Samples LiTbF$_4$ and TbF$_3$ were synthesized by modified methods of colloidal chemistry. The magnetization of these samples was measured in the external magnetic field at 100 Oe and 1 T and in temperature range 2-300 K. Temperatures of phase transition to the magnetic ordering dipolar ferromagnet state were determined for synthesized samples.

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\textbf{Keywords:} synthesis, magnetization, Curie temperature

1. Introduction

The complex fluorides LiReF$_4$ (Re is rare earth ion) represent a class of crystal materials used as model objects in physics of dipolar magnets. Crystals of rare-earth tetrafluoride compounds with a controlled size, shape, structure and surface have unique optical, electronic, magnetic and catalytic properties important for practical applications \cite{1}. The LiTbF$_4$ crystal has a scheelite (CaWO$_4$) structure with the space group $\text{C}_{6}^{4}h$ \cite{2} and TbF$_3$ crystallizes in orthorhombic $D_{16}^{2h}$ space group \cite{3}. Trifluorides exhibit distinct magnetic properties at low temperatures and they are of interest as model systems for the theoretical study of magnetic ordering in condition of competition between the dipole-dipole and exchange interactions \cite{4}.

Measurements of magnetic DC-susceptibility in LiTbF$_4$ single crystals showed that LiTbF$_4$ is the dipole Ising dielectric ferromagnet at temperatures below the $T_c = 2.89$ K \cite{5,6}. Magnetic moments of Tb$^{3+}$ are ordered along the c-easy axis at $T < T_c$. The TbF$_3$ is also a dipolar ferromagnet, as it was found in the magnetization measurements \cite{4}. The phase transition in TbF$_3$ single crystal from the paramagnetic to the ferromagnetic state occurs at temperatures below $T_c = 3.95$ K \cite{4}. At low temperatures ($T < T_c$) magnetic moments of ions Tb$^{3+}$ (∼9 µB) are ordered in two magnetically non-equivalent sublattices in the crystallographic ac plane at angles $\varphi = \pm 28^\circ$ to the a-axis. Due to large values of the magnetic moments, the magnetic ordering is induced mainly by classical dipole-dipole interaction between the terbium ions, which dominates the magnetic exchange interaction.

The tetrafluorides LiTbF$_4$, LiDyF$_4$, LiHoF$_4$ and LiErF$_4$ have been the subject of intensive studies by NMR and EPR. The nuclear magnetic resonance rotation spectra were observed and contributions to the local magnetic fields from the dipole and the transferred hyperfine interactions were distinguished \cite{7}. The fluorine and lithium NMR line shifts have been measured \cite{8} in the temperature range from 300 to 1.3 K and in the fields up to 40 kG for LiTbF$_4$, LiHoF$_4$. Angular dependences of fluorine NMR spectra have been observed in the external magnetic field,

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Magnetic properties of powders LiTbF$_4$ and TbF$_3$

![Figure 1. XRD patterns of samples 1-3.](image)

Table 1. Characteristic size of samples 1-3 obtained by Photocor Complex.

<table>
<thead>
<tr>
<th>Sample</th>
<th>Characteristic size (diameter)</th>
</tr>
</thead>
<tbody>
<tr>
<td>LiTbF$_4$(79.9%)+TbF$_3$(20%)</td>
<td>20 mkm</td>
</tr>
<tr>
<td>LiTbF$_4$(6.9%)+TbF$_3$(81.2%)</td>
<td>1.5 mkm</td>
</tr>
<tr>
<td>TbF$_3$</td>
<td>585 nm</td>
</tr>
</tbody>
</table>

Influence of nanoparticle size on the magnetic, optical and electronic properties of the fluorides is one of the fundamental problems in the physics of the rare-earth compounds and fundamental physics. Nanoparticles of DyF$_3$ were synthesized by the colloidal chemistry methods [12]. The influence of the microwave-assisted hydrothermal treatment on magnetic properties and structure of PrF$_3$ and LaF$_3$ nanoparticles was investigated by XRD, TEM, NMR and EPR [12–21]. In this paper we present the few synthesis technique for TbF$_3$ and LiTbF$_4$ materials. Magnetic and structure properties of the TbF$_3$ and LiTbF$_4$ powders are investigated by DC-susceptibility measurements.

2. Syntesis of micro- and nano-sized powders

The synthesis of LiTbF$_4$ is complex procedure [1,22]. The method of synthesis with trifluoroacetate precursors is chemically aggressive, the methods of hydrothermal synthesis and thermal decomposition demand high temperatures and high pressure in autoclave [22–28].

The samples of LiTbF$_4$ were synthesized using following techniques:

**Sample 1:** LiNO$_3$(aq)+Tb(NO$_3$)$_3$+NH$_4$(aq)$\rightarrow$powder$^{400^\circ,4h}$LiTbF$_4$(s)+TbF$_3$(s).

**Sample 2:** LiOH(aq)+Tb(NO$_3$)$_3$+NH$_4$(aq)$\rightarrow$powder$^{400^\circ,4h,\text{CF}}$LiTbF$_4$(s)+TbF$_3$(s).

The TbF$_3$ nanoparticles were synthesized using standard technologies [10,11].

