



## Dynactin 1 (N-term)

### Mouse monoclonal Antibody

#53110

**Catalog Number:** 53110

**Amount:** 100µg/100µl

**Swiss-Prot No. :**Q14203

**Gene name:**dctn1

**Gene id:**1639

**Clone Number:** 3D5-C6-D5

**Form of Antibody:**Purified mouse monoclonal in buffer containing 0.1M Tris-Glycine (pH 7.4, 150 mM NaCl) with 0.2% sodium azide, 50% glycerol

**Storage/Stability:** Store at -20°C/1 year

**Immunogen:** Purified recombinant human Dynactin 1 protein fragments expressed in E.coli.

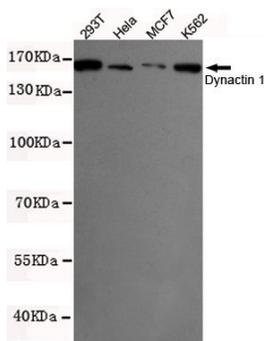
**Purification:** affinity-chromatography

**Specificity/Sensitivity:**This antibody detects endogenous levels of Dynactin 1 and does not cross-react with related proteins

**Reactivity:** Human

**Applications:**

Predicted MW: 150 kd WB: 1:500



Western blot detection of Dynactin 1 in K562, MCF7, 293T and HeLa cell lysates using Dynactin 1 mouse mAb (1:500 diluted). Predicted band size: 150 kDa. Observed band size: 150 kDa.

**Background :**

This gene encodes the largest subunit of dynactin, a macromolecular complex consisting of 10 subunits ranging in size from 22 to 150 kD. Dynactin binds to both microtubules and cytoplasmic dynein. Dynactin is involved in a diverse array of cellular functions, including ER-to-Golgi transport, the centripetal movement of lysosomes and endosomes, spindle formation, chromosome movement, nuclear positioning, and axonogenesis. This subunit interacts with dynein intermediate chain by its domains directly binding to dynein and binds to microtubules via a highly conserved glycine-rich cytoskeleton-associated protein (CAP-Gly) domain in its N-terminus. Alternative splicing of this gene results in multiple transcript variants encoding distinct isoforms. Mutations in this gene cause distal hereditary motor neuropathy type VII B (HMN7B) which is also known as distal spinal and bulbar muscular atrophy (dsBMA).