



Jülich Centre for Neutron Science



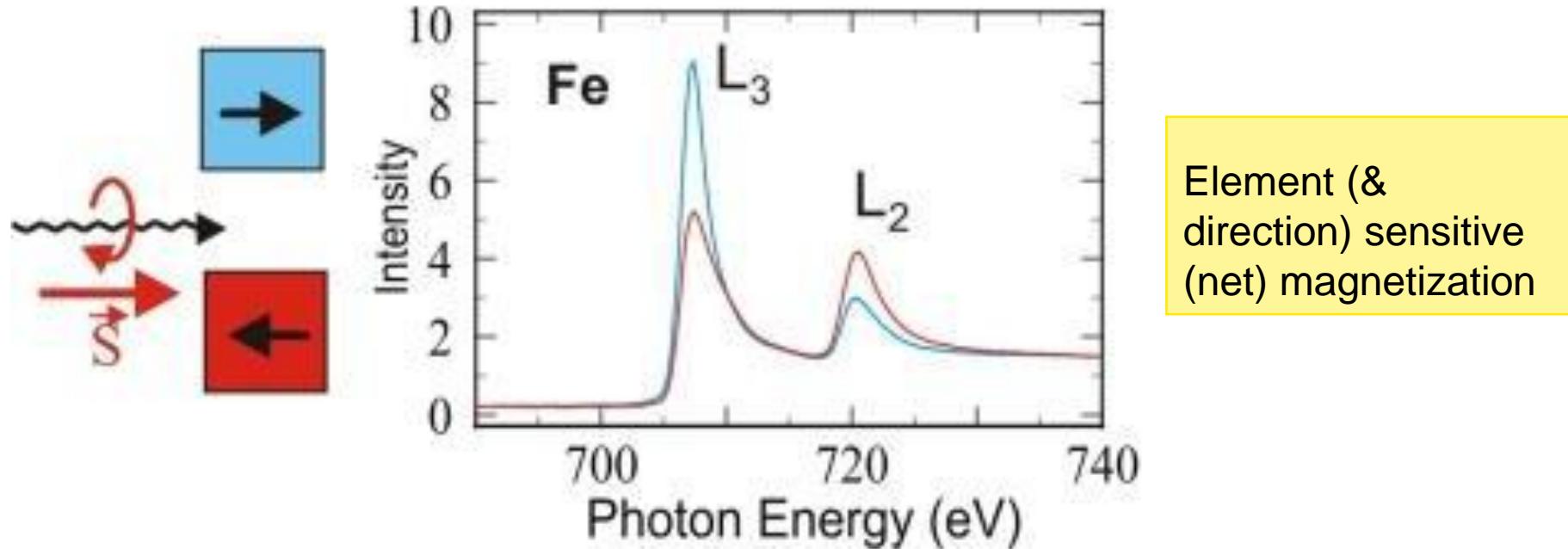
Soft X-ray absorption Study of Magnetic Materials

09.09.2022 | Sabreen Hammouda

s.hammouda@fz-juelich.de

X-ray Magnetic Circular Dichroism (XMCD)

Dichroism: dependence of absorption on the photon polarisation



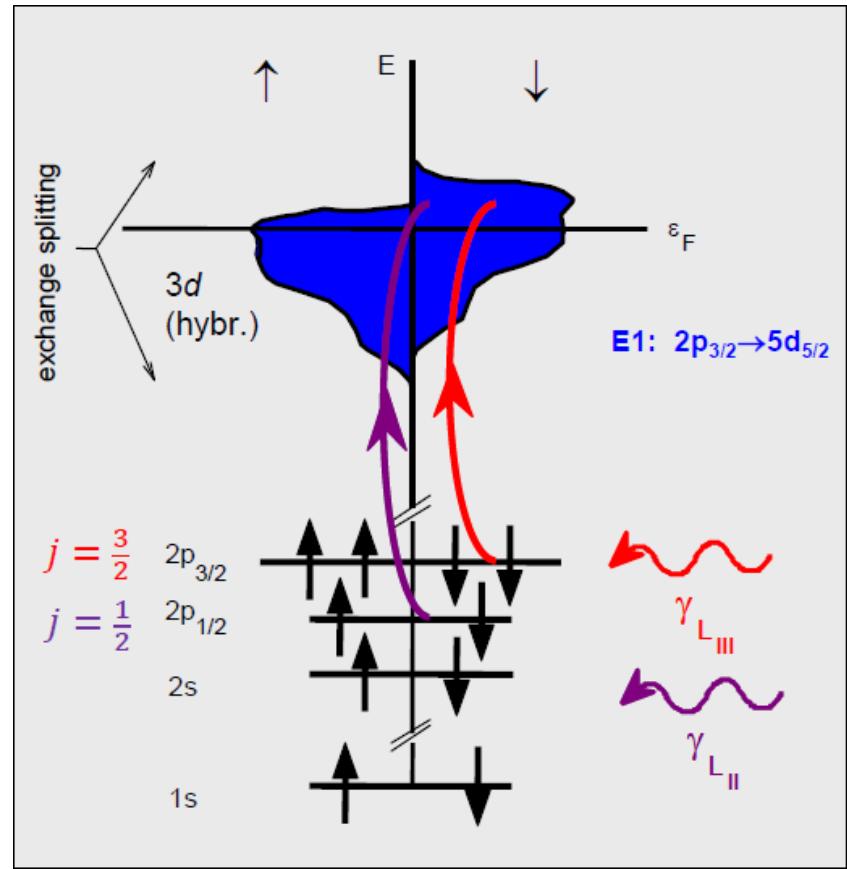
XMCD:

Absorption \propto available final states

Two-step model:

Transfer of the photon helicity (angular momentum) onto the photo electron – due to spin-orbit coupling this goes partially onto the spin.

