

## Datasheet

### MYH9 monoclonal antibody (M03), clone 2B3

**Catalog Number:** H00004627-M03

**Regulation Status:** For research use only (RUO)

**Product Description:** Mouse monoclonal antibody raised against a partial recombinant MYH9.

**Clone Name:** 2B3

**Immunogen:** MYH9 (AAH11915, 131 a.a. ~ 220 a.a) partial recombinant protein with GST tag. MW of the GST tag alone is 26 KDa.

**Sequence:**

RLKQLKRQLEEAEEEEAQRANASRRKLQRELEDATETA  
DAMNREVSSLKNLRRGDLFPVPRRMARKGAGDGS  
DEEVDGKADGAEAKPAE

**Host:** Mouse

**Reactivity:** Human

**Applications:** ELISA, IF, IHC-P, S-ELISA, WB-Ce, WB-Re, WB-Ti, WB-Tr  
(See our web site product page for detailed applications information)

**Protocols:** See our web site at <http://www.abnova.com/support/protocols.asp> or product page for detailed protocols

**Isotype:** IgG2b Kappa

**Storage Buffer:** In 1x PBS, pH 7.4

**Storage Instruction:** Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

**Entrez GeneID:** 4627

**Gene Symbol:** MYH9

**Gene Alias:** DFNA17, EPSTS, FTNS, MGC104539, MHA, NMHC-II-A, NMMHCA

**Gene Summary:** This gene encodes a myosin IIA heavy chain that contains an IQ domain and a myosin head-like

domain. The protein is involved in several important functions, including cytokinesis, cell motility and maintenance of cell shape. Defects in MYH9 are the cause of non-syndromic sensorineural deafness autosomal dominant type 17, Epstein syndrome, Alport syndrome with macrothrombocytopenia, Sebastian syndrome, Fechtner syndrome and macrothrombocytopenia with progressive sensorineural deafness. [provided by RefSeq]