

DESCRIPTION

Source Mouse myeloma cell line, NS0-derived
Met129-Tyr950 & Ser79-Tyr950, both with C-terminal 6-His tag
Accession # Q76M96

N-terminal Sequence Analysis Met129 & Ser79

Predicted Molecular Mass 100.1 kDa

SPECIFICATIONS

SDS-PAGE 125-150 kDa, reducing conditions

Activity Measured by its ability to inhibit proliferation of HT-29 human colon adenocarcinoma cells.
The ED₅₀ for this effect is 1-4 µg/mL.

Endotoxin Level <0.10 EU per 1 µg of the protein by the LAL method.

Purity >95%, by SDS-PAGE under reducing conditions and visualized by silver stain.

Formulation Lyophilized from a 0.2 µm filtered solution in PBS. See Certificate of Analysis for details.

PREPARATION AND STORAGE

Reconstitution Reconstitute at 250 µg/mL in sterile PBS.

Shipping The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below.

Stability & Storage Use a manual defrost freezer and avoid repeated freeze-thaw cycles.

- 12 months from date of receipt, -20 to -70 °C as supplied.
- 1 month, 2 to 8 °C under sterile conditions after reconstitution.
- 3 months, -20 to -70 °C under sterile conditions after reconstitution.

BACKGROUND

URB (upregulated in BRS-3 deficient mice) is a 150 kDa, secreted glycoprotein that belongs to the sushi-repeat-containing protein superfamily (1, 2). Sushi repeats, otherwise known as short consensus repeats (SCRs), are 60 amino acid (aa) sequences usually involved in protein-protein interaction. They are characterized by the presence of four Cys, two Pro, one Gly and one Trp (3). Human URB is synthesized as a 950 aa precursor that contains a 21 aa signal sequence and a 929 aa mature region. The mature molecule contains three extended sushi/SCR domains of approximately 140 aa each. They bear resemblance to the fifth sushi-repeat in human SPRX (4). The three lie between aa 141 - 281, 615 - 760, and 771 - 913, respectively. Between the first and second SCR lie two amino acid-rich regions, a Thr-rich domain (aa 347 - 404), and a Lys-rich domain (aa 487 - 588). Three potential N-linked glycosylation sites exist in the last two SCR's, while six potential bipartite nuclear localization signals (NLS) occur between aa 420 - 780. There are two potential alternate splice forms for human URB. One is 594 aa in length, and shows a simple truncation at Ser594. This effectively removes the second and third SCRs and two bipartite NLS (5). The second is 553 aa in length and shows a simple truncation after Lys553. This eliminates four bipartite NLSs, the second and third SCRs, and part of the Lys-rich domain (6). Full-length human URB is 83%, 84% and 87% aa identical to rat, mouse and bovine URB, respectively. URB is found in chondrocytes and appears to be downregulated upon CFU-Fibroblast differentiation (1). Thus, it may play a role in skeletogenesis.

References:

1. Liu, Y. *et al.* (2004) *Biochem. Biophys. Res. Commun.* **322**:497.
2. Aoki, K. *et al.* (2002) *Biochem. Biophys. Res. Commun.* **290**:1282.
3. Anatova, J. *et al.* (1989) *Biochemistry* **28**:4754.
4. Meindl, A. *et al.* (1995) *Hum. Mol. Genet.* **4**:2339.
5. Isogai, T. *et al.* (2002) GenBank Accession # BAC11475.
6. Strausberg, R.L. *et al.* (2002) GenBank Accession # AAH86876.