2. Transition with dipole selection rules ($\Delta l = \pm 1, \Delta s = 0$) probes free final states for the spin direction of the photo electron.



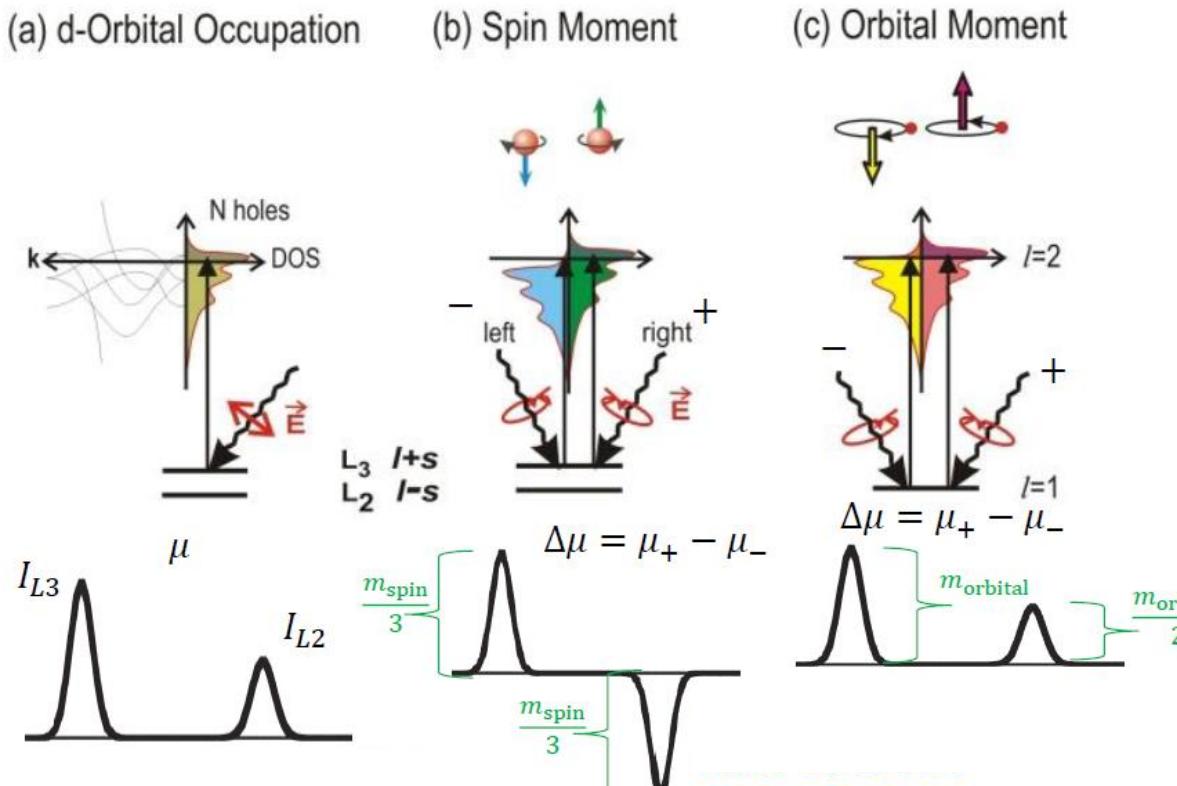
XMCD: sum rules

Also dependence on orbital moment (directly, helicity transferred to Δm_ℓ)

Opposite spin-orbit coupling in $p_{3/2}$ ($L3; j=\ell+s$) und $p_{1/2}$ ($L2; j=\ell-s$)

