

BOOK OF ABSTRACTS

JULY 15-20, 2018 MOSCONE CENTER icm2018sf.org









W3-09. Variation of magnetic properties of bilayered and trilayered NiFe and IrMn-based thin film structures. C. Gritsenko¹,
V. Rodionova¹, A. Berg¹, G. Babaytsev², I. Dzhun²,
N. Chechenin², M. Volochaev³ and A. Sokolov³ 1. Laboratory of Novel Magnetic Material, Immanuel Kant Baltic Federal University, Kaliningrad, Russian Federation; 2. Skobeltsyn Institute of Nuclear Physics, Lomonosov Moscow State University, Moscow, Russian Federation; 3. Kirensky Institute of Physics, Federal Research Center KSC SB RAS, Krasnovarsk, Russian Federation

FRIDAY Morning 10:00 SAN FRANCISCO BALLROOM

Session W4 MAGNETIC STRUCTURES AND MAGNETIC PHASE DIAGRAMS (Poster Session)

Alannah Hallas, Co-Chair Rice University, Houston, TX, United States Franziska Weickert, Co-Chair Florida State University, Los Alamos, NM, United States

W4-01. Critical properties of a generalized XY model with

competing nematic-like couplings. *M. Zukovic*¹ and G. Kalagov¹ *I. Institute of Physics, P. J. Šafárik University, Košice, Slovakia*

W4-02. Magnetic Transitions in the Heusler Compounds Fe_{3,x}Mn_xSi. T. Nonoyama¹, M. Hiroi², I. Shigeta¹, R. Kato¹, H. Manaka² and N. Terada² 1. Department of Physics and Astronomy, Graduate School of Science and Engineering, Kagoshima University, Kagoshima, Japan; 2. Department of Electrical and Electronics Engineering, Graduate School of Science and Engineering, Kagoshima University, Kagoshima, Japan

W4-03. Withdrawn

- W4-04. Novel Quantum and Thermal Phase Transitions of Easy-Axis Triangular Antiferromagnets in a Transverse Field. D. Yamamoto¹, G. Marmorini², M. Tabata¹ and I. Danshita³ 1. Department of Physics and Mathematics, Aoyama-Gakuin University, Sagamihara-shi, Kanagawa, Japan; 2. Research and Education Center for Natural Sciences, Keio University, Kanagawa, Japan; 3. Yukawa Institute for Theoretical Physics, Kyoto University, Kyoto, Japan
- W4-05. Non-equilibrium phase transitions in magnetic systems. A. Galda^{1,2} and V. Vinokur² 1. James Franck Institute, University of Chicago, Chicago, IL, United States; 2. Materials Science Division, Argonne National Laboratory, Lemont, IL, United States
- W4-06. Magnetic and electrical properties of R_sIr₃ (R = Tb, Er). K. Ueda¹ and T. Tsutaoka¹ I. Graduate School of Education, Hiroshima University, Higashi-Hiroshima, Japan

W4-07. Magnetic properties of Li⁵⁷Fe_{0.01}Mn_{0.99}PO₄ investigated by using external field Mössbauer spectroscopy. *H. Choi*¹ and C. Kim¹ I. Department of Physics, Kookmin University, Seoul, The Republic of Korea

- W4-08. Magnetic study on inverse spinel compounds Co₂XO₄ (X = Ti, Sn) with partial substitution of Zn for Co. H. Ohta¹, S. Takada¹, G. Ueda¹, K. Yamagishi¹ and H. Aruga Katori¹ I. Department of Applied Physics, Tokyo University of Agriculture and Technology, Koganei, Japan
- W4-09. Magnetic Property of Polymorphs (Al_xFe_{2,x})GeO₅. H. Aruga Katori¹ 1. Department of Applied Physics, Tokyo University of Agriculture and Technology, Koganei, Japan

