

High-Accuracy Ion Range Measurements using Fluorescent Nuclear Track Detectors



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INTRODUCTION

- Fluorescent Nuclear Track Detectors (FNTDs) (Fig. 1) are based on biocompatible single aluminium oxide crystals doped with carbon and magnesium (Al_2O_3 ; C, Mg).
- Their superior spatial resolution allows for **monitoring single particle tracks** with a detection efficiency close to 100% for ions with LET greater than approximately $0.2 \text{ keV}/\mu\text{m}$ [1].
- Implanted detectors or detectors in body cavities can help **accessing direct information on a radiation treatment** such as ion fluences, energies or ranges.
- We therefore measured ion ranges for particle beams of low and clinical fluence in order to investigate the **feasibility of future in-vivo FNTD applications**.

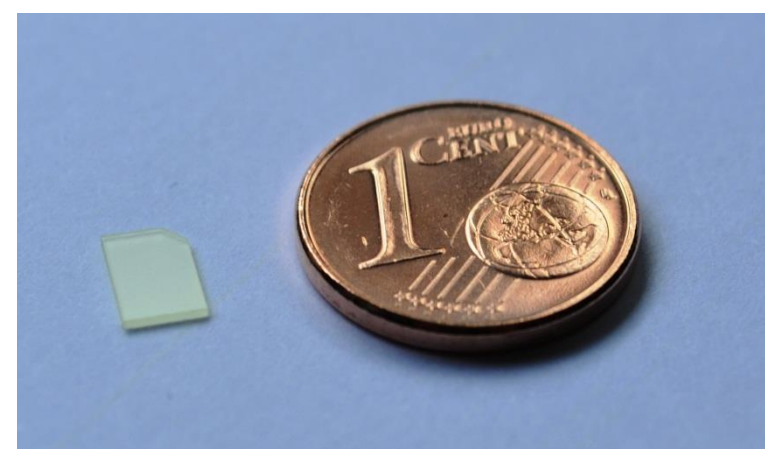


Fig. 1 Size of a FNTD compared to a one cent coin

FNTDs

- Because FNTDs are grown in a highly reduced atmosphere, they contain **high concentration of clustered oxygen vacancy defects** charge compensated by magnesium ions substituting aluminium ions in the crystal lattice (Fig. 2).