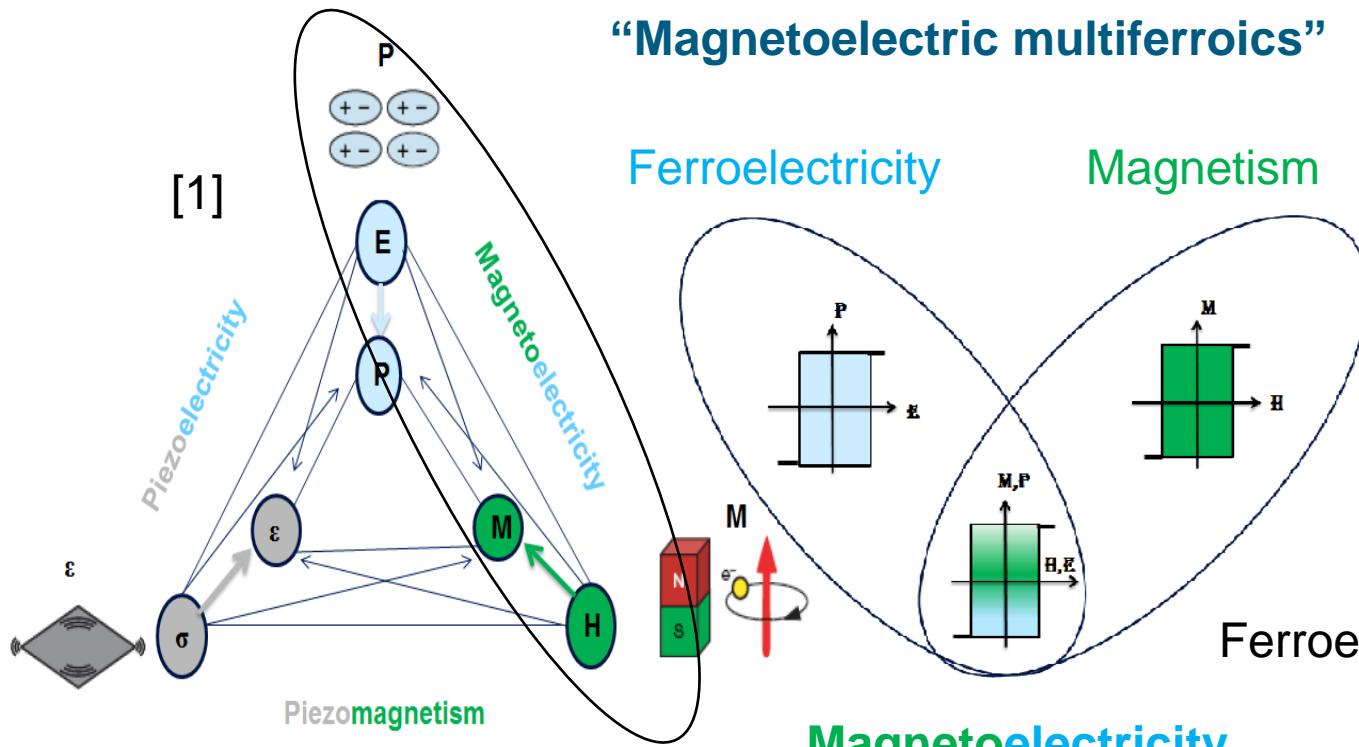


Spin- and orbital moments can be distinguished

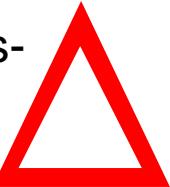


Multiferroics

“Magnetolectric multiferroics”



Multiferroics-
‘Contra-
indication’



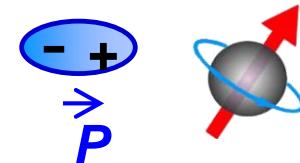
Ferromagnetism

Partially full d shell

Ferroelectricity

Empty d shell

Ferroelectricity: Charge (Dipoles)



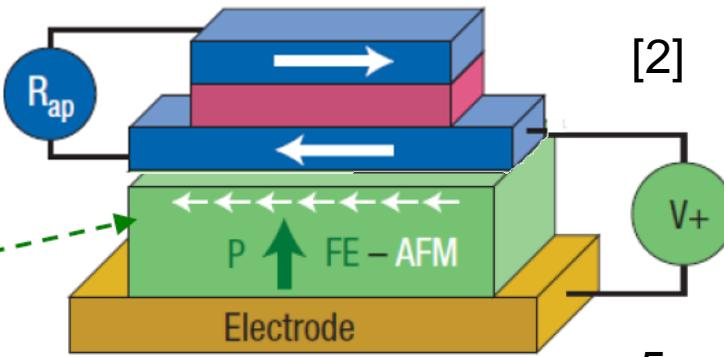
Magnetoelectricity

[1] Velev et al., Phil. Trans. R. Soc. (2011)

[2] Bibes and Barthelemy, Nat. Mater. (2008)

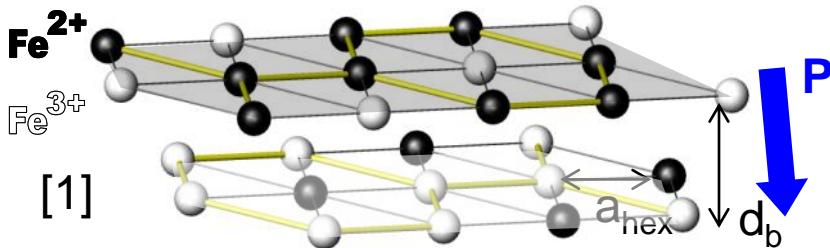
Multiferroic

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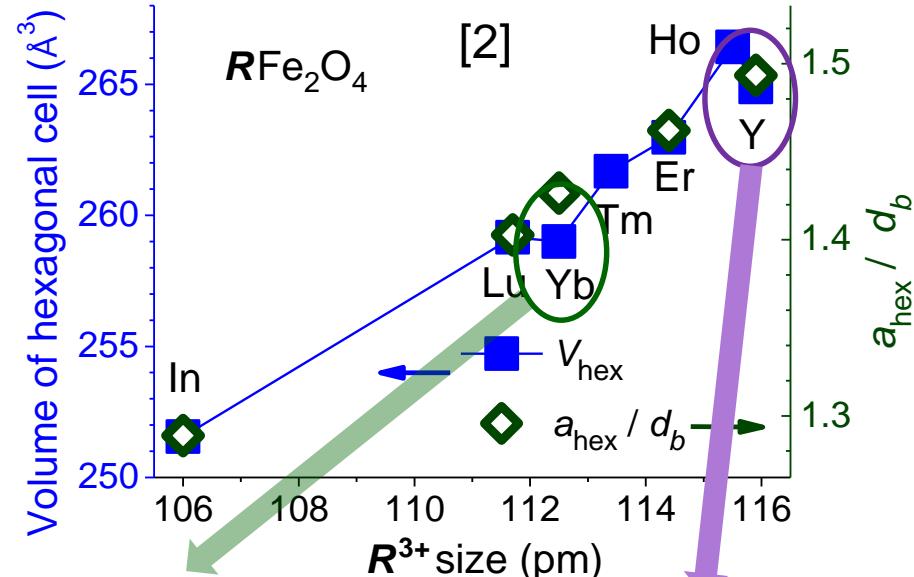


“Magnetic random-
access memory”

Ferroelectricity through charge order?



