

# S- Adenosylmethionine Antibody (Clone # 118-6)

<b>ALTERNATE NAMES:</b>	S- Adenosylmethionine
<b>CATALOG #:</b>	6940-25
<b>AMOUNT:</b>	25 µl
<b>HOST/ISOTYPE:</b>	Mouse IgG2b
<b>IMMUNOGEN:</b>	S-Adenosylmethionine analog Aza-SAM conjugated to KLH
<b>INTERNAL ID:</b>	DM-09
<b>FORM:</b>	Liquid
<b>FORMULATION:</b>	10 mM PBS (pH 7.4), 0.02% Sodium azide, 50% Glycerol and 1% BSA
<b>PURIFICATION:</b>	>95% Purified from mouse ascites fluid by affinity chromatograph
<b>SPECIES REACTIVITY:</b>	All
<b>STORAGE CONDITIONS:</b>	Store at 4°C; for long storage, store at -20°C. Avoid multiple freeze-thaw cycles.

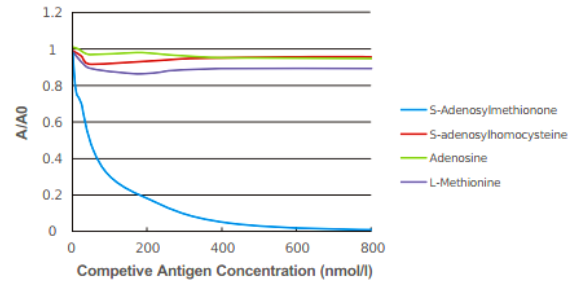
**DESCRIPTION:** S-Adenosylmethionine (SAM) is a naturally occurring compound that is found in almost every tissue and fluid in the body. It is a common co-substrate involved in methyl group transfers. It is made from adenosine triphosphate (ATP) and methionine by methionine adenosyl transferase. Transmethylation, transsulfuration, and aminopropylation are the metabolic pathways that use SAM. Although these anabolic reactions occur throughout the body, most SAM is produced and consumed in the liver. SAM plays a role in the immune system, maintains cell membranes, and helps produce and break down brain chemicals, such as serotonin, melatonin, and dopamine. It works with vitamin B12 and folate (vitamin B9). Being deficient in either vitamin B12 or folate may reduce levels of SAM in your body.

**APPLICATION:** cELISA: 1:10000, FCM: 1:200/400, IHC: 1:200/400.

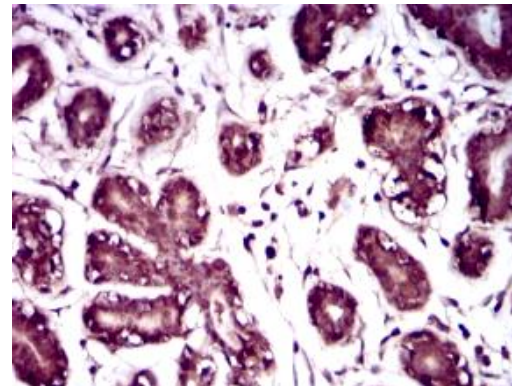
**SPECIFICITY:** Shows the following reactivities with related compounds: S-Adenosylmethionine: 100%, S-Adenosylhomocysteine: <1%, Adenosine: <1%, L-Methionine: <1%.

**Note:** This information is only intended as a guide. The optimal dilutions must be determined by the user.

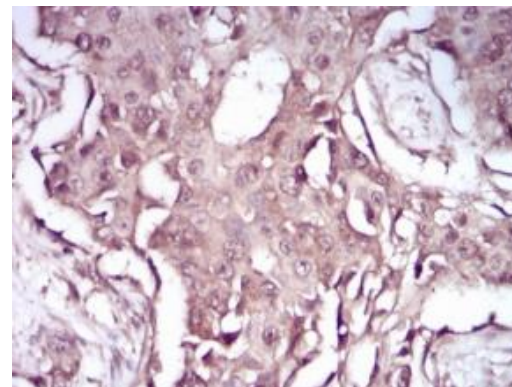
**FOR RESEARCH USE ONLY! Not to be used on humans.**