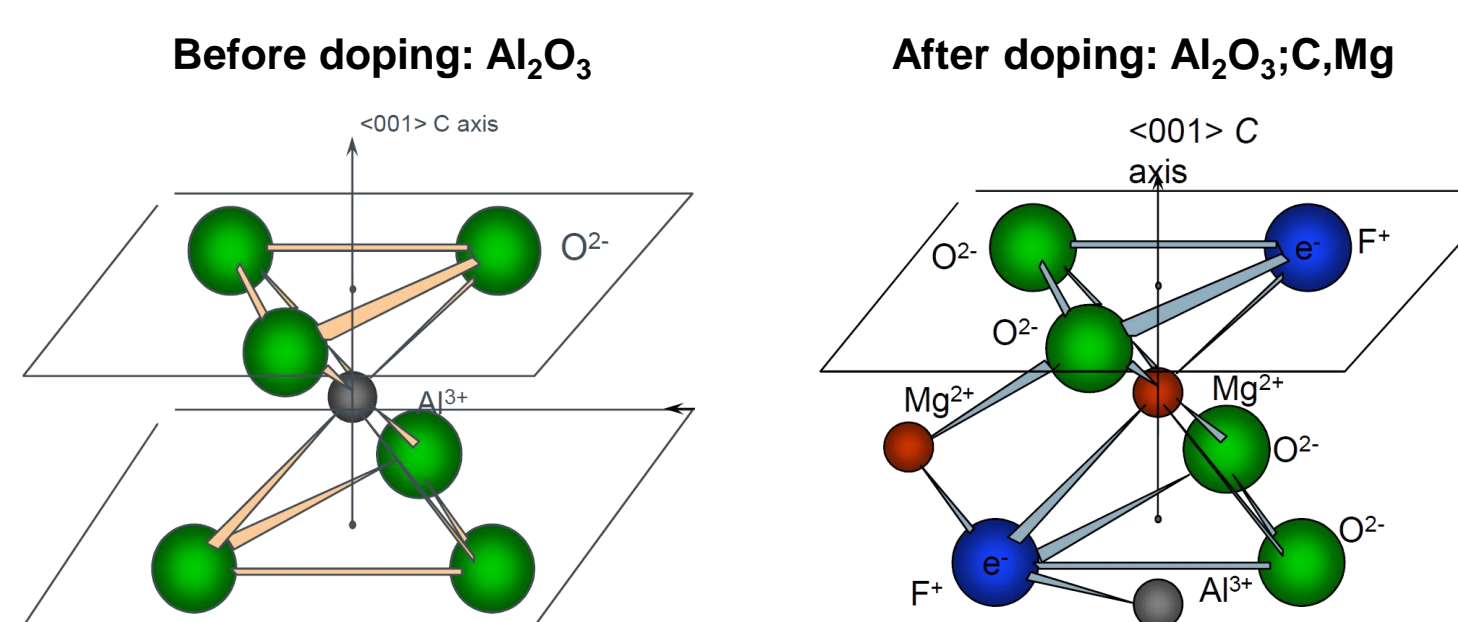


Fig. 2 Crystal structure of corundum (left) and model of an aggregate oxygen vacancy defect (right); image courtesy of M.S. Akselrod (Landauer Inc.) [2]

- Depicted F_2^{2+} (2Mg) colour centres undergo **radiochromic transformation** under ionising radiation (Fig. 3) by capturing secondary electrons: $\text{F}_2^{2+} (2\text{Mg}) + e^- \rightarrow \text{F}_2^+ (2\text{Mg})$.