Bilayer subunit in $R\text{Fe}_2\text{O}_4$ compounds.



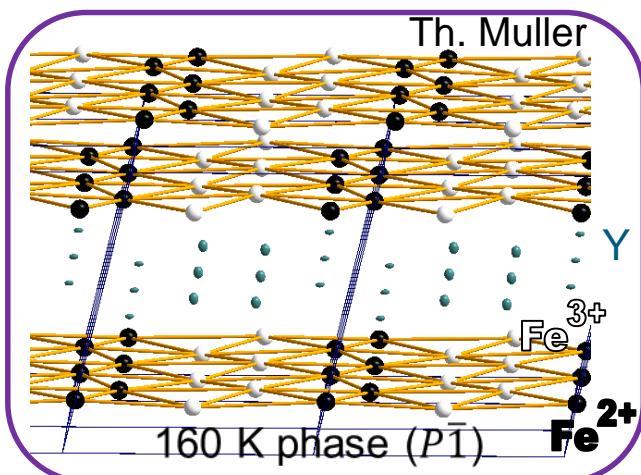
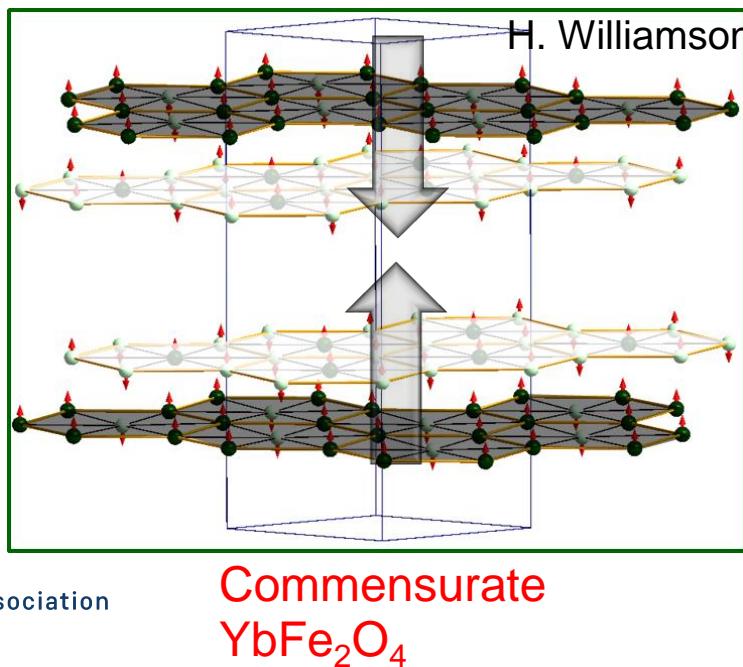
[1] Ikeda et al., J. Phys: Condens. Matter (2008)

[2] Angst, Phys. Status Solidi RRL (2013)

