SANTA CRUZ BIOTECHNOLOGY, INC.

FANCC (C-14): sc-18110



BACKGROUND

Fanconi anemia (FA) is an autosomal recessive disorder characterized by bone marrow failure, birth defects and chromsomal instability. The FA Group C complementation group gene encodes the protein FANCC, which is located in both cytoplasmic and nuclear compartments. FANCC is expressed in a cell cycle-dependent manner, with the lowest levels at the G1/S boundary and the highest levels in the M phase. The FANCC protein interacts with other FA complementation group proteins as well as non-FA proteins. A human α spectrin II acts as a scaffold to enhance interactions between FANCC and FANCA to form a nuclear complex. Another binding partner of FANCC is the BTB/POZ domain containing protein FAZF, which is a transcriptional repressor. In hematopoietic cells expressing mutant FANCC, PKR is constitutively phosphorylated and has increased binding affinity for double-stranded RNA, which suggests that FANCC indirectly suppresses the activity of PKR. These cells are apoptotic and are hypersensitive to IFN- γ and TNF α . In addition, FANCC protein is involved in the activation of Stat1 through receptors for at least three hematopoietic growth and survival factors.

CHROMOSOMAL LOCATION

Genetic locus: FANCC (human) mapping to 9q22.32; Fancc (mouse) mapping to 13 B3.

SOURCE

FANCC (C-14) is available as either goat (sc-18110) or rabbit (sc18110-R) polyclonal affinity purified antibody raised against a peptide mapping near the C-terminus of FANCC of human origin.

PRODUCT

Each vial contains 200 μg IgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-18110 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

APPLICATIONS

FANCC (C-14) is recommended for detection of FANCC of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 µg per 100-500 µg of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500), immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

FANCC (C-14) is also recommended for detection of FANCC in additional species, including equine, canine and porcine.

Suitable for use as control antibody for FANCC siRNA (h): sc-35354, FANCC siRNA (m): sc-35355, FANCC shRNA Plasmid (h): sc-35354-SH, FANCC shRNA Plasmid (m): sc-35355-SH, FANCC shRNA (h) Lentiviral Particles: sc-35354-V and FANCC shRNA (m) Lentiviral Particles: sc-35355-V.

Molecular Weight of FANCC: 60 kDa.

Positive Controls: K-562 nuclear extract: sc-2130, MOLT-4 nuclear extract: sc-2151 or Jurkat nuclear extract: sc-2132.

STORAGE

Store at 4° C, **DO NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

DATA





FANCC (C-14): sc-18110. Western blot analysis of FANCC expression in K-562 $({\bm A}),$ MOLT-4 $({\bm B})$ and Jurkat $({\bm C})$ nuclear extracts.

FANCC (C-14): sc-18110. Immunoperoxidase staining of formalin fixed, paraffin-embedded human tonsil tissue showing nuclear staining of cells in germinal centers and cells in non-germinal centers.

SELECT PRODUCT CITATIONS

- 1. Gordon, S., et al. 2005. FANCC, FANCE, and FANCD2 form a ternary complex essential to the integrity of the Fanconi anemia DNA damage response pathway. J. Biol. Chem. 280: 36118-36125.
- Sinha, S., et al. 2008. Alterations in candidate genes PHF2, FANCC, PTCH1 and XPA at chromosomal 9q22.3 region: pathological significance in earlyand late-onset breast carcinoma. Mol. Cancer 7: 84.

RESEARCH USE

For research use only, not for use in diagnostic procedures.

PROTOCOLS

See our web site at www.scbt.com or our catalog for detailed protocols and support products.

and suppo

MONOS Satisfation Guaranteed

Try **FANCC (6E7): sc-293308**, our highly recommended monoclonal alternative to FANCC (C-14).