# abcam

### Product datasheet

# Recombinant Human GNMT protein ab79190

#### 画像数1

#### 製品の詳細

製品名 Recombinant Human GNMT protein

精製度 > 95 % SDS-PAGE.

ab79190 is purified using conventional chromatography techniques. Endotoxin Level: < 1.0 EU

per 1µg of protein (determined by LAL method)

**発現系** Escherichia coli

タンパク質長 Full length protein

Animal free No

**由来** Recombinant

生物種 Human

配列 MGSSHHHHHH SSGLVPRGSH MVDSVYRTRS

LGVAAEGLPD QYADGEAARV WQLYIGDTRS
RTAEYKAWLL GLLRQHGCQR VLDVACGTGV
DSIMLVEEGF SVTSVDASDK MLKYALKERW
NRRHEPAFDK WVIEEANWMT LDKDVPQSAE
GGFDAVICLG NSFAHLPDCK GDQSEHRLAL
KNIASMVRAG GLLVIDHRNY DHILSTGCAP
PGKNIYYKSD LTKDVTTSVL IVNNKAHMVT
LDYTVQVPGA GQDGSPGLSK FRLSYYPHCL
ASFTELLQAA FGGKCQHSVL GDFKPYKPGQ

TYIPCYFIHV LKRTD

### 特性

Our **Abpromise guarantee** covers the use of **ab79190** in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

アプリケーション SDS-PAGE

製品の状態 Liquid

前処理および保存

保存方法および安定性 Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.

pH: 8.00

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#### 関連情報

機能 Catalyzes the methylation of glycine by using S-adenosylmethionine (AdoMet) to form N-

methylglycine (sarcosine) with the concomitant production of S-adenosylhomocysteine (AdoHcy).

Possible crucial role in the regulation of tissue concentration of AdoMet and of metabolism of

methionine.

組織特異性 Abundant in liver.

関連疾患 Defects in GNMT are the cause of glycine N-methyltransferase deficiency (GNMT deficiency)

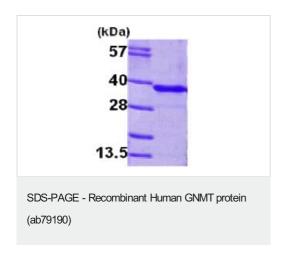
[MIM:606664]; also known as hypermethioninemia. The only clinical abnormalities in patients with

this deficiency are mild hepatomegaly and chronic elevation of serum transaminases.

配列類似性 Belongs to the glycine N-methyltransferase family.

細胞内局在 Cytoplasm.

#### 画像



15% SDS-PAGE showing ab79190 at approximately 35kDa (3µg).

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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