Mössbauer Spectroscopy of Iron-based Chalcogenides

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Abstract-Soon after the discovery of superconductivity in LaFeAsO with $T_{\rm C} = 26$ K in 2008, many other iron-based superconductors were synthetized. They are all based on the layers which contain iron and a pnictogen (As, P) or a chalcogen (S, Se, Te) element. Due to the connection between superconductivity and magnetism these novel unconventional high-T_C superconductors have attracted tremendous interest in the scientific community. A particularly well studied is tetragonal FeSe in the PbO type structure (11 family). The improvement of $T_{\rm C}$ was achieved by the intercalation of an additional layer such as perovskite-like blocks or alkaline metals into the Fe-based chalcogenide lavered systems. This led to creation of new superconducting compounds, $A_y \text{Fe}_{2-x} \text{Se}_2$ (A is an alkaline element) named 122 family whose physical and structural properties are found to be very sensitive on details of composition. Unlike layered the chemical cuprate superconductors, a cationic disorder arisen from a substitution in an Fe-layer improves $T_{\rm C}$. The highest $T_{\rm C}$ in the Fe-based chalcogenide superconductors is accomplished by suppression of both long range crystallographic and magnetic order. Mössbauer spectroscopy is a very useful tool for studies of structural phase transitions, structure defects, and chemical and structural inhomogeneities. This lecture is devoted to the local structure studies of FeSexS1 - x, K0.7Na0.1Fe2Se2, KFe1 xCoxSe2, and similar Fe-based chalcogenide compounds using the Mössbauer spectroscopy.

Index Terms—Superconductors; iron-based chalcogenides; Mössbauer spectroscopy; local structures.

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