W4-10. Withdrawn

W4-11. Origin of Magnetocrystalline Anisotropy in Trigonal Magnetics with Zero Orbital Moment. J. Kliava¹,
K. Seleznyova², M. Strugatsky², A. Drovosekov³, S. Yagupov² and V. Zubov⁴ I. LOMA, Université de Bordeaux, Talence, France; 2. Physics and Technology Institute, Crimean Federal University, Simferopol, Russian Federation; 3. P.L. Kapitza Institute for Physical Problems, RAS, Moscow, Russian Federation; 4. Lomonosov Moscow State University, Moscow, Russian Federation

FRIDAY MORNING 10:00

SAN FRANCISCO BALLROOM

Session W5 MULTIFERROICS III (Poster Session) Helen Walker, Chair

Helen Walker, Chair STFC, Didcot, United Kingdom

- W5-01. Spin waves in multiferroic Ni₃TeO₆. J. Lass^{1,2}, C.R. Andersen¹, J.O. Birk¹, H.K. Leerberg¹, S. Birkemose¹, S. Toth², U. Stuhr², M. Bartoviak², C. Niedermayer², Z. Lu³, R. Toft-Petersen⁴, M. Retuerto⁵ and K. Lefmann¹ 1. Niels Bohr Institute, University of Copenhagen, København Ø, Denmark; 2. Paul Scherrer Institut, Villigen, Switzerland; 3. Helmholtz-Zentrum Berlin, Berlin, Germany; 4. Physics, Technical University of Denmark, Lyngby, Denmark; 5. Instituto de Catálisis y Petroleoquímica, Consejo Superior de Investigaciones Cientificas, Madrid, Spain
- W5-02. Effect of the Eu³⁺ ion Substitution at the Nd site of the NdCrTiO₅: Structural, Magnetic, Electrical And Electronic Structure Studies. K. Gautam¹, A. Ahad², K. Dey¹, S. Majid², S.K. Sharma⁵, J. Coaquira³, S. Francoual⁴ and D. Shukla¹
 1. Material Science, UGC-DAE, CSR, Indore, India; 2. Physics, Aligarh Muslim University, Aligarh, ALIGARH, India;
 3. Laboratory of Magnetic Materials, University of Brasilia, NFA, Brazil; 4. Photon Science, Deutsches Elektronen-Synchrotron (DESY), Hamburg, Germany; 5. Physics, Universidade Federal do Maranhão, Sao Luis, Brazil

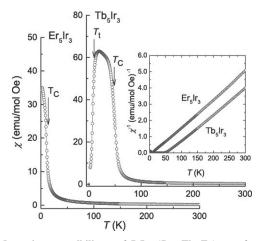
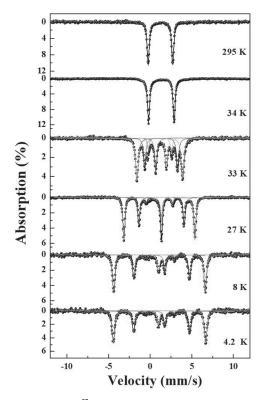


Fig.2 Magnetic susceptibility χ of R_5 Ir₃ (R = Tb, Er) as a function of temperature.

W4-07. Magnetic properties of Li⁵⁷Fe_{0.01}Mn_{0.99}PO₄ investigated by using external field Mössbauer spectroscopy. *H. Choi*¹ and C. Kim¹ *I. Department of Physics, Kookmin University, Seoul, The Republic of Korea*

