# MYH9 Antibody

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| **Product Code** | CSB-PA015303GA01HU |
| **Storage** | Upon receipt, store at -20°C or -80°C. Avoid repeated freeze. |
| **Uniprot No.** | P35579 |
| **Immunogen** | Human MYH9 |
| **Raised In** | Rabbit |
| **Species Reactivity** | Human,Mouse,Rat |
| **Tested Applications** | ELISA,WB,IHC,IF |
| **Storage Buffer** | PBS with 0.1% Sodium Azide, 50% Glycerol, pH 7.3. -20°C, Avoid freeze / thaw cycles. |
| **Purification Method** | Antigen Affinity Purified |
| **Isotype** | IgG |
| **Alias** | myosin, heavy chain 9, non-muscle;MYH9;DFNA17;EPSTS;FTNS;MGC104539;MHA;NMHC-II-A;NMMHCA ; |
| **Product Type** | Purified Rabbit Anti human PolyClonal Antibody |
| **Immunogen Species** | Homo sapiens (Human) |
| **Target Names** | MYH9 |
| **Target Details** | 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. |