



Mössbauer study of

$Y_3AI_5O_{12}$ (YAG), $Y_3Fe_5O_{12}$ (YIG) and $Gd_3Ga_5O_{12}$ (GGG) single crystal garnets following dilute implantation of ⁵⁷Mn⁺.

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Motivation



Synthetic crystalline garnets: Yttrium aluminium garnet (YAG, $Y_3AI_5O_{12}$), Yttrium iron garnet (YIG, $Y_3Fe_5O_{12}$) and Gadolinium Gallium Garnet (GGG, $Gd_3Ga_5O_{12}$) are commonly used as a host materials in solid-state lasers, data storage, acoustic transmitters and numerous nonlinear optics applications (light-emitting diodes, scintillators, magneto-optical films etc.).

Despite the similar chemical structure the three garnets types posses different physical properties. In order to normalize our research we use only single crystal layers without additional dopant ions concentrations.

Yttrium Iron garnet (YIG, Y, Fe, O,)



Yttrium Aluminium garnet (YAG, Y₃Al₅O₁₂)



Gadolinium Gallium garnet (GGG, Gd₃Ga₅O₁₂)



Preliminary view

All the spectra in this presentation have been fitted with "Vinda" software. Spectra calibration have been provided with SS detector test. Velocity calibration gives emission temperature dependent scale relative to α-Fe.



$Y_3AI_5O_{12}$ (YAG)

DB 3, "**Voigt lines shape**": Specific of the this model doublet is the possibility two legs of the quadrupole are allowed to have different Gaussian broadening.

BT1 and **BT2**, "**Blume-Tjoin line-shape**" doublets. This model describe the ⁵⁷Fe emission Mössbauer relaxation spectra for the indicated relaxation times in a magnetic hyperfine field.



Spectra analysis YAG



DB3 associated with Fe³⁺ in YAG garnet could be assigned to the slow spin relaxation.



$Gd_3Ga_5O_{12}$ (GGG)

- **DB1** suggest Fe in highly distorted lattice environment.
- DB2 Probably DB2 is due to intersitial Fe.
- DB3 hyperfine parameters are consistent with an assignment to high-spin Fe³⁺.
- There is no annealing stage between RT and 563 K is observed.
- High-spin Fe³⁺ shows fast spin relaxation, presumably due to exchange interactions with Gd³⁺.



YAG spectra analysis



Quenching Spectra_YAG

								BT1 + BT2			
T (°C)	Corr.T.	Bck	δ (mm/s)	ΔEq (mm/s)	σs(ð) (mm/s)	σp(δ) (mm/s)	Area (%, mm/s)	Bscale	dSOD	р	Area
RT	300	0.505	-0.74898	1.7762229	0.396546534	0.505065461	-253.9839174	1	-0.238	0.251	-191.6144
LN ₂ RT	110	1.074	-0.82391	1.90131086	0.310070863	0.509220693	-226.0838609	1	-0.127	0.034	-215.757
3,2A	343.6	0.672	-0.74496	1.71105056	0.368821652	0.478184123	-244.2782615	1	-0.268	0.329	-200.6874
LN2_3,2A	110	1.258	-0.82631	1.91906488	0.315147475	0.534034122	-233.4107599	1	-0.127	0.034	-190.8429
6,0A	517.6	0.734	-0.65381	1.53633502	0.361109233	0.470246915	-192.3856868	1	-0.389	0.747	-183.4272
LN2_6,0A	110	0.236	-0.8325	1.98047189	0.3515473	0.51802749	-215.2768744	1	-0.127	0.034	-186.5385





514 K

414 K

298 K

11.11

8

Veločity (mm³/s)

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YAG: Data from Mössbauer spectra

parameters





Spectra analysis YIG



Complex Mössbauer spectra fiited with distribution two models DIST.1 and DIST.2 at temperature range 298 K ÷ 798 K. An additional two doublets have added been to accomplish the fitting analysis at 704 K and 798 K.



Spectra analysis YIG



Hyperfine fields (H_{in} , T), isomer shifts (IS, mm/s), quadrupole splitting (QS, mm/s), linewidths (W, mm/s), intensity ratio of magnetic peaks (PR = $P_{2.5}/P_{1.6}$), area ratio (16a/24d) of the octahedral (a) and tetrahedral (d) sublattices, and lattice constant (Å) of YIG films.

	Films	H _{in} (a) H _{in} (d) IS(a) IS(d)	QS(a)	QS(d)	W(a)	W(d)	PR(a)	PR (d)	16a/24d	Lattice C
1.	YIG	47.9 38.	7 0.32	0.16	0.02	- 0.01	0.42	0.56	1.06	1.20	0.53	12.3744
2.	$Bi_{0.2}Y_{2.8}Fe_5O_{12}$	49.2 39.	7 0.36	0.15	- 0.02	0.01	0.35	0.54	1.06	1.30	0.53	12.3802

K. Nomura et al. / Conversion electron Mössbauer spectroscopy of YIG Hyperfine Interactions 84(1994)421-426



YIG: data from Mössbauer spectra parameters



Summarize of the results

Summarize of the result for YAG

- Definite signs of an annealing stage with temperature range RT to 440 °C are not observed. Annealing at higher temperature is needed to incorporate all ions on regular lattice sites.
- High spin Fe³⁺ in this material shows slow spin relaxation presumably due to exchange interaction of Fe ions with Al ³⁺.

Summarize of the result for YIG

- Although limited to six ranges, the temperature dependence of the Isomer shifts and Quadrupole splitting's of the YIG garnet without impurity atoms concentration are consistent with the values for iron (III).
- The YIG single crystal garnet show clear temperature dependence at temperatures above the RT. Temperature range from 298K to 798 K is indicative for changes in crystal structure. Curie temperature for YIG is 560 K. After 604 K we observe cleary shrinking of the two line shapes distributions and predomination of the paramagnetic structure.