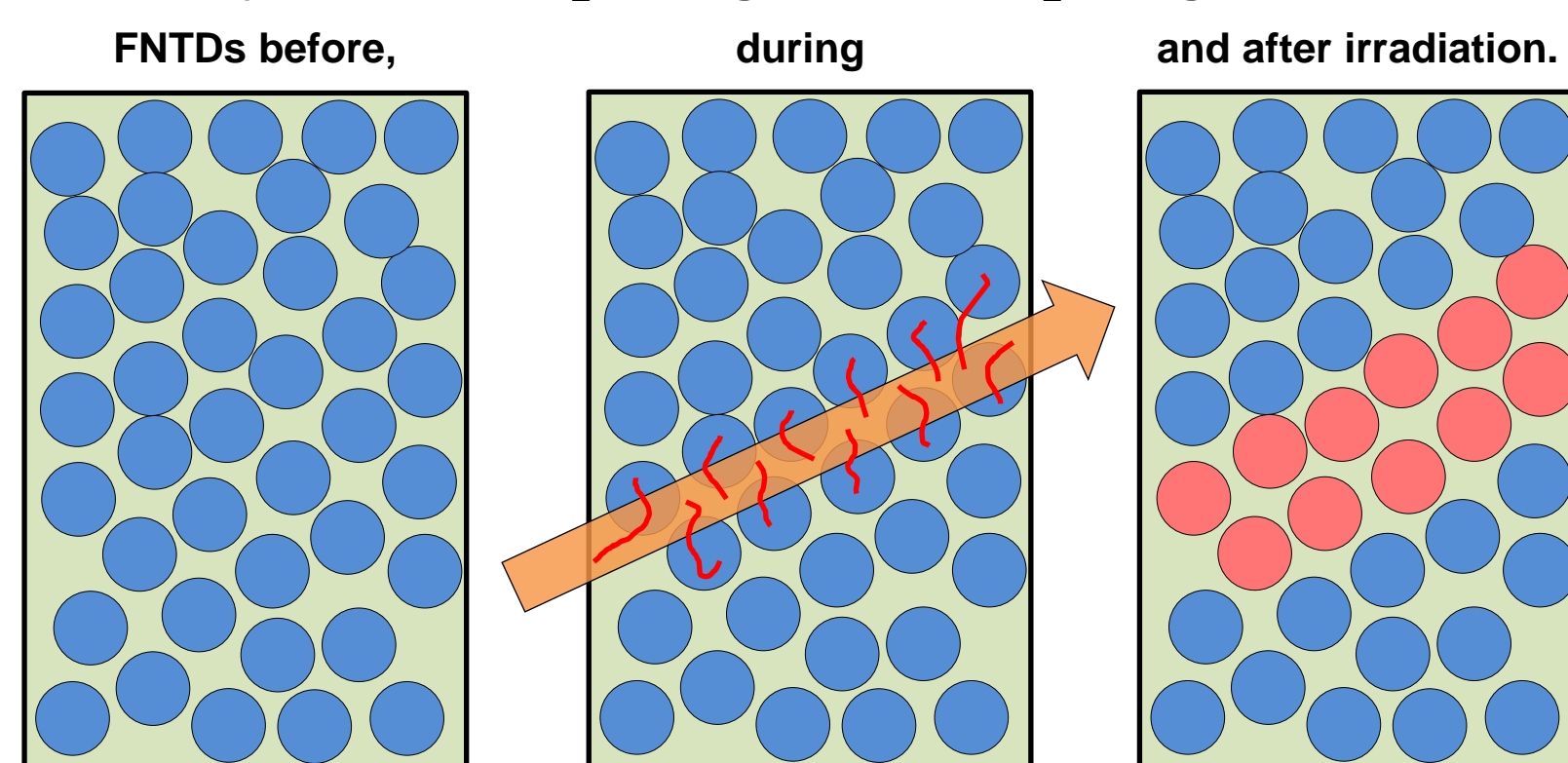