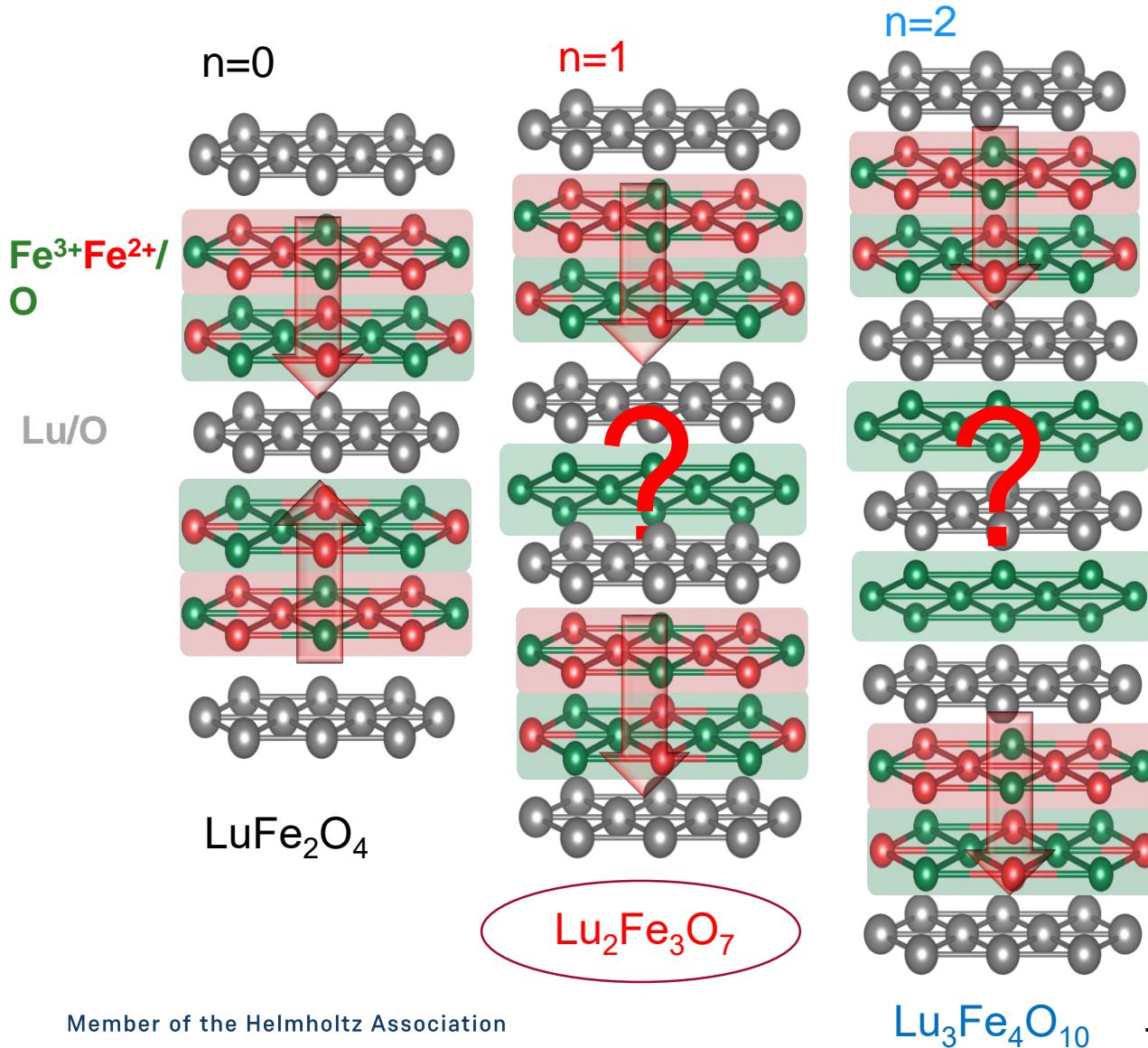
[3] Angst presentation

LuFe_2O_4
proposed to be a
magnetoelectric
multiferroic
through valence
ordering.

Fe_3O_4 is a proper
example.



Structure of $\text{LuFe}_2\text{O}_4(\text{LuFeO}_3)_n$ ($n=0,1,2$)

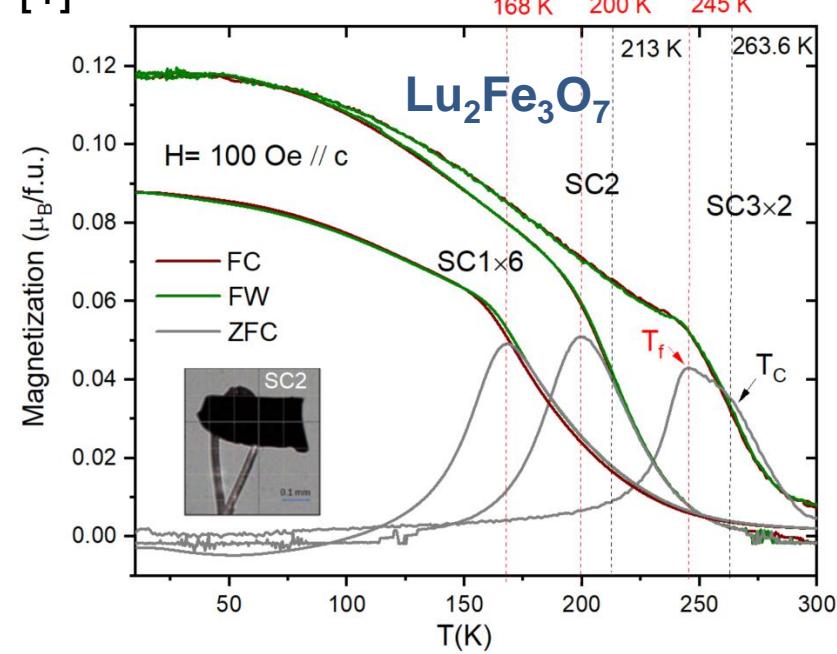


- Rhombohedral ($\text{R}\bar{3}\text{m}$, n even) [1]
- Hexagonal ($\text{P}6_3/\text{mmc}$, n odd) [2]
- Mössbauer spectroscopy studies : Fe-O mono-layer Fe^{3+} ions, while the bilayer contains $\text{Fe}^{2.5+}$ [3,4]

- [1] Matsui, J.Appl. Cryst. (1980)
[2] Kimizuka et al., Acta Crysta. (1976)
[2] Matsui et al., Acta Crysta. (1979)
[3] Tanaka et al., J. Magn. Magn. Mater. (1983)
[4] Tanaka and Iida, Hyperne Interact. (1994)

Magnetic properties vs stoichiometry

[1]



