## Phospho-eNOS (Ser1177) Antibody

For Research Use Only. Not for Use in Diagnostic Procedures.

| Applications: | Reactivity: | Sensitivity: | MW (kDa): | Source: | UniProt ID: |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| WB | H B Pg | Endogenous | 140 | Rabbit | Entrez-Gene Id: |
|  |  |  |  |  |  |
|  |  |  |  |  |  |

## Product Usage <br> Information

Storage
Specificity / Sensitivity

Species predicted to react based on 100\% sequence homology:

Source / Purification

Background

| Application | Dilution |
| :--- | :--- |
| Western Blotting | 1:1000 |

Supplied in 10 mM sodium HEPES (pH 7.5), $150 \mathrm{mM} \mathrm{NaCl}, 100 \mu \mathrm{~g} / \mathrm{ml}$ BSA and $50 \%$ glycerol. Store at $20^{\circ} \mathrm{C}$. Do not aliquot the antibody.

Phospho-eNOS (Ser1177) Antibody detects endogenous levels of eNOS only when phosphorylated at Ser1177.

Mouse, Rat

Polyclonal antibodies are produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Ser1177 of human eNOS. Antibodies are purified by protein A and peptide affinity chromatography.

Endothelial nitric-oxide synthase (eNOS) is an important enzyme in the cardiovascular system. It catalyzes the production of nitric oxide (NO), a key regulator of blood pressure, vascular remodeling, and angiogenesis $(1,2)$. The activity of eNOS is regulated by phosphorylation at multiple sites. The two most thoroughly studied sites are the activation site Ser1177 and the inhibitory site Thr495 (3). Several protein kinases including Akt/PKB, PKA, and AMPK activate eNOS by phosphorylating Ser1177 in response to various stimuli $(4,5)$. In contrast, bradykinin and $\mathrm{H}_{2} \mathrm{O}_{2}$ activate eNOS activity by promoting both Ser1177 phosphorylation and Thr495 dephosphorylation (6,7).

Background References

1. Fulton, D. et al. (2001) J Pharmacol Exp Ther 299, 818-24.
2. Shaul, P.W. (2002) Annu Rev Physiol 64, 749-74.
3. Chen, Z.P. et al. (1999) FEBS Lett 443, 285-9.
4. Dimmeler, S. et al. (1999) Nature 399, 601-5.
5. Fulton, D. et al. (1999) Nature 399, 597-601.
6. Harris, M.B. et al. (2001) J Biol Chem 276, 16587-91.
7. Thomas, S.R. et al. (2002) J Biol Chem 277, 6017-24.

## Species Reactivity

Western Blot Buffer

## Applications Key

Cross-Reactivity Key

## Trademarks and Patents

Species reactivity is determined by testing in at least one approved application (e.g., western blot).

IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5\% w/v BSA, 1X TBS, $0.1 \%$ Tween $® 20$ at $4^{\circ} \mathrm{C}$ with gentle shaking, overnight.

WB: Western Blotting
H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse GP: Guinea Pig Rab: rabbit All: all species expected

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