

TMEM43 Polyclonal Antibody

Catalog No: #31580



Package Size: #31580-1 50ul #31580-2 100ul

Orders: order@signalwayantibody.com
Support: tech@signalwayantibody.com

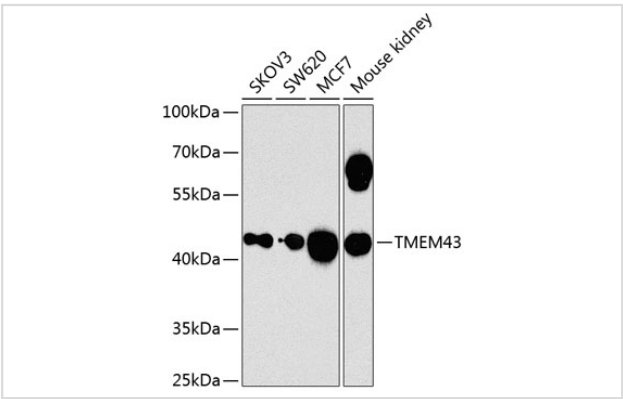
Description

| | |
|-----------------------|---|
| Product Name | TMEM43 Polyclonal Antibody |
| Host Species | Rabbit |
| Clonality | Polyclonal |
| Isotype | IgG |
| Purification | Affinity purification |
| Applications | WB |
| Species Reactivity | Human,Mouse |
| Immunogen Description | Recombinant fusion protein of human TMEM43 (NP_077310.1). |
| Other Names | TMEM43;ARVC5;ARVD5;EDMD7;LUMA |
| Accession No. | Uniprot:Q9BTV4GeneID:79188 |
| Uniprot | Q9BTV4 |
| GeneID | 79188 |
| Calculated MW | 45kDa |
| SDS-PAGE MW | 45kDa |
| Formulation | PBS with 0.02% sodium azide,50% glycerol,pH7.3. |
| Storage | Store at -20°C. Avoid freeze / thaw cycles. |

Application Details

WB 1:500 - 1:2000

Images



Western blot analysis of extracts of various cell lines, using TMEM43 antibody.

Background

This gene belongs to the TMEM43 family. Defects in this gene are the cause of familial arrhythmogenic right ventricular dysplasia type 5 (ARVD5), also known as arrhythmogenic right ventricular cardiomyopathy type 5 (ARVC5). Arrhythmogenic right ventricular dysplasia is an inherited disorder, often involving both ventricles, and is characterized by ventricular tachycardia, heart failure, sudden cardiac death, and fibrofatty replacement of

cardiomyocytes. This gene contains a response element for PPAR gamma (an adipogenic transcription factor), which may explain the fibrofatty replacement of the myocardium, a characteristic pathological finding in ARVC.

Note: This product is for in vitro research use only