$0.4 * 0.20 * 0.05 \text{ mm}^3$

Crystal : $\text{CO}_2:\text{CO}$

SC1:50

SC2:85

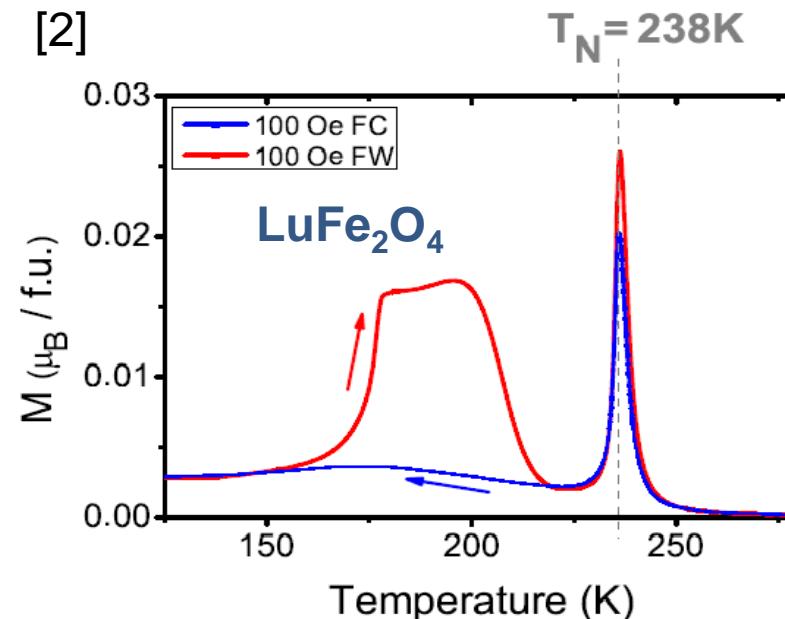
SC3:85

S1:90

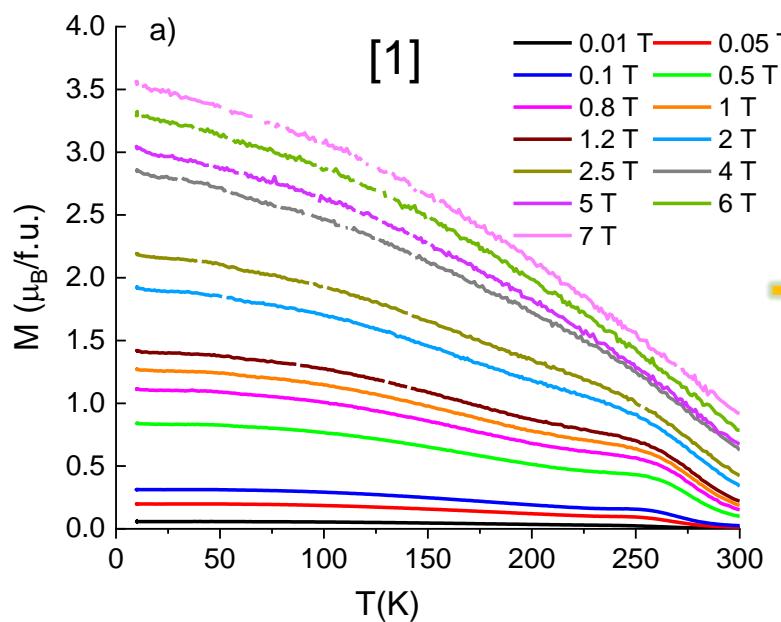
S2:85

- Best $M(T)$ curves still have no sharp features as observed for not intercalated compounds
- Variations even within one batch.
- Crystals with higher oxygen content are smaller

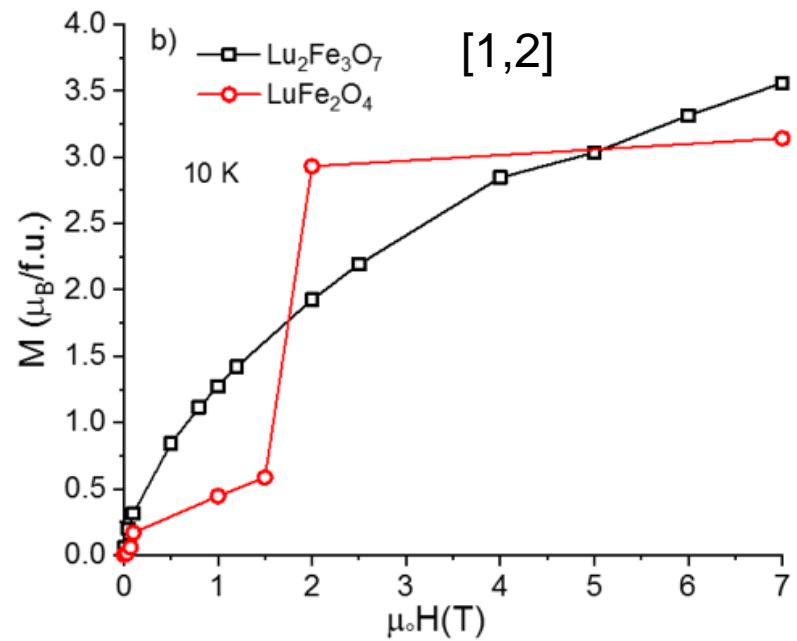
[2]



The high field studies on $\text{Lu}_2\text{Fe}_3\text{O}_7$ (SC3)



Field-cooled/warmed magnetization



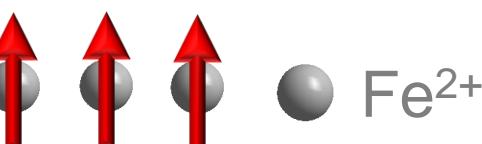
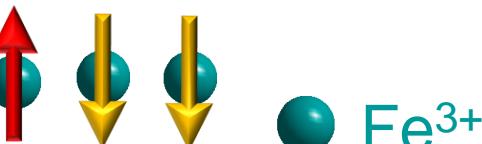
Low -T field cooled magnetization

- Moment of $\sim 3 \mu_B/\text{f.u.}$ arises from specific spin arrangement and indicates similar spin arrangement as in the bilayer of LuFe_2O_4
- An additional H-induced contribution is from the single layer spins.

