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## **MOCOS (N) Antibody, Rabbit Polyclonal**

**Cat#: R1047-1**

**Quantity: 100 ul**

**Predicted M.W.: 98 kDa**

**Lot#: Refer to vial**

**Application: WB**

**Uniprot ID: Q96EN8**

### **Background:**

Molybdenum cofactor sulfurase (MOCOS) belongs to the class-V pyridoxal-phosphate-dependent aminotransferase family. It sulfurates the molybdenum cofactor. Sulfation of molybdenum is essential for xanthine dehydrogenase (XDH) and aldehyde oxidase (ADO) enzymes in which molybdenum cofactor is liganded by 1 oxygen and 1 sulfur atom in active form. In vitro, the C-terminal domain is able to reduce N-hydroxylated prodrugs, such as benzamidoxime. Defects in MOCOS are the cause of xanthinuria type 2 (XU2). Patients suffering XU2 cannot metabolize allopurinol into oxypurinol due to dual deficiency of xanthine dehydrogenase and aldehyde oxidase.

### **Other Names:**

Molybdenum cofactor sulfurase, MCS, MOS, MoCo sulfurase, hMCS

### **Source and Purity:**

Rabbit polyclonal antibodies were produced by immunizing animals with a GST-fusion protein containing the N-terminal region of human MOCOS. Antibodies were purified by affinity purification using immunogen.

### **Storage Buffer and Condition:**

Supplied in 1 x PBS (pH 7.4), 100 ug/ml BSA, 40% Glycerol, 0.01% NaN<sub>3</sub>. Store at -20 °C. Stable for 6 months from date of receipt.

### **Species Specificity:**

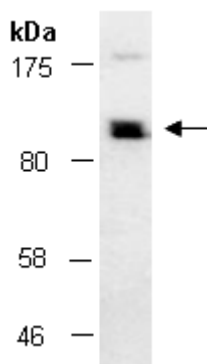
Human

### **Tested Applications:**

WB: 1:500-1:1,000 (detect endogenous protein\*)

\*: The apparent protein size on WB may be different from the calculated M.W. due to modifications.

**Product Data:**



**Fig 1.** Western blot of total cell extracts from human HepG2, using Ab (R1047-1) at RT for 2 h.