Endogenous localization of TOP-2 in *C. elegans* using a C-terminal GFP-tag

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Abstract

To investigate the dynamic localization of Topoisomerase II in live *C. elegans* we have generated a C-terminally GFP-tagged version of TOP-2 at the endogenous locus. We found that TOP-2::GFP localizes in a similar pattern to the previously published TOP-2::3XFLAG strain and does not disrupt the meiotic chromosome segregation functions of this enzyme.

Figure 1. Endogenous TOP-2 localization using a GFP-tag: (A-F) Live imaging of the top-2::gfp *C. elegans* strain using confocal microscopy. (A) A complete representation of TOP-2 localization in the germ line of top-2::gfp worms. Scale bar= 50 µm. (B, C) TOP-2 localization in the head and tail, respectively, of top-2::gfp worms. Scale bar= 50 µm (D) TOP-2 localization during pachytene of top-2::gfp worms. White arrowheads point to examples of TOP-2::GFP foci. Scale bar= 5 µm. (E) TOP-2 localization in -1 and -2 oocytes in top-2::gfp worms. Scale bar= 5 µm. (F) TOP-2 localization in developing embryos in top-2::gfp worms. Scale bar= 50 µm. (G) Brood size comparison of endogenously tagged top-2::gfp to wild type (N2). (H) Percent embryonic viability from the top-2::gfp strain compared to wild type (N2). Statistics were conducted using a two-tailed Student’s T-test, n.s. indicates not significant, *p=0.0001.

Description

Topoisomerase II is a homodimeric, ATP-dependent enzyme that is conserved across species (Dong and Berger 2007; Nitiss 2009; Kim, Lee, and Koo 2000; Pommier et al. 2016; Vos et al. 2011). Topoisomerase II (Topo II) is expressed in all eukaryotic cells and is essential for many cellular processes including replication, transcription, chromosome structure, and chromosome segregation during mitosis (Nitiss 2009). In meiosis, Topo II is required for proper homologous chromosome segregation at the first meiotic division (Jaramillo-Lambert et al. 2016;
Generation of C-terminally tagged TOP-2::GFP via CRISPR-Cas9: A C-terminally meGFP-tagged version of TOP-2 (top-2::gfp) was created via the direct delivery method (Paix et al. 2017) using dpy-10 as a co-CRISPR marker (Arribere et al. 2014). The injection mix contained Cas9 protein (10 mg), dpy-10 CRISPR RNA (crRNA) (3.2 mg), dpy-10 (crRNA) repair oligonucleotide (0.2 mg), universal trans-activating crRNA (tracrRNA) (20 mg, Dharmacon, GE Life Sciences), and a crRNA targeting the c-terminal sequence of top-2 (GACGCGUCGUCGACUCCGACUGUUUUAGAGCUAUGCUGUUUG, 8 mg). The repair template was generated by PCR using primers containing meGFP (pAP969, Paix et al. 2017) and top-2 forward (5’-TTACGACGTGAGCTGACAGATCCAGATCCAGATCCAGAACAGAGAGAGACGGCGCTGT CGACCTCGGACTCAGGATTCAAGGAGAGAGCTTTCCACGCGG3’) and top-2 reverse sequences (5’-GGAAAGAAGAATAATATGAAACATCTAAGGAGGTTGGAAACAAGAATTATTTACTTGAGACGCTCCATCGAGG3’) and a reverse primer (5’-GTCTCCCTTGGATGTGCAGCGGTGAGGTCCTATCGG3’). The crRNA was designed to have a low possibility of off-target sites using http://crispr.mit.edu. Two independent strains were created (AG330 and AG332). Strain AG332 was the focus of this study.

Live imaging was conducted by placing 10 worms on a slide with a fresh 2% agarose pad and 10 µl of anesthetic (0.2 mM tetramisole in M9 buffer). A glass coverslip was then placed on top of the worms and agarose pad and the space between the agarose pad and the edges of the coverslip were filled with pipetting M9 under the coverslip. Live images were taken on a Zeiss LSM880 confocal microscope using either the 40x (Figure 1A, B, C, & F) or the 63x (Figure 1D & E) objectives. Image processing and analysis was done on Fiji Is Just ImageJ (Schindelin et al. 2012).

Embryonic viability assays for the wild type (N2) and top-2 (av87) (top-2::gfp) strains were conducted at 20°C. Both strains were grown on MYOB petri dishes inoculated with the E. coli (OP50). Ten L4 hermaphrodite larvae of each strain were put onto separate plates and allowed to grow and lay eggs for 16-24 h (one worm per plate). After each 24 h period, the adult (parent) worm was transferred onto a fresh plate until no additional embryos were produced. The total number of progeny (larvae and dead embryos) were counted one day after the parent was transferred. Three independent trials were conducted for each strain.

Brood sizes were determined by placing a single L4 hermaphrodite for both strains [N2 and top-2 (av87) (top-2::gfp)] on a 35 mm MYOB plate spotted with E. coli (OP50). Each hermaphrodite was allowed to lay embryos for 24 h at 20°C. After 24 h the hermaphrodite was transferred to a fresh plate until no more embryos were produced (two additional transfers). Total brood size was calculated by adding together live larvae and dead embryos produced from a single hermaphrodite.

Reagents
AG332 top-2 (av87) (top-2::gfp) is available from the Jaramillo-Lambert Lab.
N2 is available from the CGC.

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