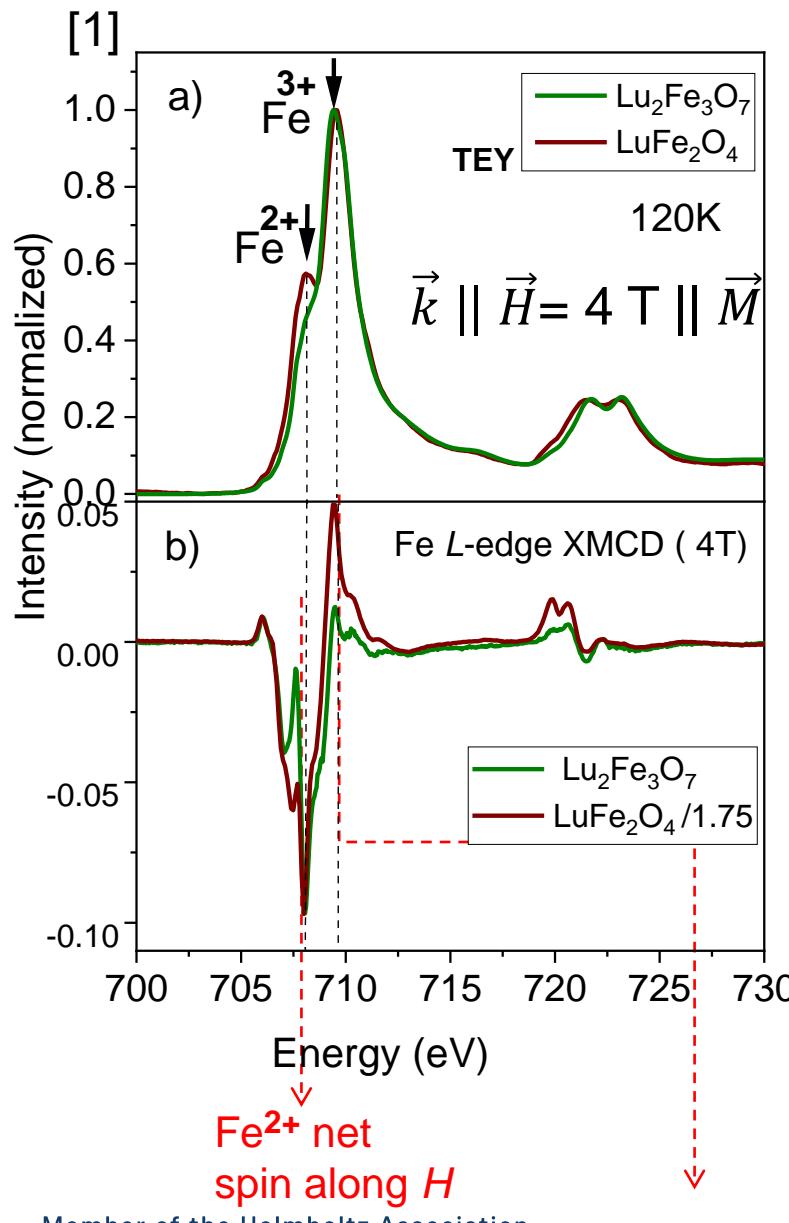
[1] Hammouda et al., submitted to PRB.

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[2] LuFe_2O_4 data from de Groot, Ph.D. thesis, RWTH Aachen university. (2012)



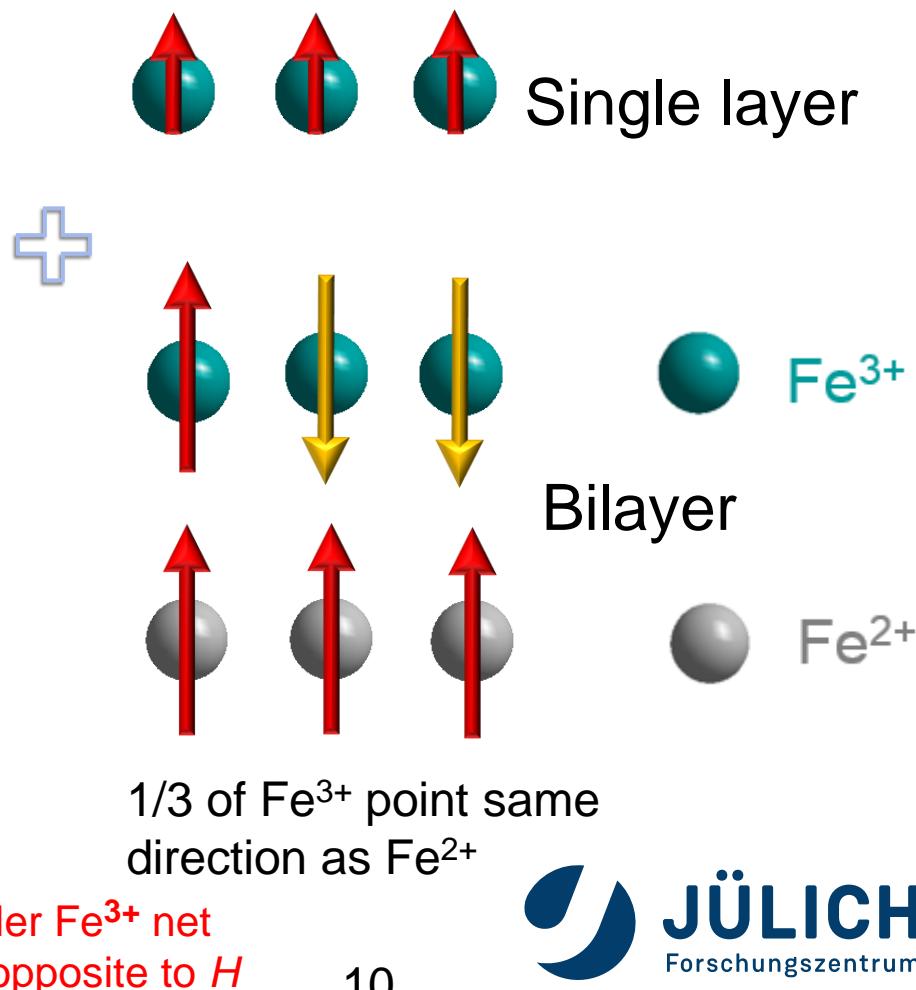
X-ray magnetic circular dichroism (XMCD) on SC3



