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## Produktinformation



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Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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### Zuschläge

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- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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# GALE (h4): 293T Lysate: sc-170673

## BACKGROUND

GALE, also known as galactowaldenase, UDP-galactose-4-epimerase or SDR1E1, is a 348 amino acid protein that functions as the third enzyme in the Leloir pathway of galactose metabolism. A member of the sugar epimerase family, GALE exists as a homodimer, binds FAD as a cofactor and catalyzes the epimerization of UDP-N-acetylglucosamine to UDP-N-acetylgalactosamine and UDP-glucose to UDP-galactose. The gene encoding GALE maps to human chromosome 1p36.11 and mutations in this gene lead to the development of complex disorder known as epimerase-deficiency galactosemia (EDG) or galactosemia type 3, which is characterized by mental retardation, liver damage, cataracts and deafness.

## REFERENCES

1. Reuser, A.J., Koster, J.F., Hoogeveen, A. and Galjaard, H. 1978. Biochemical, immunological, and cell genetic studies in glycogenosis type II. *Am. J. Hum. Genet.* 30: 132-143.
2. Holton, J.B., Gillett, M.G., MacFaul, R. and Young, R. 1981. Galactosaemia: a new severe variant due to uridine diphosphate galactose-4-epimerase deficiency. *Arch. Dis. Child.* 56: 885-887.
3. Henderson, M.J., Holton, J.B. and MacFaul, R. 1983. Further observations in a case of uridine diphosphate galactose-4-epimerase deficiency with a severe clinical presentation. *J. Inher. Metab. Dis.* 6: 17-20.
4. Kingsley, D.M., Kozarsky, K.F., Hobbie, L. and Krieger, M. 1986. Reversible defects in O-linked glycosylation and LDL receptor expression in a UDP-Gal/UDP-GalNAc 4-epimerase deficient mutant. *Cell* 44: 749-759.
5. Alano, A., Almashanu, S., Chinsky, J.M., Costeas, P., Blitzer, M.G., Wulfsberg, E.A. and Cowan, T.M. 1998. Molecular characterization of a unique patient with epimerase-deficiency galactosaemia. *J. Inher. Metab. Dis.* 21: 341-350.
6. Maceratesi, P., Daude, N., Dallapiccola, B., Novelli, G., Allen, R., Okano, Y. and Reichardt, J. 1998. Human UDP-galactose 4' epimerase (GALE) gene and identification of five missense mutations in patients with epimerase-deficiency galactosemia. *Mol. Genet. Metab.* 63: 26-30.
7. Wohlers, T.M., Christacos, N.C., Harreman, M.T. and Fridovich-Keil, J.L. 1999. Identification and characterization of a mutation, in the human UDP-galactose-4-epimerase gene, associated with generalized epimerase-deficiency galactosemia. *Am. J. Hum. Genet.* 64: 462-470.
8. Thoden, J.B., Wohlers, T.M., Fridovich-Keil, J.L. and Holden, H.M. 2001. Human UDP-galactose 4-epimerase. Accommodation of UDP-N-acetylglucosamine within the active site. *J. Biol. Chem.* 276: 15131-15136.
9. Schulz, J.M., Watson, A.L., Sanders, R., Ross, K.L., Thoden, J.B., Holden, H.M. and Fridovich-Keil, J.L. 2004. Determinants of function and substrate specificity in human UDP-galactose 4'-epimerase. *J. Biol. Chem.* 279: 32796-32803.

## STORAGE

Store at -20° C. Repeated freezing and thawing should be minimized. Sample vial should be boiled once prior to use. Non-hazardous. No MSDS required.

## CHROMOSOMAL LOCATION

Genetic locus: GALE (human) mapping to 1p36.11.

## PRODUCT

GALE (h4): 293T Lysate represents a lysate of human GALE transfected 293T cells and is provided as 100 µg protein in 200 µl SDS-PAGE buffer.

## APPLICATIONS

GALE (h4): 293T Lysate is suitable as a Western Blotting positive control for human reactive GALE antibodies. Recommended use: 10-20 µl per lane.

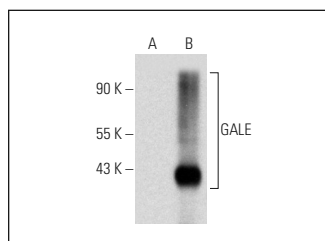
Control 293T Lysate: sc-117752 is available as a Western Blotting negative control lysate derived from non-transfected 293T cells.

GALE (C-4): sc-390407 is recommended as a positive control antibody for Western Blot analysis of enhanced human GALE expression in GALE transfected 293T cells (starting dilution 1:100, dilution range 1:100-1:1,000).

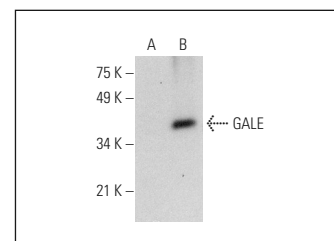
## RECOMMENDED SUPPORT REAGENTS

To ensure optimal results, the following support reagents are recommended:  
1) Western Blotting: use m-IgGκ BP-HRP: sc-516102 or m-IgGκ BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz Marker™ Molecular Weight Standards: sc-2035, UltraCruz® Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048.

## DATA



GALE (C-4): sc-390407. Western blot analysis of GALE expression in non-transfected: sc-117752 (A) and human GALE transfected: sc-170673 (B) 293T whole cell lysates.



GALE (D-11): sc-390460. Western blot analysis of expression in non-transfected: sc-117752 (A) and human GALE transfected: sc-170673 (B) 293T whole cell lysates.

## RESEARCH USE

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

## PROTOCOLS

See our web site at [www.scbt.com](http://www.scbt.com) for detailed protocols and support products.