For Research Use Only

Piezo2 Polyclonal antibody

Catalog Number:26205-1-AP 5 Publications

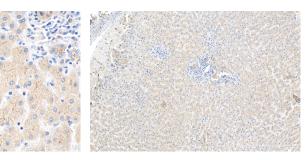


| Basic Information | Catalog Number: 26205-1-AP | GenBank Accession Number: AB527139 | Purification Method: Antigen affinity purification | |
|------------------------|---|---|--|--|
| | Size: 400 µg/ml | Genel D (NCBI): 63895 | Recommended Dilutions: WB 1:500-1:2000 | |
| | Source: Rabbit | UNIPROT ID: Q9H5I5 | IHC 1:200-1:800 | |
| | Isotype: IgG Immunogen Catalog Number: AG24528 | Full Name: family with sequence similarity 38 member B | | |
| | | Calculated MW: 318 kDa | | |
| | | Observed MW: 80 kDa | | |
| Applications | Tested Applications: | Positive Controls: | | |
| | IHC, WB, ELISA | WB: HUVEC cells, Neuro-2a cells, SK-N-SH cells | | |
| | Cited Applications: IHC, WB | IHC : human | n liver tissue, mouse eye tissue | |
| | Species Specificity: human, mouse | | | |
| | Cited Species: human, mouse | | | |
| | Note-IHC: suggested antigen retrieval with TE buffer pH 9.0; (*) Alternatively, antigen retrieval may be performed with citrate buffer pH 6.0 | | | |
| | FAM38B, also named as PIEZO2, is a mechanosensitive, rapidly inactivating (RI) ion channel which is open and converts the mechanical stimulus signals into bioelectrical signals after stimulated by mechanical signals. FAM3: has been recently identified in dorsal root ganglion (DRG) neurons to mediate tactile transduction. It plays an important role in the biological process, maintaining cell metabolism and cell migration. Loss-of-function mutations in the human FAM38B gene cause an autosomal recessive syndrome of muscular atrophy with perinata respiratory distress, arthrogryposis, and scoliosis.The 80 kDa band detected by SDS-PAGE can be caused by alternative splicing (PMID: 34335288, 37227654). | | | |
| Background Information | important role in the biological pr mutations in the human FAM38B g respiratory distress, arthrogryposi | rocess, maintaining cell metabolism and gene cause an autosomal recessive sync s, and scoliosis.The 80 kDa band detecte | l cell migration. Loss-of-function rome of muscular atrophy with perinat | |
| | important role in the biological pr mutations in the human FAM388 g respiratory distress, arthrogryposi alternative splicing (PMID: 34335) | rocess, maintaining cell metabolism and gene cause an autosomal recessive sync s, and scoliosis.The 80 kDa band detecte | I cell migration. Loss-of-function rome of muscular atrophy with perinat d by SDS-PAGE can be caused by | |
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For technical support and original validation data for this product please contact:T: 4006900926E: Proteintech-CN@ptglab.comW: ptgcn.com

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Selected Validation Data





Immunohistochemical analysis of paraffinembedded human liver tissue slide using 26205-1-AP (Piezo2 antibody) at dilution of 1:400 (under 40x lens). Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0). Immunohistochemical analysis of paraffinembedded human liver tissue slide using 26205-1-AP (Piezo2 antibody) at dilution of 1:400 (under 10x lens). Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0). Various lysates were subjected to SDS PAGE followed by western blot with 26205-1-AP (Piezo2 antibody) at dilution of 1:1000 incubated at room temperature for 1.5 hours.