# γ-GCSc (L-20): sc-15087



The Power to Question

## **BACKGROUND**

The GCLC gene consists of 16 exons and encodes the 636 amino acid protein  $\gamma\text{-GCSc}$  ( $\gamma\text{-glutamylcysteine}$  synthetase heavy subunit), also designated  $\gamma\text{-L-glutamate-L-cysteine}$  ligase catalytic subunit (GLCLC).  $\gamma\text{-GCSc}$  is expressed in hemocytes, brain, liver and kidney.  $\gamma\text{-GCSc}$  associates with a regulatory or modifier subunit,  $\gamma\text{-GCSm}$  ( $\gamma\text{-glutamylcysteine}$  synthetase light subunit), to form a heterodimer,  $\gamma\text{-GCS}$ .  $\gamma\text{-GCS}$  is the first enzyme involved and the rate determining step in glutathione biosynthesis. Oxidants, cadium and methyl mercury upregulate the transcription of  $\gamma\text{-GCS}$ .  $H_2O_2$  regulation depends on the Yap1 protein and the presence of glutamate, glutamine and lysine. Cadium regulates transcription through proteins Met-4, Met-31 and Met-32. Cbf1, a DNA binding protein, inhibits transcription of  $\gamma\text{-GCS}$ . Chemopreventive compounds cause increased levels of  $\gamma\text{-GCSc}$  in kidney tissues, which may protect against chemically induced carcinogenesis. A His 370 Leu amino acid change in  $\gamma\text{-GCSc}$  causes deficiencies in activity which are responsible for hemolytic anemia and low red blood cell glutathione levels.

## **REFERENCES**

- 1. Lunn, G., et al. 1979. Transport accounts for glutathione turnover in human erythrocytes. Blood 54: 238.
- Sierra-Rivera, E., et al. 1995. Assignment of the gene (GLCLC) that encodes the heavy subunit of γ-glutamylcysteine synthetase to human chromosome 6. Cytogenet. Cell Genet. 70: 278-279.
- Walsh, A.C., et al. 1996. Genetic mapping of GLCLC, the human gene encoding the catalytic subunit of γ-glutamyl-cysteine synthetase, to chromosome band 6p12 and characterization of a polymorphic trinucleotide repeat within its 5-prime untranslated region. Cytogenet. Cell Genet. 75: 14-16.
- Stephen, D.W., et al. 1997. Amino acid-dependent regulation of the Saccharomyces cerevisiae GSH1 gene by hydrogen peroxide. Mol. Microbiol. 23: 203-210.
- Thompson, S.A., et al. 1999. Induction of glutamate-cysteine ligase (gamma-glutamylcysteine synthetase) in the brains of adult female mice subchronically exposed to methylmercury. Toxicol. Lett. 110: 1-9.
- Beutler, E., et al. 1999. The molecular basis of a case of γ-glutamylcysteine synthetase deficiency. Blood 94:2890-2894.
- 6. Gipp, J.J., et al. 2000. Structure of the human glutamate-L-cysteine ligase catalytic (GLCLC) subunit gene. Cytogenet. Cell Genet. 88: 130-132.
- 7. Dormer, U.H., et al. 2000. Cadmium-inducible expression of the yeast GSH1 gene requires a functional sulfur-amino acid regulatory network. J. Biol. Chem. 275: 32611-32616.

## **CHROMOSOMAL LOCATION**

Genetic locus: GCLC (human) mapping to 6p12.1; Gclc (mouse) mapping to 9 E1.

# SOURCE

 $\gamma$ -GCSc (L-20) is an affinity purified goat polyclonal antibody raised against a peptide mapping within an internal region of  $\gamma$ -GCSc of human origin.

#### **PRODUCT**

Each vial contains 200  $\mu g$  lgG in 1.0 ml of PBS with <0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-15087 P, (100  $\mu$ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

## **APPLICATIONS**

 $\gamma$ -GCSc (L-20) is recommended for detection of  $\gamma$ -GCSc of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

 $\gamma$ -GCSc (L-20) is also recommended for detection of  $\gamma$ -GCSc in additional species, including equine, canine, bovine, porcine and avian.

Suitable for use as control antibody for  $\gamma$ -GCSc siRNA (h): sc-41978,  $\gamma$ -GCSc siRNA (m): sc-41979,  $\gamma$ -GCSc shRNA Plasmid (h): sc-41978-SH,  $\gamma$ -GCSc shRNA Plasmid (m): sc-41979-SH,  $\gamma$ -GCSc shRNA (h) Lentiviral Particles: sc-41978-V and  $\gamma$ -GCSc shRNA (m) Lentiviral Particles: sc-41979-V.

Molecular Weight of γ-GCSc: 73 kDa.

Positive Controls: Hep G2 cell lysate: sc-2227, A549 cell lysate: sc-2413 or A-431 whole cell lysate: sc-2201.

# **RECOMMENDED SECONDARY REAGENTS**

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use donkey anti-goat IgG-HRP: sc-2020 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible donkey anti-goat IgG-HRP: sc-2033 (dilution range: 1:2000-1:5000), Cruz Marker™ Molecular Weight Standards: sc-2035, TBS Blotto A Blocking Reagent: sc-2333 and Western Blotting Luminol Reagent: sc-2048. 2) Immunofluorescence: use donkey anti-goat IgG-FITC: sc-2024 (dilution range: 1:100-1:400) or donkey anti-goat IgG-TR: sc-2783 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

#### **STORAGE**

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

## **RESEARCH USE**

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



Try  $\gamma$ -GCSc (H-5): sc-390811 or  $\gamma$ -GCSc (F-9): sc-166356, our highly recommended monoclonal aternatives to  $\gamma$ -GCSc (L-20).

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