## **Certificate of Analysis**

## Actin, beta Human (non-muscle) Western Blot Positive **Control**

## **Purified**

Catalog No		CAS No	Molecular Formula	Molecular	Weight	Storage
MBS634566						-20°C
Lot No	<b>Control No</b>	<b>Revision No</b>	Revised By A	Approved By		
L20030554	C20030554					

Actin and myosin are the two major cytoskeleton proteins implicated in cellular movement, secretion, phagocytosis, and kinesis. Actin is one of the most conserved cellular protein. At least 6 actin isoforms have been identified by protein sequence analyses. identified by protein sequence analyses. Four actin isoforms represent the differentiation markers of muscle tissues.

**Purity:** 

There are three a-actins: a-skeletal, a-cardiac, and a-smooth muscle), one b-actin (b-non-muscle), and two g-actins b-actin (b-non-muscle), and two g-actins (g-smooth muscle and g-non-muscle). Actin isoform are >90% conserved, except in the N-terminal 18-aa (50-60% homology). Beta-actin protein and mRNA levels are often used as a reference for comparing changes in cellular protein/mRNA levels by Western or Northern blots. Northern blots.

Purified by affinity chromatography to >95% purity.

Supplied as a liquid in SDS-PAGE sample buffer (reduced). Form:

researcher.

Suitable for use in Western Blot. Other **Applications:** applications not tested.

Western Blot: load 10ul/lane. **Recommended Dilutions:** Note: SDS may crystallize in cold conditions. It should redissolve by

warming before taking it from the stock. It should be heated once prior to loading on gels. If the product has been stored for several weeks, then it may be preferable to add 5ul of fresh 2x sample buffer per 10ul of the A0760-47B solution prior to heating and loading on gels. This preparation is not biologically active. Optimal dilutions to be determined by the researcher.

Storage and Stability:
May be stored at 4°C for short-term only. Aliquot to avoid repeated freezing and thawing. Store at -20°C. Aliquots are stable for 6 months after receipt at -20°C. For maximum recovery of product, centrifuge the original vial after thawing and prior to removing the cap. Further dilutions can be made in assay buffer.

1. Ohmuri H (1995) Gene Accession #S38782; Vandekerchove, J et al (1978) Eur. J. Biochem. 90, **451**. 2. Lessard J et al (1988) Cell. Motil Cytoskel. 10, **349**. 3. North JA et al (1994) J. Cell Sci. **107**, 437.