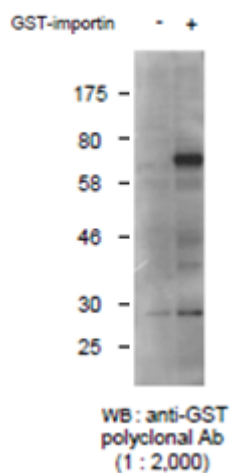


## Anti-GST antibody, rabbit serum

<b>Product code</b>	60-021
<b>Size</b>	100 µl
<b>Storage</b>	Store at 4°C for short term. For long term storage store at -20°C. Aliquot to avoid repeated freezing thawing.
<b>Concentration</b>	N/A
<b>Buffer</b>	0.05% sodium azide
<b>Purity</b>	Rabbit antiserum
<b>Immunogen</b>	Recombinant full-size GST (aa 1-212)
<b>Isotype</b>	Rabbit IgG
<b>Reactivity</b>	Specific to GST and GST-tagged proteins
<b>Special notes</b>	N/A
<b>Application</b>	<ol style="list-style-type: none"> <li>1. Western blotting (dilution: 1/2,000~1/10,000)</li> <li>2. Immunoprecipitation (assay dependent)</li> <li>3. ELISA</li> </ol> <p>Other applications have not been tested.</p>
<b>Background</b>	<p>Glutathione S transferase (GST) from <i>Schistosoma japonicum</i> is commonly used to create fusion proteins. GST-tag has the size of 220 amino acids (roughly 26kDa) and is fused to the N-terminus of a protein. GST fusion proteins can be produced in <i>Escherichia coli</i>, as recombinant proteins and are used to purify and detect proteins of interest. The GST part binds its substrate, glutathione. GST-fusions protein can be easily purified from cell extracts by affinity chromatography with glutathione resin.</p>
<b>Data Link</b>	NCBI Protein Data <a href="https://www.ncbi.nlm.nih.gov/protein/AAA57089">AAA57089</a>
Please note: All products are FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES. NOT FOR MILITARY USE.	

**Data Images:** 60-021 Anti-GST antibody, rabbit serum

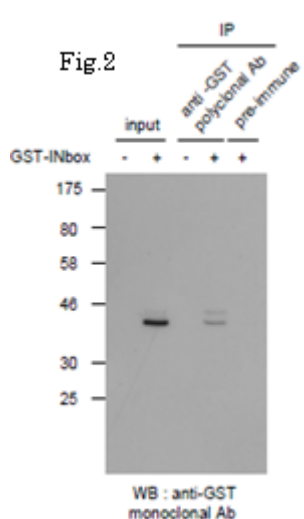
Fig.1



**Fig.1 Detection of GST-tagged protein with this antibody by Western blotting.**

-: Lysate of 293T cells transfected with an empty vector

+: Lysate of 293T cells transfected with the plasmid carrying the GST-tagged importin gene



**Fig.2 Immunoprecipitation of GST-tagged protein with this antibody followed by Western blotting.**

-: Lysate of 293T cells transfected with an empty vector

+: Lysate of 293T cells transfected with the plasmid carrying the GST-tagged INbox gene

### References:

1. Smith DB & Johnson KS (1988) "Single-step purification of polypeptides expressed in *Escherichia coli* as fusions of glutathione-S-transferase." *Gene* **67**:31-40 PMID: [3047011](#)
2. Kaelin WG Jr *et al* (1991) "Identification of cellular proteins that can interact specifically with the T/E1A-binding region of the retinoblastoma gene product." *Cell* **64**:521-532 PMID: [1825028](#)
3. *Molecular Cloning: A laboratory Manual* (eds. Sambrook, J., Russell, D.W. Cold Spring Harbor Laboratory Press, Cold Spring Harbor, New York, USA, 2001) pp.15.36-15.39, pp.18.48-18.59.