

Product datasheet for TA372403S

WB

Q9H161

Primary Antibodies

ALX4 Rabbit Polyclonal Antibody

Product data:

Product Type:

Applications:

OriGene Technologies, Inc.

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| Recommended Dilution: | WB: 200-1000 WB positive control: Mouse brain tissue and HEPG2 cell lysates |
|-------------------------|--|
| Reactivity: | Human, Mouse |
| Host: | Rabbit |
| lsotype: | lgG |
| Clonality: | Polyclonal |
| Immunogen: | Synthetic peptide of human ALX4 |
| Formulation: | pH7.4 PBS, 0.05% NaN3, 40% Glycerol |
| Concentration: | lot specific |
| Purification: | Antigen affinity purification |
| Conjugation: | Unconjugated |
| Storage: | Store at -20°C. |
| Stability: | 1 year |
| Predicted Protein Size: | 44 kDa |
| Gene Name: | ALX homeobox 4 |
| Database Link: | Entrez Gene 60529 Human |



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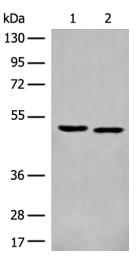
Serigene ALX4 Rabbit Polyclonal Antibody – TA372403S

Background: This gene encodes a paired-like homeodomain transcription factor expressed in the mesenchyme of developing bones, limbs, hair, teeth, and mammary tissue. Mutations in this gene cause parietal foramina 2 (PFM2); an autosomal dominant disease characterized by deficient ossification of the parietal bones. Mutations in this gene also cause a form of frontonasal dysplasia with alopecia and hypogonadism; suggesting a role for this gene in craniofacial development, mesenchymal-epithelial communication, and hair follicle development. Deletion of a segment of chromosome 11 containing this gene, del(11)(p11p12), causes Potocki-Shaffer syndrome (PSS); a syndrome characterized by craniofacial anomalies, mental retardation, multiple exostoses, and genital abnormalities in males. In mouse, this gene has been shown to use dual translation initiation sites located 16 codons apart.

Synonyms:

FPP; KIAA1788; PFM; PFM2

Product images:



Gel: 8%SDS-PAGE Lysate: 40 µg Lane 1-2: Mouse brain tissue and HEPG2 cell lysates Primary antibody: [TA372403] (ALX4 Antibody) at dilution 1/250 Secondary antibody: Goat anti rabbit IgG at 1/8000 dilution Exposure time: 2 minutes

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