www.innocyto.com.cn

InnoCyto Inc.

15375 Barranca Pkwy, Suite I-103
Irvine, CA 92618
Email: info@innocyto.com
Phone: 1-949-418-7543

Anti-Human VEGF (Fab) (Ranibizumab Biosimilar)

Catalog Number:	505101, 505102
Size:	2 mg, 10 mg
Target Name:	Vascular endothelial growth factor, VEGF-A, VEGFA, VEGF
Regulatory Status:	RUO

PRODUCT DETAILS

Clone:	Ranibizumab
Application:	Flow cytometry, animal model study
Reactivity:	Human
Format:	Liquid
Product Description:	Anti-Human VEGF (Fab) (Ranibizumab Biosimilar)
Isotype:	Human IgG1
Clonality:	Recombinant
Immunogen:	Human VEGF
Species specificity:	Human
Purity:	>95% by reducing SDS-PAGE
Grade:	In vivo
Storage Conditions:	4ºC
Maximal Shelf Life:	12 months
Synonyms:	VEGF-A
RRID:	AB_3739327
Antibody Type:	Recombinant
Reactivity:	Human

BACKGROUND INFORMATION

Ranibizumab is a recombinant humanized monoclonal antibody fragment (Fab) designed to specifically bind and neutralize vascular endothelial growth factor A (VEGF-A). Structurally, it is a single antigen-binding fragment derived from the same parent murine antibody as bevacizumab but lacks the Fc (fragment crystallizable) region, which distinguishes it from full-length immunoglobulin G (IgG) antibodies. The absence of the Fc domain results in a smaller, monovalent molecule with a molecular weight of approximately 48 kilodaltons (kDa).

The molecular structure of Ranibizumab comprises a variable heavy (VH) domain and a variable light (VL) domain connected to their respective constant regions (CH1 and CL). Together, these domains form a single high-affinity antigen-binding site, with

complementarity-determining regions (CDRs) mediating specific binding to VEGF-A isoforms, including VEGF121, VEGF165, and VEGF189. The CDRs establish strong non-covalent interactions with residues located within the heparin-binding domain of VEGF-A. This high-affinity binding occurs in the picomolar range, effectively sequestering VEGF-A and preventing it from binding to its cell surface receptors, VEGFR-1 (Flt-1) and VEGFR-2 (KDR). The disruption of VEGF-receptor interactions inhibits downstream receptor dimerization and phosphorylation, blocking intracellular signaling pathways such as MAPK and PI3K-AKT that mediate vascular permeability, endothelial proliferation, and angiogenesis in experimental systems.

Functionally, Ranibizumab's small molecular size enhances tissue penetration and allows rapid diffusion through extracellular matrices, making it suitable for localized VEGF inhibition at target sites. Although it lacks the Fc fragment, and thus cannot engage Fc gamma receptors (FcγRs) or activate complement, it retains high structural stability and solubility due to optimized domain folding and disulfide bridge formation. Overall, Ranibizumab exemplifies precision antibody engineering, combining molecular compactness and epitope specificity to investigate VEGF-mediated signaling and angiogenic processes in molecular biology research.

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