The Li⁵⁷Fe_{0.01}Mn_{0.99}PO₄ polycrystalline sample has been studied by x-ray diffraction (XRD), vibrating sample magnetometer(VSM), and Mössbauer spectroscopy. The crystal structure is found to be orthorhombic with space group *Pmnb*. The lattice constants are a_0 =6.1009, b_0 =10.4435, and c_0 = 4.7427 Å. The magnetic properties of the sample were measured by using a VSM. The temperature dependences of ZFC and FC magnetizations at an applied field of 100 Oe. They show that magnetic Néel temperature (T_N) and spin-reorientation temperature (T_8) is 34 and 8 K, respectively. Mössbauer spectra of Li⁵⁷Fe_{0.01}Mn_{0.99}PO₄ have been taken at various temperatures ranging from 4.2 to 295 K. Magnetic hyperfine and quadrupole interaction in Li⁵⁷Fe_{0.01}Mn_{0.99}PO₄ at 4.2 K have been studied, yielding the following result; hyperfine field $H_{\rm hf}$ = 320.78 kOe, electric quadrupole splitting $\Delta E_{\rm Q} = 2.81$ mm/s, $\theta = 90^{\circ}$, $\phi = 0^{\circ}$, $\eta = 0.75$, and R = 1.2. Also, we were performed the Mössbuaer measurements with a high external field of 4.8 T. The value of magnetic hyperfine field (329.97 kOe) with external field of 4.8 T at 4.2 K was larger than that 320.78 kOe with zero magnetic field. While, the electric quadrupole splitting (2.62 mm/s) with external field 4.8 T at 4.2 K was smaller than that 2.81 mm/s with zero applied field. The abrupt increase in $H_{\rm hf}$ and magnetization was caused by orbital angular moment contribution from spin-orbit coupling.

Y. Deng, et al, Adv. Energy Mater. 7, 1601958 (2017).



Mössbauer spectra of $Li^{57}Fe_{0.01}Mn_{0.99}PO_4$ at various temperatures from 4.2 to 295 K

W4-08. Magnetic study on inverse spinel compounds Co₂XO₄ (X = Ti, Sn) with partial substitution of Zn for Co. *H. Ohta*¹, S. Takada¹, G. Ueda¹, K. Yamagishi¹ and H. Aruga Katori¹ *I. Department of Applied Physics, Tokyo University of Agriculture and Technology, Koganei, Japan*

Inverse spinels Co_2XO_4 (X = Ti, Sn) are known as compensated ferrimagnets with their ferrimagnetic transition temperature $T_{\rm N} = 48$ K and 41 K for X = Ti and Sn, respectively [1-3]. In their ferrimagnetic state, a spin-glasslike behavior is also observed. Thus, these compounds are categorized as the semi spin glass[3,4]. Although the magnetic structure in the ordered state has been revealed, the reason why the moments show both collinear ordering and spin glass freezing is still unclear. In Co₂XO₄, Co ions occupy both the whole of the A site and a half of the B site, and X ions occupy the rest of the B site. Thus, the B site, also known as the pyrochlore lattice, is just half-filled with magnetic ions. In this situation, spins are distributed onto the pyrochlore lattice at perfectly random, and this may lead to the spin glass. To understand the origin of semi spin glass of Co2XO4 and physics of the half-filled pyrochlore lattice, we studied Co₂XO₄ by partially substituting Zn for Co. Here, Zn preferentially replaces the A site rather than the B site. From the analysis results of the powder X-ray diffraction measurements, we confirm that Co atoms on the A site are selectively replaced with Zn atoms in both cases of X. After Zn atoms fully occupy the A site, the Co atoms on the B site are replaced with Zn atoms. The magnetic measurement results show that $T_{\rm N}$ of both compounds decreases with the increase of the ratio of Zn to Co, and their magnetism changes from ferrimagnetism to spin glass when all the Co atoms on the A site are replaced with Zn atoms. These results indicate that the pyrochlore lattice which is half-filled with Co atoms shows spin glass due to its randomness, and semi spin glass of Co2XO4 originates from this nature of the half-filled pyrochlore lattice.

 S. Thota, V. Narang, S. Nayak, S. Sambasivam, B. C. Choi, T. Sarkar, M. S. Andersson, R. Mathieu, and M. S. Seehra, J. Phys.:Condends. Matter 27 166001 (2015). [2] S. Nayak, S. Thota, D. C. Joshi, M. Krautz, A. Waske, A. Behler, J. Eckert, T. Sarkar, M. S. Andersson, R. Mathieu, V. Narang, and M. S. Seehra, Phys. Rev. B 92, 214434 (2015). [3] J. Hubsch, and