## **Investigating the Proximity Effect in h-BN/ 2D Magnet Heterostructures**

## <u>Katarzyna Ludwiczak</u><sup>1</sup>, Johannes Binder<sup>1</sup>, Aleksandra Krystyna Dąbrowska<sup>1</sup>, Piotr Tatarczak<sup>1</sup>, Joanna Sitnicka<sup>1</sup>, Jacek Jasiński<sup>2</sup>, Roman Stępniewski<sup>1</sup>, Andrzej Wysmołek<sup>1</sup>

<sup>1</sup>University of Warsaw, Pasteura 5, 02-093 Warsaw, Poland; <sup>2</sup>University of Louisville, 2301 S 3rd St, Louisville, KY 40292, United States of America

Hexagonal boron nitride (h-BN) is a two-dimensional crystal widely used in layered materials research due to its flat surface, high chemical stability, and lack of dangling bonds. Recent advances in the study of color centers in h-BN have made it an excellent platform for investigating basic physical phenomena, including optically addressable spin defects, rendering it a promising candidate for quantum technologies [1].

We focus on the growth of high quality h-BN samples using Metalorganic Vapour Phase Epitaxy (MOVPE) technique. The samples are grown on large ( $\sim$ 2 inches) sapphire substrates and cover them uniformly [2,3]. This method allows us to engineer material parameters such as its thickness, structure and consequently - type and concentration of defects. We observe emission from colour centres in epitaxially grown h-BN at different energies, indicating the presence of various defects, like boron vacancies or carbon-related defect complexes.

In this study, we explore the proximity effect between h-BN and 2D magnets (e.g. chromium trihalides). Such an approach enables the investigation of paramagnetic centres in h-BN using locally applied magnetic fields. We studied the photoluminescence of h-BN/CrBr<sub>3</sub>/h-BN heterostructures and found an additional peak in the spectrum at low (liquid helium) temperature. We believe that the optical response is correlated with the magnetic ordering in CrBr<sub>3</sub> crystals (Fig 1).

In the future, the creation of such heterostructures can become feasible using a subsequential growth of 2D magnets on top of the h-BN epitaxial layers [3]. Such an approach could be a reliable solution for scaling up ultrathin heterostructures and allowing their future development for real-life applications.

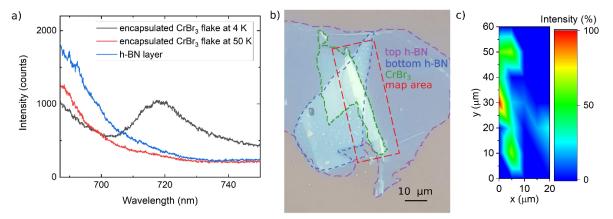


Fig. 1. a) Photoluminescence spectra of a h-BN/CrBr<sub>3</sub>/h-BN heterostructure. b) Optical microscope image of the studied heterostructure. c) Intensity (peak area) map of the additional photoluminescence peak of the studied heterostructure.

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