Human Topoisomerase II beta Microplate Assay

Product Description (Product Number TRTB201 and TRT202)

The kit is supplied with sufficient human topo II beta enzyme , plasmid DNA substrate (supercoiled pNO1; supplied at 1 mg/ml), 10X Assay Buffer, ATP, Enzyme Dilution Buffer, DNA-detection dye (Promega Diamond dye) and TFO1 oligo for 100 assays. The enzyme is supplied at a concentration of 5-10 U/µl in Dilution Buffer (see Note 1). The kit is also supplied with sufficient Wash Buffer, TF buffer and T10 buffer for one 96-well plate. These buffers are supplied as 20X concentrates and must be diluted prior to use with ultra-pure water.

Store at -80 °C. (Stable for 3 months undiluted.) The enzyme should be centrifuged briefly before opening and it is recommended that it is aliquoted (undiluted) to avoid repeated freeze-thaw cycles.

For *in vitro* laboratory research use only.

Dilution Buffer

50 mM Tris.HCl (pH 7.5)
100 mM NaCl
1 mM DTT
0.5 mM EDTA
50 % (v/v) glycerol
50 µg/ml albumin

Assay Buffer
(supplied as 10X stock)

50 mM Tris.HCl (pH7.5)
125 mM NaCl
10 mM $MgCl_2$
5 mM DTT
100 µg/ml albumin

ATP (supplied as 30X stock)

30 mM ATP

TF Buffer
(supplied as a 20X stock)

50 mM sodium acetate (pH 5.0)
50 mM NaCl
50 mM $MgCl_2$

Wash Buffer
(supplied as a 20X stock)

20 mM Tris.HCl (pH 7.6)
137 mM NaCl
0.005 % (w/v) BSA
0.05 % (v/v) Tween-20

DNA-detection Dye
(supplied as a 1,000X stock)

Supplied in 100% DMSO

T10 Buffer
(supplied as a 20X stock)

10 mM Tris-HCl (pH 8)
1 mM EDTA

Preparation of Plate and Relaxation Assay

Rehydrate wells with 3 x 200 µl Wash Buffer (diluted from 20X stock before use).

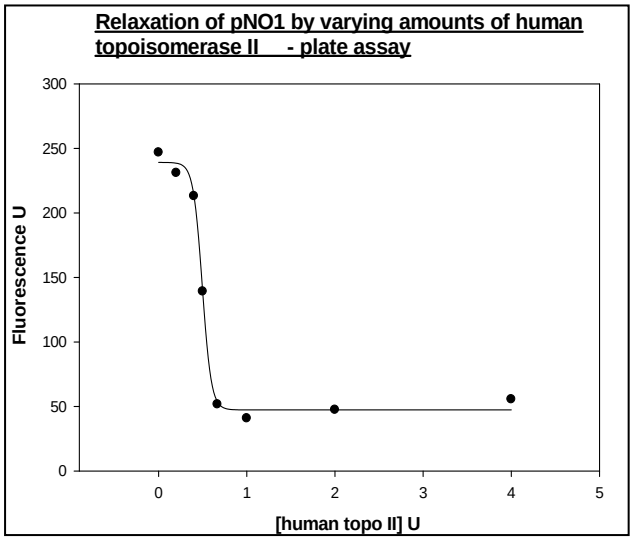
Immobilize 100 µl of 500nM TFO1 oligo in well (5 µl of 10 µM TFO1 in 95 µl Wash Buffer), 5 minutes at room temperature. Wash off excess oligo with 3 x 200 µl Wash Buffer.

Incubate 1.5 U of human topo II with 0.75 µg of supercoiled pNO1 in a reaction volume of 30 µl at 37°C for 30 minutes in Assay Buffer. Incubate reaction in well.

Add 100 µl TF Buffer (diluted from 20X stock before use) to well and incubate for a further 30 minutes at room temperature to allow triplex formation.

Remove liquid from well and wash with 3 x 200 µl TF Buffer to remove unbound plasmid.

Stain with DNA-detection dye (diluted to 1X with T10 buffer. Add 200 µl per well. Incubate for 10 - 20 minutes, **mix** and read in fluorescence plate reader; Ex: 495 nm; Em: 537 nM).



Quality Control

1) Purity: Human topoisomerase II is purified to > 95 % purity as judged by SDS-polyacrylamide gel electrophoresis. 2) Tests for human topoisomerase I contamination by looking for relaxation of supercoiled pBR322 under topoisomerase I assay conditions were negative. 3) kDNA or pBR322 were also incubated for 4 hrs in assay buffer + 10 mM $MgCl_2$ at 37 °C. These tests were negative for the formation of linear products, indicating the absence of nuclease contamination.

References

Maxwell, A., Burton, N.P. and O'Hagan, N. (2006). High-throughput assays for DNA gyrase and other topoisomerases. *Nucleic Acid Res*. **34(15)**, e104

Notes.

1. A unit of enzyme activity is defined as the amount of enzyme that just supercoils 0.5 μg of pNO1 in 30 minutes at 37ºC as judged by agarose gel electrophoresis.
2. It is important to mix the samples before reading or they will be greatly reduced.