



# Anti Equine Endoglin (CD105)

## Background

Endoglin (CD105) is mainly expressed on endothelial cells of newly formed vessels and increased under hypoxic environment. Thus, endoglin could be used as a powerful marker of neovascularization in solid malignancies including mammary carcinoma as well as regenerating stage of injury or inflammation.

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Product type	Primary Antibody
Immunogen	Equine Endoglin(a.a.1 - 112)
Raised in	Mouse
Myeloma	P3U1
Clone number	2A8B3
Isotype	IgG2a, $\kappa$
Source	Culture supernatant
Purification	-----
Buffer	Supernatant with 0.002% $\text{NaN}_3$ as preservative
Concentration	1 mg/mL
Volume	50 $\mu\text{L}$
Label	Unlabeled
Specificity	Equine Endoglin
Storage	Store at 4°C
Reference	-----

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Recommended Dilutions	Western blotting (1 : 1,000) , Immunohistochemistry (1 : 50), ELISA (1 : 100,000)
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Other applications have not been tested or not reactive.  
Optimal dilutions/concentrations should be determined by the end user.

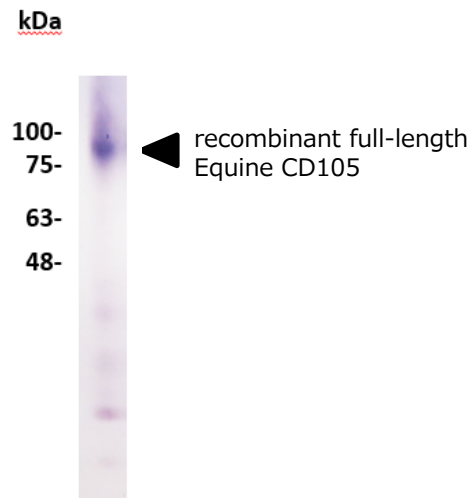


Fig 1. Western blot analysis

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Blocking : 0.5% Casein / Tris-saline (150 mM NaCl/10 mM Tris-HCl, pH 7.6)

Primary antibody : 0.5% Casein / Tris-saline (150 mM NaCl/10 mM Tris-HCl, pH 7.6)

Secondary antibody : ALP-labeled 2nd antibody in PBS-T

Detection : NBT/BCIP

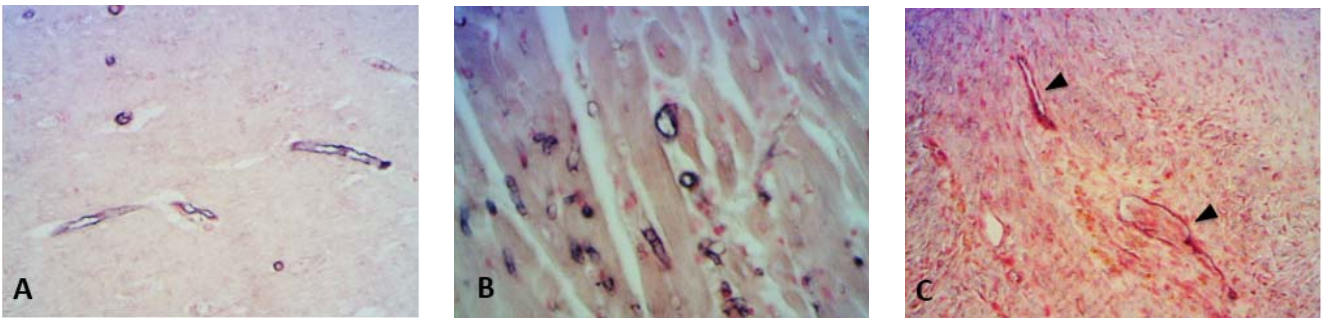


Fig 2. Immunohistochemistry

All tissues were fixed with 10% formalin and embedded in paraffin wax. After deparaffinization, sections were pretreated with 6 M urea/50 mM Tris-HCl, pH 7.6 for 30 min at room temperature.

A. The antibody reacted endothelial cells in capillary vessels of equine normal brain (ALP-labeled 2nd antibody/nuclear fast red).

B. The antibody reacted endothelial cells in capillary vessels of the mitral bulb in equine normal heart (ALP-labeled 2nd antibody/nuclear fast red).

C. The antibody reacted endothelial cells in newly generated capillary vessels (neovascularization) observed in granulation tissue in equine healing tendon (arrowheads) (ALP-labeled 2nd antibody/nuclear fast red).

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