

# Anti-Mouse IgG kappa, AlpSdAbs<sup>®</sup> VHH (iFluor647)

## Summary

<b>Code</b>	001-106-009
<b>Immunogen</b>	Fab region of Mouse IgG
<b>Host</b>	Alpaca pacous
<b>Isotype</b>	VHH domain of alpaca IgG2b/2c
<b>Conjugate</b>	iFluor647(Ex:651nm, Em:667nm)
<b>Specificity</b>	Mouse IgG kappa chain
<b>Cross-Reactivity</b>	No cross-reactivity with rabbit, human, cynomolgus, rat, goat IgG
<b>Purity</b>	Recombinant Expression and Affinity purified
<b>Concentration</b>	0.5mg/mL
<b>Formation</b>	Liquid, 10mM PBS (pH 7.5), 0.05% sucrose, 0.1% trehalose, 0.01% proclin300, 50% glycerol
<b>Storage</b>	Store at -20 °C(Avoid freeze / thaw cycles) , Protect from light.

## Description

Anti-Mouse IgG kappa, AlpSdAbs<sup>®</sup> VHH(iFluor647) is designed for detecting mouse IgG kappa chain specifically, and Anti-Mouse IgG kappa, AlpSdAbs<sup>®</sup> VHH(iFluor647) is useful for super-resolution microscopy. Anti-Mouse IgG kappa, AlpSdAbs<sup>®</sup> VHH(iFluor647) is based on recombinant single domain antibody to mouse IgG kappa chain coupled to iFluor647. Based on immunoelectrophoresis and/or ELISA, Anti-Mouse IgG kappa, AlpSdAbs<sup>®</sup> VHH(iFluor647) reacts with the mouse IgG kappa chain selectively, no reactivity with the mouse IgG lambda chain or heavy chain of mouse IgG.

## Background

VHH are single-domain antibodies derived from the variable regions of heavy chain of Camelidae immunoglobulin. The size of VHH is extremely small(<15KDa) compared to other forms of antibody fragment, which significantly increase the permeability of VHH.

The smaller size of the VHH decreases linkage error and increases staining accuracy effectively. Standard immunodetection approaches use typically a primary antibody (1.Ab) which binds the protein of interest (POI) and a secondary antibody (2.Ab) that binds to the 1.Ab and carries a detection element. The complex formed by the primary antibody and the secondary antibody (1.Ab-2.Ab) is widely used because it is a cost effective and flexible approach since only the 2.Abs need to be coupled to the detection element. However, the use of this complex carries some relevant limitations. The 1.Ab-2.Ab can measure up to 30 nm, leading to a large distance between the targeted molecule and the detection element, causing the so called “linkage” or “displacement” error. While this might not influence the results in some applications (e.g. epifluorescence, ELISA or FACS), it is of major relevance for super-resolution microscopy techniques where the localization precision can be as high as 1 nm. The linkage error can be reduced by using directly labelled small affinity probes like camelid single domain antibodies (sdAbs) also known as nanobodies (Nbs), which have sizes below 3 nm.

## Benefits

High lot-to-lot consistency  
 Increased sensitivity and higher affinity  
 Animal-free production

## Suggested Working Concentration

<b>ELISA</b>	1:10000-1:50000
<b>WB</b>	1:10000-1:50000
<b>ICC/IF</b>	1:200-1:2000
<b>Flow Cyt</b>	1:200-1:2000

Super-resolution microscopy

Dilution factors are presented in the form of a range because the optimal dilution is a function of many factors, such as antigen density, permeability, etc. The actual dilution used must be determined empirically.

This product is for research use only and is not approved for use in humans or in clinical