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## Mouse Monoclonal antibody anti-human A2AR

#Cat: NB-19-0002    Size: 100 uL

### Description

Adenosine is an endogenous nucleoside released from cells by facilitated diffusion and is also produced extracellularly by degradation of ATP. Adenosine receptors are subdivided into four G-coupled protein receptors subtypes (A1, A2A, A2B and A3) depending on their pharmacological properties to exert numerous effects on various tissues including the central nervous system.

Most of the anti-inflammatory effects of adenosine have been assigned to the A2A receptor subtype (A2AR).

A2AR is expressed in many immune and inflammatory cells and is up-regulated by T- helper cell type 1 cytokines. Because changes of peripheral A2AR reflect changes that occur at the injured tissue. A2AR assay appears as a valuable marker for monitoring treatment in patients with inflammatory cells infiltrating the failing organ.

### Product Information

<b>Host:</b>	Mouse
<b>Applications:</b>	ELISA, WB
<b>Reactivity:</b>	Human
<b>Clonality:</b>	Monoclonal
<b>Clone ID:</b>	ADONIS
<b>Conjugation:</b>	Unconjugated
<b>Isotype:</b>	IgM
<b>Formulation:</b>	Antibody obtained by ammonium sulphate precipitation.
<b>Constituent:</b>	PBS pH 7.4
<b>Concentration:</b>	>1 mg/mL (exact concentration is lot-dependent)
<b>Storage Instruction:</b>	For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

### Target

<b>Protein Name:</b>	A2A receptor subtype (A2AR)
<b>Immunogen:</b>	C-Terminal part of second extracellular loop of the A2A receptor

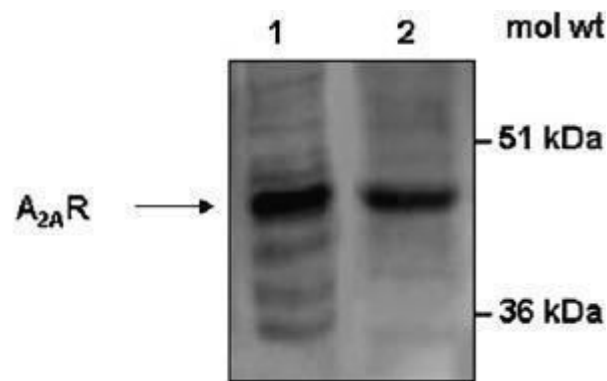
### Applications

This antibody is recommended for detection of A2A receptor subtype (A2AR) of human origin by ELISA, Western Blotting. For functional activity, a custom size azide-free is available.

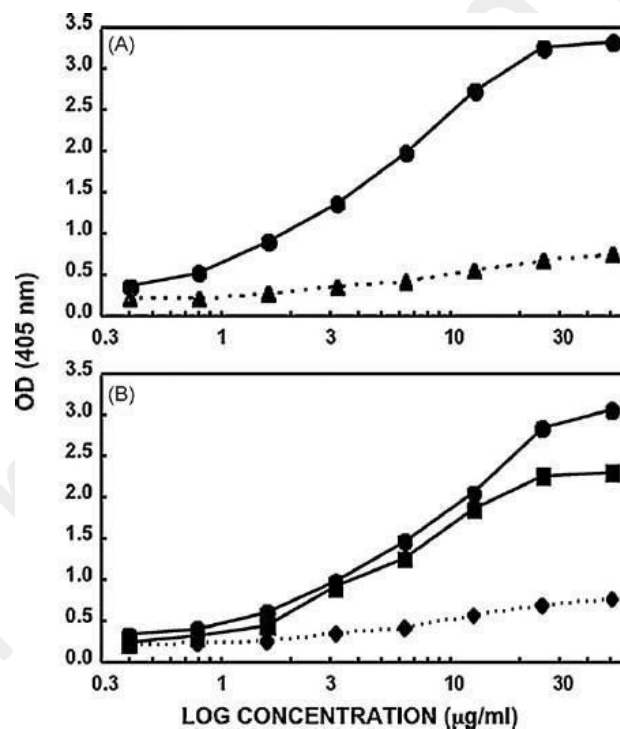
### Precaution

**For professional users.** Proper handling of this product as with any product derived from biological sources according to local and applicable regulations.

## Data



Western blots of the Adonis binding on reduced lysate of A<sub>2A</sub>-Chem-3 cells (lane 1) and human PBMC (lane 2).



Dose-response curves of Adonis binding to: (A) the immunogen peptide, uncoated wells served as blanks and

(B) glutaraldehyde fixed A<sub>2A</sub>-Chem-3 cells (circle) and normal human PBMC (square), glutaraldehyde-treated uncoated wells (lozenge) served as blanks. Results are given in optical density read at 405nm and are the mean values of duplicates.

## References

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For reference only