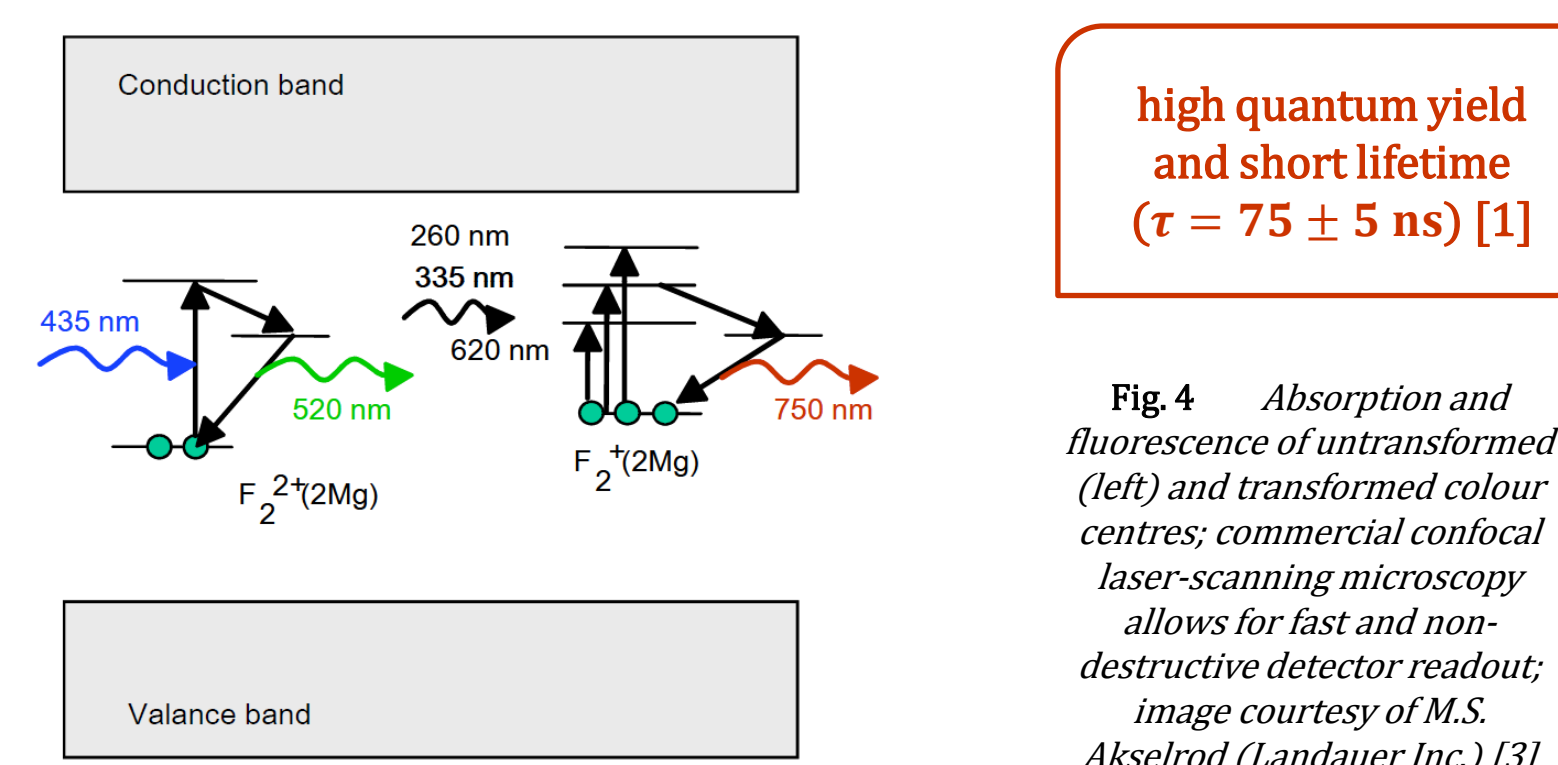