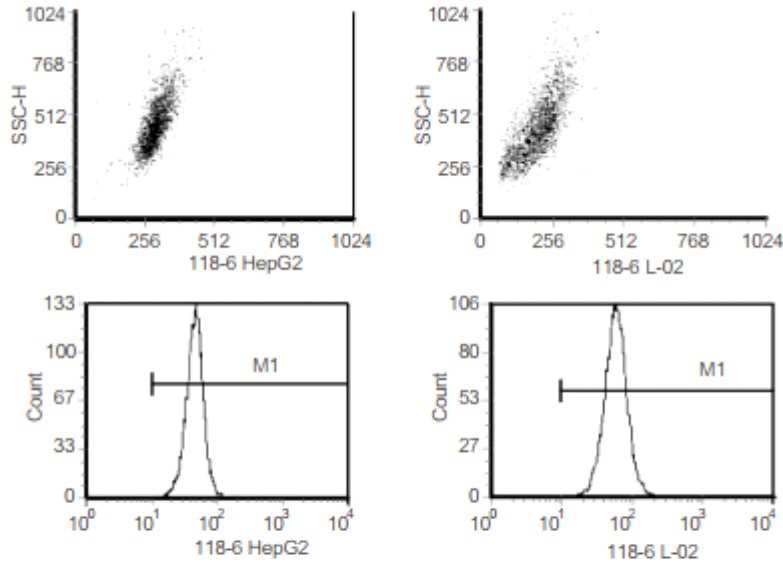
**Competitive ELISA:** 0.1 g/ml of SAM coating standard was coated into 96 wells. Serial dilution of SAM standard, S-Adenosylhomocysteine (SAH), Adenosine, L-Methionine and antibody were added. HRP conjugated Goat anti-Mouse IgG antibody was used to develop the color. The A is the OD450 value of the test well and the A0 is the OD450 of the well without competitive antigen.



**Immunohistochemistry** staining was performed using the antibody with benign breast tissue adjacent to carcinoma. Brown areas indicated strong positive staining in nuclear and cytoplasmic areas.



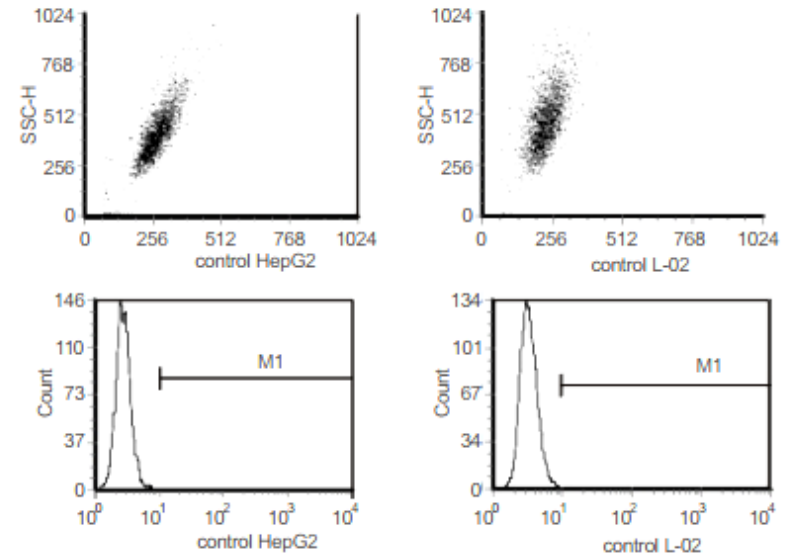
The same samples as in above figure from breast cancer area. Cytoplasmic and nuclear areas showed negative or much weak or background staining.



Marker	Low bound	High bound	# of Events	# of gated cells	Geometric Mean
ALL	1	10000	10000	100.0	45.34
M1	10	10000	9979	99.79	45.55

Marker	Low bound	High bound	# of Events	# of gated cells	Geometric Mean
ALL	1	10000	10000	100.0	58.22
M1	10	10000	9992	99.92	58.34

**FCM** results from normal liver cell line L02 and hepatocyte carcinoma cell line Hep G2 stained with anti-SAM monoclonal antibody from clone 118-6. Average fluorescence signal in Hep G2 cells was reduced compared to that in L02 cells, indicating SAM level is reduced during carcinogenesis.



Marker	Low bound	High bound	# of Events	# of gated cells	Geometric Mean
ALL	1	10000	10000	100.0	2.72
M1	10	10000	17	0.17	48.98

Marker	Low bound	High bound	# of Events	# of gated cells	Geometric Mean
ALL	1	10000	10000	100.0	3.33
M1	10	10000	44	0.44	17.12

**FCM analysis control.** Normal liver cells L02 and carcinoma cells Hep G2 were stained with the buffer without any antibody.

**RELATED PRODUCTS:**

- S- Adenosylmethionine Antibody (Clone # 84-3) (Cat # 6941-25)
- S- Adenosylmethionine Antibody (Clone # 118-18) (Cat # 6942-25)
- S- Adenosylmethionine Antibody (Clone # 84-19) (Cat # 6943-25)
- S- Adenosylmethionine Antibody (Cat # 6944-25)
- S- Adenosylhomocysteine Antibody (Clone # 301-10) (Cat # 6945-25)