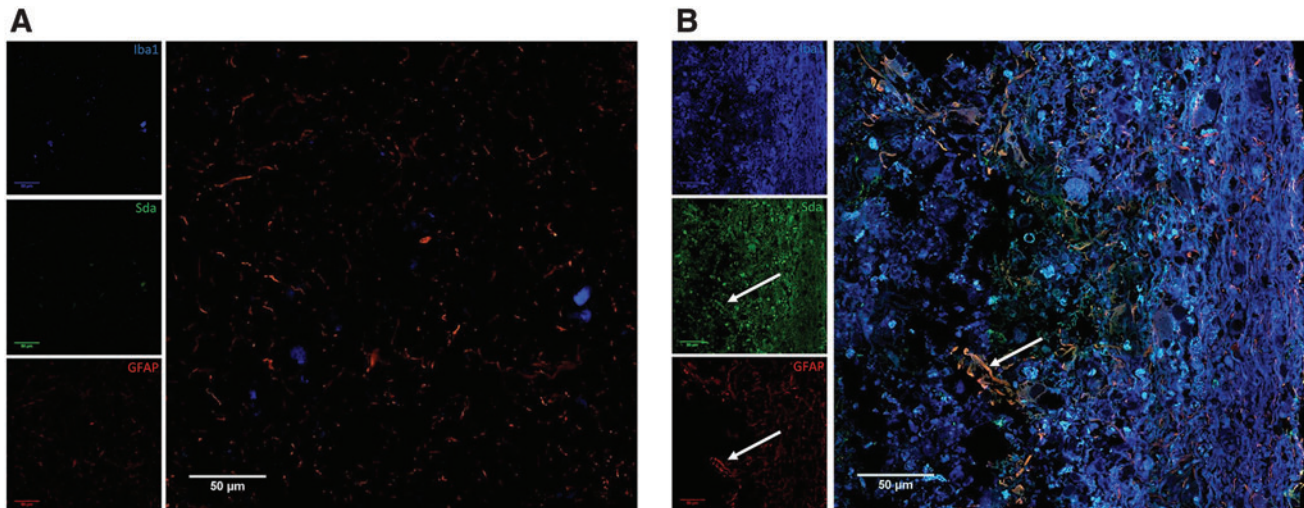


SUPPLEMENTARY FIG. S4. MALDI-TOF mass spectra of O-linked glycosylation from 3 DPI rat spinal cord sample. Per-methylated O-linked glycosylation derived from the 35% acetonitrile fraction (see Methods section). Indicated areas (m/z 1300–2300) in the spectra have a 10-fold magnification. All molecular ions are $[M+Na]^+$. Putative structures are based on composition, tandem MS, and biosynthetic knowledge. DPI, days post-injury; MALDI-TOF, matrix-assisted laser desorption ionization time-of-flight; MS, mass spectrometry.



SUPPLEMENTARY FIG. S5. Limited co-localization of Sda with GFAP-positive cells. (A) High-magnification image to show co-labeling Iba1 (blue), Sda (green), and GFAP (red) for sham. (B) High-magnification image to show co-labeling Iba1 (blue), Sda (green), and GFAP (red) for 14 DPI. Arrow points to a GFAP-positive cell with Sda. DPI, days post-injury; GFAP, glial fibrillary acidic protein; Iba1, anti-ionized calcium-binding adapter molecule 1.