Fig. 3 FNTD containing only untransformed colour centres (left); liberation and capture of secondary electrons under ionising irradiation (middle); radiochromic transformation (right)

- Untransformed F_2^{2+} (2Mg) (light blue) and untransformed F_2^+ (2Mg) colour centres (light red) have different absorption and emission bands (Fig. 4) which can be stimulated using **confocal microscopy**.



high quantum yield
($\tau = 75 \pm 5 \text{ ns}$) [1]

EXPERIMENT 1: Range measurements

FNTDs were irradiated with **mono-energetic ions** using a broad range of particle types (hydrogen to xenon), kinetic energies, and particle fluences Φ (from 4.5×10^5 up to $1.0 \times 10^{11} \text{ cm}^{-2}$) (Fig. 5).

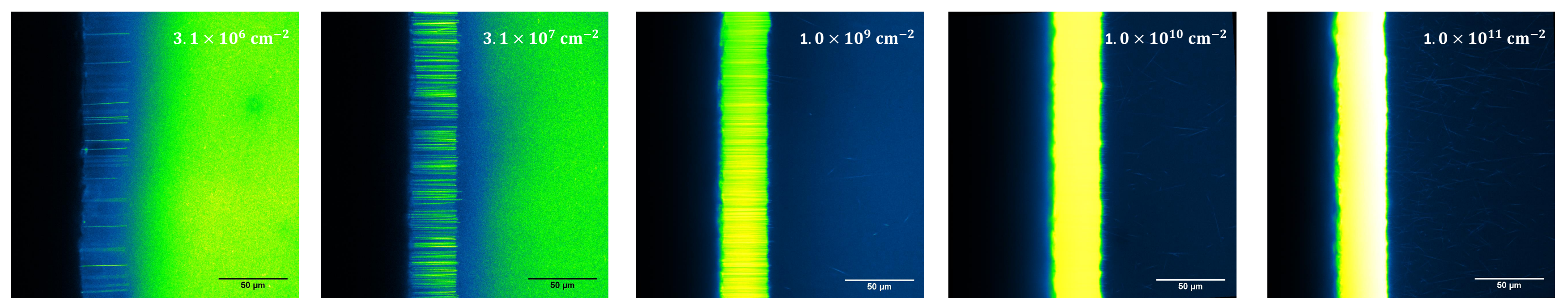
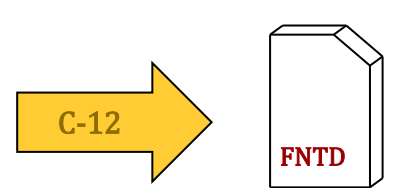


Fig. 5 Comparison of carbon irradiated FNTDs; theoretical SRIM range of 33.05 μm [4] at a total kinetic beam energy of 49 MeV