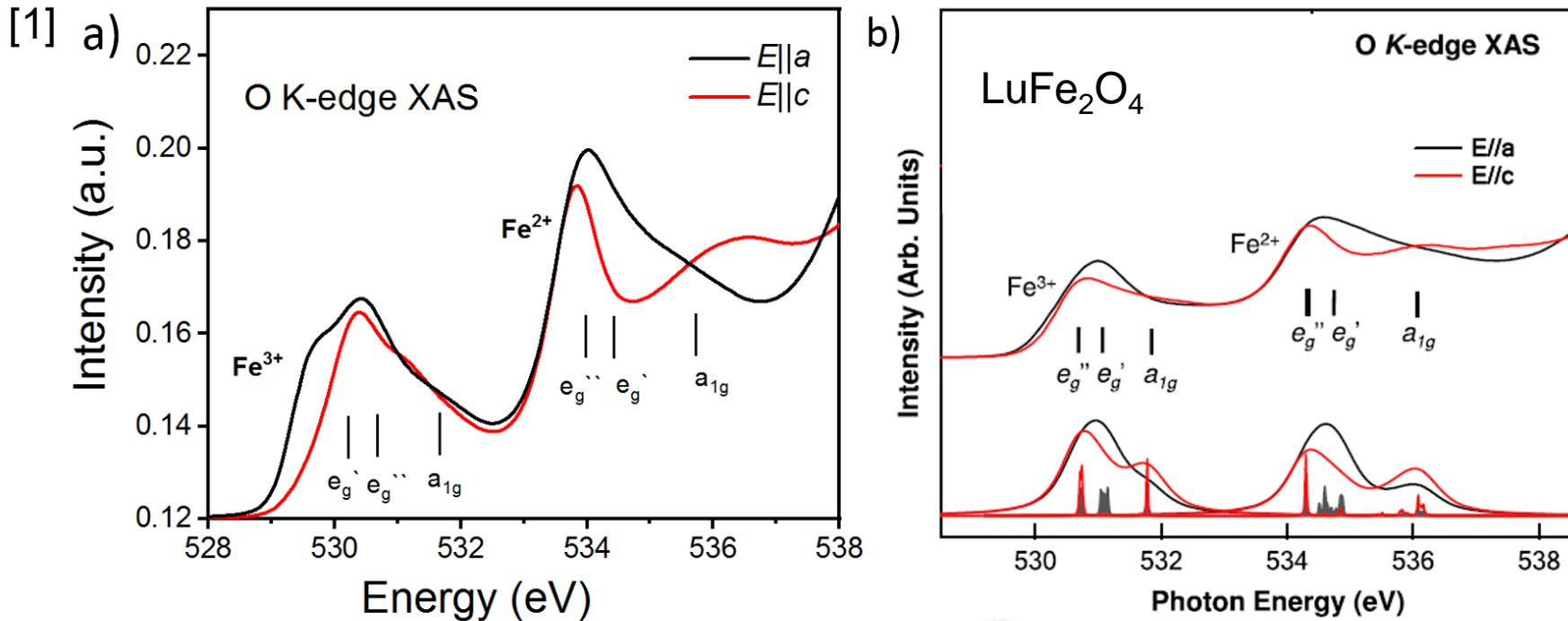
XMCD data probing Fe edge $L_{2/3}$

$$\Delta\mu = \mu_+ - \mu_-$$

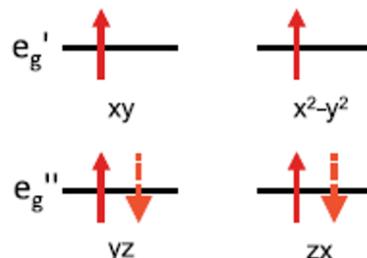
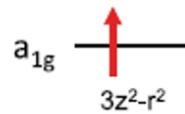
to deduce the net spins belonging to the different valances



Linear X-ray absorption spectra (LXAS) on SC3



[Ko et al., Phys.Rev.Lett. (2009)]



The super exchange interactions are very similar in the bilayer

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Summary

1. Single crystals Lu₂Fe₃O₇ exhibit no long-range spin order.
2. Similar spin order in the bilayer as in LuFe₂O₄:
 - comparable net moment to the bilayer of LuFe₂O₄
 - $\frac{1}{3} \frac{1}{3}$ in-plane propagation.
 - similar shape of the XMCD with net moment of the Fe²⁺ is the same
3. In the single layer, spins are paramagnetic-like in the first approximation.
4. Spin-charge coupling by XMCD, similar CO

Outlook

- Similar experiments will be done on quantum materials at HESEB.

Acknowledgements



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Thanks all who helped !

**Thanks for your
attention!**