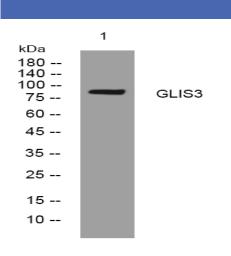


GLIS3 rabbit pAb

| Catalog No : | YT7769 |
|--------------------------|---|
| Reactivity : | Human;Mouse |
| Applications : | WB |
| Target : | GLIS3 |
| Gene Name : | GLIS3 ZNF515 |
| Protein Name : | GLIS3 |
| Human Gene Id : | 169792 |
| Human Swiss Prot | Q8NEA6 |
| No: | |
| Mouse Swiss Prot No : | Q6XP49 |
| Immunogen : | Synthesized peptide derived from human GLIS3 AA range: 381-431 |
| Specificity : | This antibody detects endogenous levels of GLIS3 at Human/Mouse |
| Formulation : | Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide. |
| Source : | Polyclonal, Rabbit,IgG |
| Dilution : | WB 1?500-2000 |
| Purification : | The antibody was affinity-purified from rabbit antiserum by affinity- chromatography using epitope-specific immunogen. |
| Concentration : | 1 mg/ml |
| Storage Stability : | -15°C to -25°C/1 year(Do not lower than -25°C) |
| Molecularweight : | 85kD |



| Background : | This gene is a member of the GLI-similar zinc finger protein family and encodes a nuclear protein with five C2H2-type zinc finger domains. This protein functions as both a repressor and activator of transcription and is specifically involved in the development of pancreatic beta cells, the thyroid, eye, liver and kidney. Mutations in this gene have been associated with neonatal diabetes and congenital hypothyroidism (NDH). Alternatively spliced variants that encode different protein isoforms have been described but the full-length nature of only two have been determined. [provided by RefSeq, Jul 2008], |
|---------------------------|--|
| Function : | disease:Defects in GLIS3 are a cause of NDH syndrome [MIM:610199]; also called neonatal diabetes mellitus with congenital hypothyroidism. NDH syndrome is a new neonatal diabetes syndrome associated with congenital hypothyroidism, congenital glaucoma, hepatic fibrosis and polycystic kidneys.,function:Acts as both a repressor and activator of transcription. Binds to the consensus sequence 5'-GACCACCCAC-3'.,similarity:Belongs to the GLI C2H2-type zinc-finger protein family.,similarity:Contains 5 C2H2-type zinc fingers.,tissue specificity:In the adult, expressed at high levels in the kidney and at lower levels in the brain, skeletal muscle, pancreas, liver, lung, thymus and ovary., |
| Subcellular Location : | Nucleus . |
| Expression : | In the adult, expressed at high levels in the kidney and at lower levels in the brain, skeletal muscle, pancreas, liver, lung, thymus and ovary. |
| Sort : | 6606 |
| No4 : | 1 |



Products Images

Western blot analysis of lysates from MDA-MB cells, primary antibody was diluted at 1:1000, $4\,^{\circ}{\rm over}$ night