- Single track evaluation ($\Phi < 10^7 \text{ cm}^{-2}$):** determine entrance and end point individually for each particle track yielding projected range and lateral straggling; time-consuming but precise measurement routine
- Track bulk evaluation ($\Phi > 10^7 \text{ cm}^{-2}$):** determine inflection points in corresponding intensity profiles yielding projected range only; fast and automated measurement routine with the same level of precision (pinhole reduction advised for very high particle fluences)

All measured ranges deviate less than 3% from tabulated SRIM data [4] (Fig. 6). Measurement accuracy does not show any significant dependency on ion type, energy, fluence or LET.

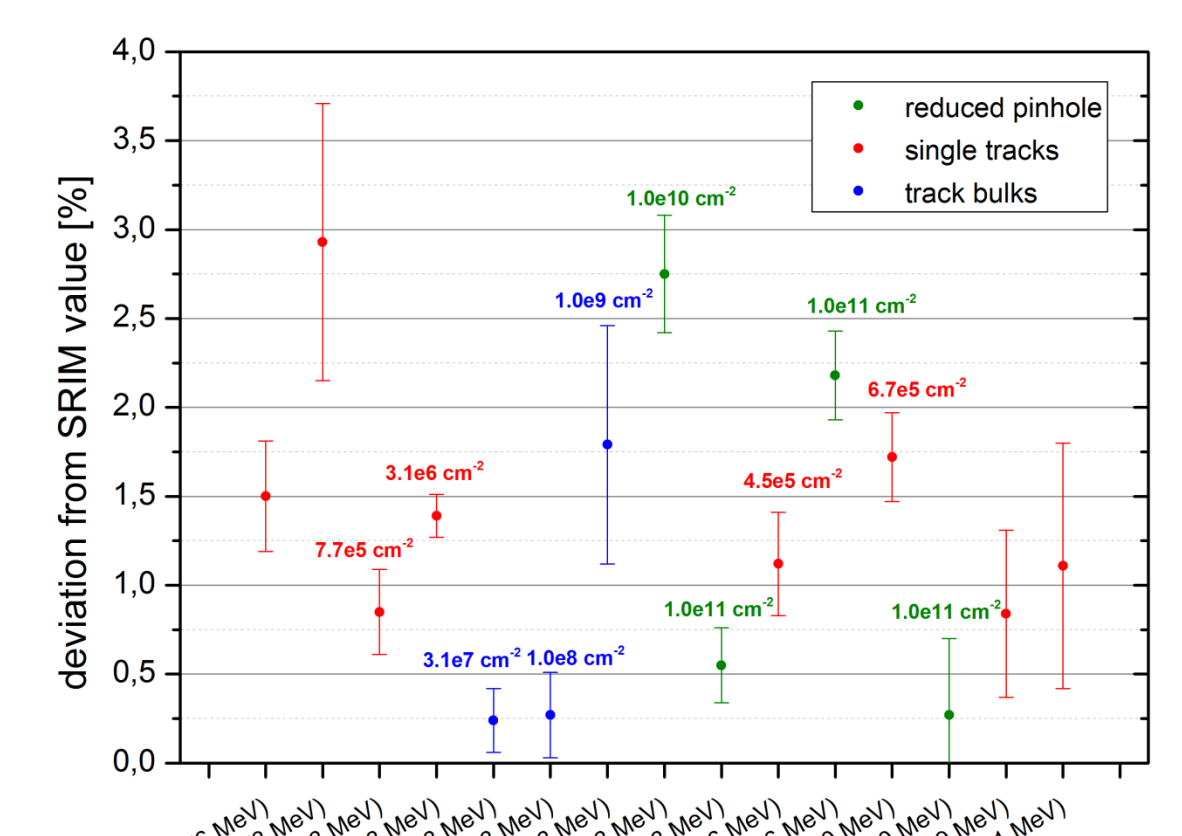
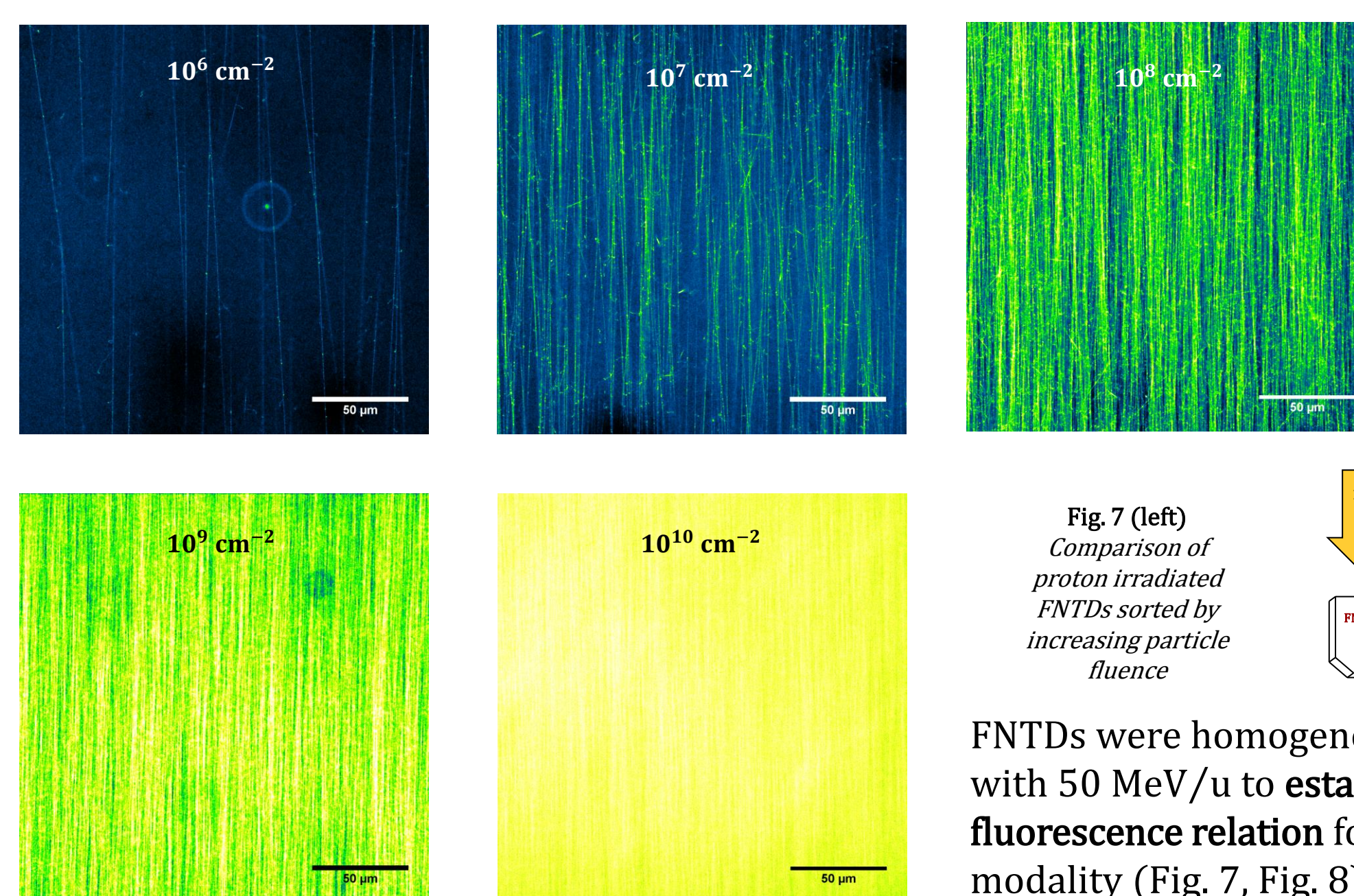
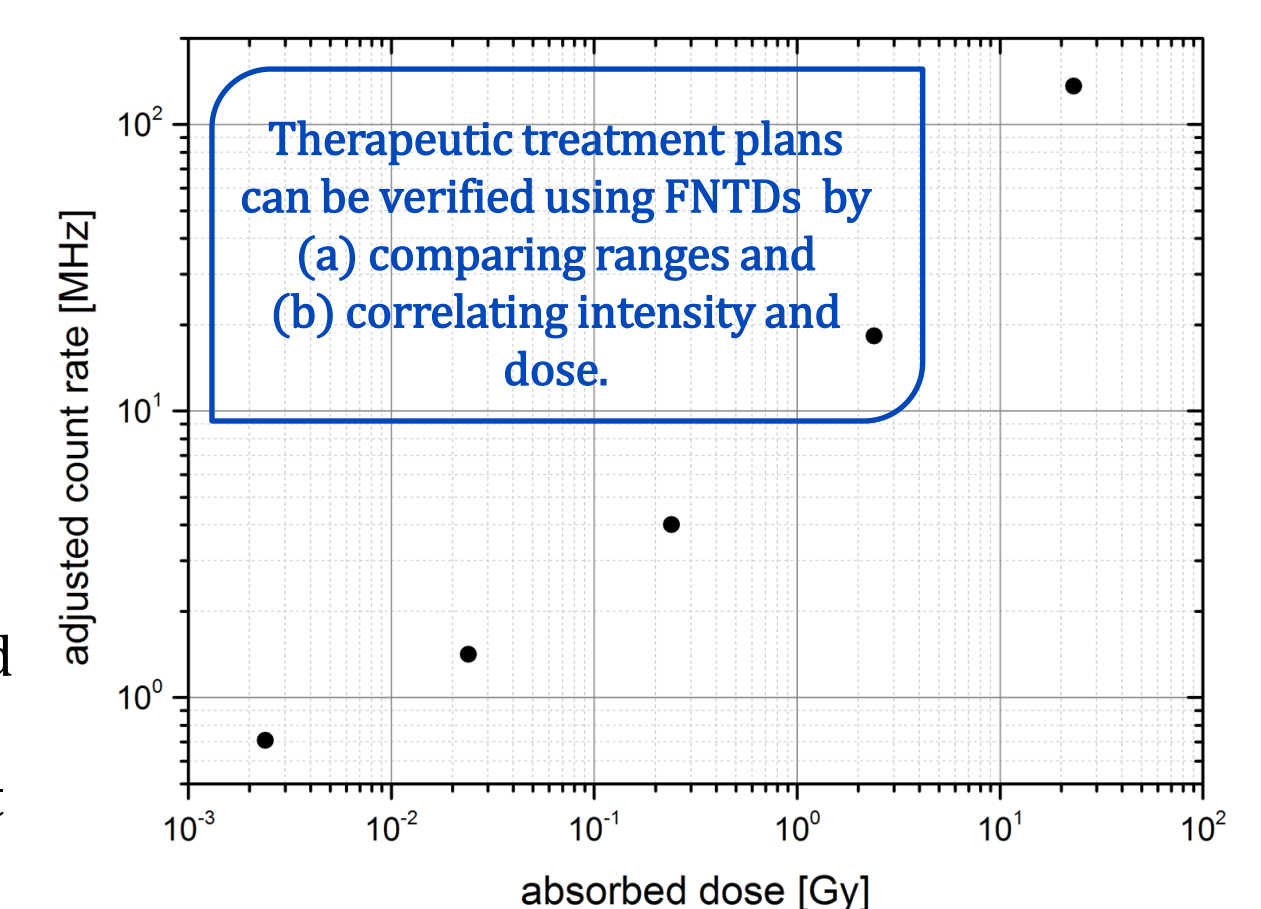