**Sample 3:** Tb(NO$_3$)$_3$+NaF(aq)$\rightarrow$TbF$_3$(s)+NaNO$_3$(aq).

The X-ray analysis was carried out by an automatic X-ray diffractometer Bruker D8 Advance with Cu K$_\alpha$ radiation ($\lambda = 1.5418$ A) at Bragg-Brentano geometry. Result of the X-ray analysis of synthesized samples is shown in Fig. 1. Phase ratio of LiTbF$_4$ and TbF$_3$ in samples 1-2 was determined using MAUD software [29].
3. Study of magnetic properties, discussion and results

The temperature dependencies of magnetic susceptibility for all synthesized samples were measured by PPMS-9 System (Quantum Design Inc.) at applied magnetic fields 100 Oe and 1 T in temperature range 2-300 K. The derivative of magnetic susceptibility versus temperature for the sample 1 is shown in Fig. 2. There are two phase transitions at temperatures $T = 2.9K$ and $T = 3.93K$, which correspond to the Curie temperature of the transitions to the ferromagnetic state for LiTbF$_4$ and TbF$_3$ single crystals, respectively. The red curve in Fig. 2 represents the dependence for single crystal LiTbF$_4$ in the magnetic field $B||c$ [30]. The total magnetization would be reduced, because all possible particles directions in external magnetic field are equivalent in powder.

The temperature dependences of magnetic susceptibility for the sample 2 (black points) and sample 3 (blue points) are shown in Fig. 3. There are no clear evidence of transitions to a magnetically ordered state for TbF$_3$ or LiTbF$_4$. This may be due to the fact that the ordering of magnetic domains happens not simultaneously.

Fig. 4 shows the measured temperature dependence of the magnetic susceptibility of the powder samples 3 (TbF$_3$ powder). The temperature dependences of the magnetic susceptibility of single crystal TbF$_3$ demonstrate a strong anisotropy (the black points correspond to the $\chi||a$ direction, the pink points correspond to the $\chi||b$ direction, the green points correspond to the $\chi||c$ direction) [4].

The chemical composition of the synthesized samples is:

Sample 1 - LiTbF$_4$(79.9±0.7%)+TbF$_3$(20.0±0.5%),
Sample 2 - TbF$_3$ (81.2±0.4%) + LiTbF$_4$(6.9±0.1%)+LiNO$_3$ H$_2$O(11.9±1.1%),
Sample 3 - TbF$_3$(99±1%).

Particle size was preliminary determined using the dynamic and static light scattering spectrometer for registration nanoparticles size and composition Photocor Complex (table 1). Such big particles size probably can be explained by agglomeration of particles in the solution.
The magnetic susceptibility of the powder can be described by \( \chi_p = (\chi_a + \chi_b + \chi_c)/3 \), where \( \chi_a, \chi_b, \chi_c \) are magnetic susceptibilities along the axes \( a, b, c \) due to the symmetry of the crystal lattice. These calculations do not well fit the experimental results, because the size and shape of the particles were not taken into account. The crystal field and local magnetic field at the rare-earth ions located near the surface of nanoparticles are also different from ones in the bulk.

Theoretical calculation of the magnetic susceptibility of \( \text{LiTbF}_4 \) is shown on Fig. 5 (red line). Here all possible orientations of the external magnetic field along the crystallographic \( c \)-axis of the nanoparticles were taken into account. The shape of the nanoparticles assumed spherical, the summation carried on sphere \( 360^\circ \). Therefore the magnetization along \( c \)-axis [30]:

\[
M_z = B_z \left[ \lambda \left( \frac{\tanh(\delta/2k_BT)}{\tanh(\delta/2k_BT)} - 1 \right) + N \right]^{-1},
\]

where \( \lambda \) is molecular field constant (\( \lambda = 5.23 \pm 0.56 \)), \( \delta \) is initial splitting in the ground state quasidoublet (\( \delta = 1 \text{ cm}^{-1} \)) of basic \( ^7\text{F}_6 \) multiplet, \( N \) is demagnetization factor (\( N = 4\pi/3 \)), \( B_z \) is external magnetic field, \( T_c \) is the Curie temperature.

The measured temperature dependences of sample 1 and single crystal \( \text{LiTbF}_4 \) at \( B_0 || a \) and \( B_0 || c \) in external magnetic field (\( B_0 = 1 \text{ T} \)) are presented in the Fig. 5.

The curve 5 in Fig. 5 shows the calculations which take into account all possible crystallographic \( c \)-axis of the particle orientations with respect to external magnetic field. The discrepancy between the theoretical calculations and experimental data can also be explained by the fact that the calculation has not considered \( \text{TbF}_3 \) presence, nanoparticles shape and size.

Curve 4 (blue line) in Fig. 5 shows the calculations which take into account the following considerations: magnetic \( \text{Tb}^{3+} \) moments in \( \text{LiTbF}_4 \) single crystal are arranged along \( c \)-axis of the crystal lattice, the magnetization along the \( a \)-axis and \( b \)-axis is almost zero. Distribution of the \( c \)-axis orientations related to the direction of the external magnetic field is of the same probability. Thus, the magnetization along the \( a \) and \( b \)-axis can be neglected and the total magnetization can be estimated as \( M = M_z/3 \). The unusual behavior of the magnetization and magnetic susceptibilities in micro- and nano-sized powders \( \text{LiTbF}_4 \) and \( \text{TbF}_3 \) will be studied in detail later.
4. Conclusion

The samples of the rare-earth fluorides were synthesized. Crystallographic structure of the synthesized samples has been identified by the X-ray diffraction, magnetic properties of the powders mainly agree with the LiTbF$_4$ magnetic properties in the bulk. However improvement of the synthesis method is needed because the undesirable TbF$_3$ phase (∼20%) was found.

Acknowledgments

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References

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