# **Technical Data Sheet**

# **Purified Mouse Anti-p53**

### **Product Information**

**Material Number:** 610183 Size: 50 μg 250 μg/ml Concentration: 80/p53 Clone:

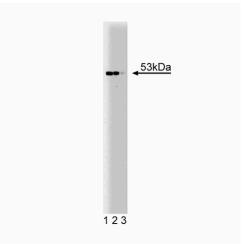
Monkey p53 aa. 195-393 Immunogen: Mouse IgG2b, κ Isotype: Reactivity: QC Testing: Human Tested in Development: Dog

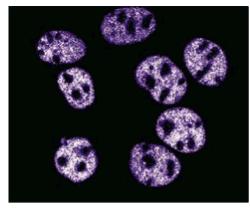
Target MW:

Storage Buffer: Aqueous buffered solution containing BSA, glycerol, and ≤0.09% sodium

#### Description

The p53 protein is critical to regulation of normal cell growth and is a suppressor of tumor cell proliferation. Inactivation of p53 by a number of mechanisms, such as missense mutations or interaction with oncogenic viral or cellular proteins, can result in tumor progression. Mutations and/or allelic loss of the p53 gene is associated with a wide variety of human tumors. Known to have a role in transcriptional regulation, p53 suppresses various promoters containing TATA elements in an apparently sequence-independent fashion. p53 also binds to DNA in a sequence-specific manner via recognition of a 20 bp consensus-binding site. This interaction stimulates the expression of genes downstream of the p53 binding site. A number of genes that contain p53-binding sites have been identified, including MDM2, GADD45, and muscle creatine kinase. It is thought that MDM2 is a feedback regulator of p53. In addition, a p53-inducible gene, Cip1, has been identified and shown to suppress tumor cell growth in culture.





Western blot analysis of p53 on A431 lysate. Lane 1: 1:500, Jane 2: 1:1000, Jane 3: 1:2000 dilution of p53.

A431

### **Preparation and Storage**

The monoclonal antibody was purified from tissue culture supernatant or ascites by affinity chromatography. Store undiluted at -20°C.

# **Application Notes**

### Application

Western blot	Routinely Tested
Immunoprecipitation	Tested During Development
Immunofluorescence	Tested During Development
Immunohistochemistry	Not Recommended

### **BD Biosciences**

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### **Product Notices**

- 1. Since applications vary, each investigator should titrate the reagent to obtain optimal results.
- 2. Please refer to www.bdbiosciences.com/pharmingen/protocols for technical protocols.
- 3. Caution: Sodium azide yields highly toxic hydrazoic acid under acidic conditions. Dilute azide compounds in running water before discarding to avoid accumulation of potentially explosive deposits in plumbing.
- 4. Source of all serum proteins is from USDA inspected abattoirs located in the United States.

#### References

Hurd C, Khattree N, Alban P. Hormonal regulation of the p53 tumor suppressor protein in T47D human breast carcinoma cell line. *J Biol Chem.* 1995; 270(48):28507-28510.(Clone-specific: Western blot)

Mercer WE. Cell cycle regulation and the p53 tumor suppressor protein. Crit Rev Eukaryot Gene Expr.. 1992; 2(3):251-263.(Biology)

Natsugoe S, Nakashima S, Matsumoto M. Expression of p21WAF1/Cip1 in the p53-dependent pathway is related to prognosis in patients with advanced esophageal carcinoma. Clin Cancer Res. 1999; 5(9):2445-2449.(Clone-specific: Immunohistochemistry)

Saitoh H, Pizzi MD, Wang J. Perturbation of SUMOlation enzyme Ubc9 by distinct domain within nucleoporin RanBP2/Nup358. *J Biol Chem.* 2002; 277(7):4755-4763.(Clone-specific: Immunofluorescence)

Stahnke K, Fulda S, Friesen C, Strauss G, Debatin KM. Activation of apoptosis pathways in peripheral blood lymphocytes by in vivo chemotherapy. *Blood.* 2001; 98(10):3066-3073.(Clone-specific: Flow cytometry, Western blot)

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