Fig. 6 Results of the high-accuracy ion range measurements for mono-energetic particle beams of various ion sources; all deviations from theory below 3%

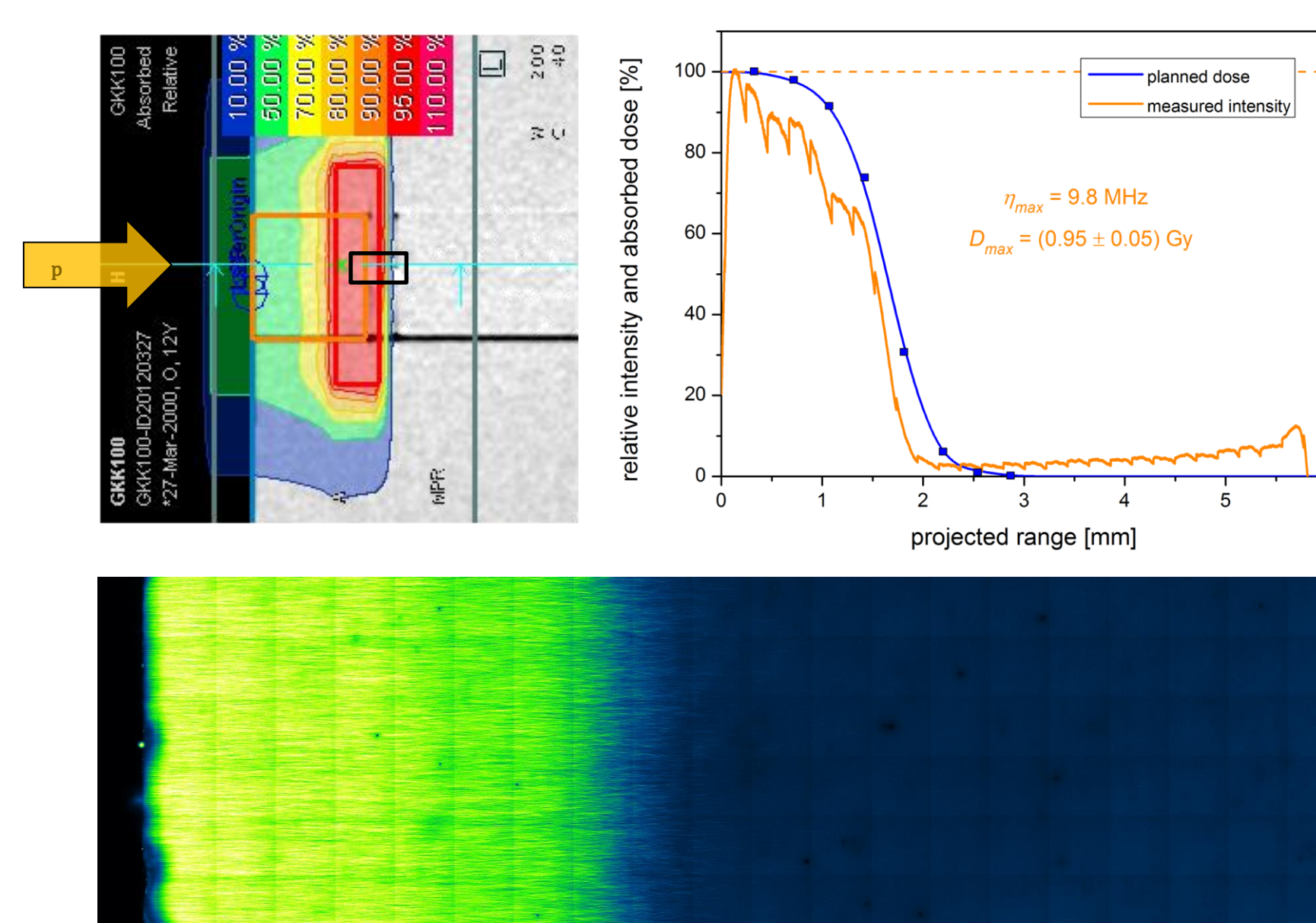
EXPERIMENT 2: Absorbed dose measurements



FNTDs were homogeneously irradiated with 50 MeV/u to establish a **dose-fluorescence relation** for this treatment modality (Fig. 7, Fig. 8).



EXPERIMENT 3: Treatment plan verification



FNTDs were placed in a PMMA cylinder as a phantom undergoing **clinical workflow**. Based on computed tomography (CT) scans, a treatment plan (applying 1 Gy protons at the detector edge) was created at HIT (Fig 9). The planned depth-dose curve at 80 % was compared to the detected intensity profile.

Detected and expected range agree within limiting CT slice thickness of 1 mm. Correlating fluorescence strength and absorbed dose verifies the maximal planned dose. FNTDs appear as potential fiducial marks in the CT.

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