



**Affinity  
Immuno**

**AffinityImmuno Inc  
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Canada  
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**Catalog:** ab-94-181 (chicken anti Rituximab idiotyp)

**Description:** Chicken polyclonal anti-Rituximab idiotyp (IgY)

**Lot:** 21-01-194A

**Product:**

Target:	Rituximab idiotyp
Host species:	Chicken
Volume:	100µl
Concentration:	1mg/ml
Total protein:	100µg
Formulation:	PBS, with 0.02% NaN <sub>3</sub> , pH7

**Production:**

Affinity purified over Rituximab resin, and then depleted using human IgG resin to remove all non anti-idiotyp reactivity. Specificity tested by comparing binding to Rituximab vs human IgG1 (Figure 1 below).

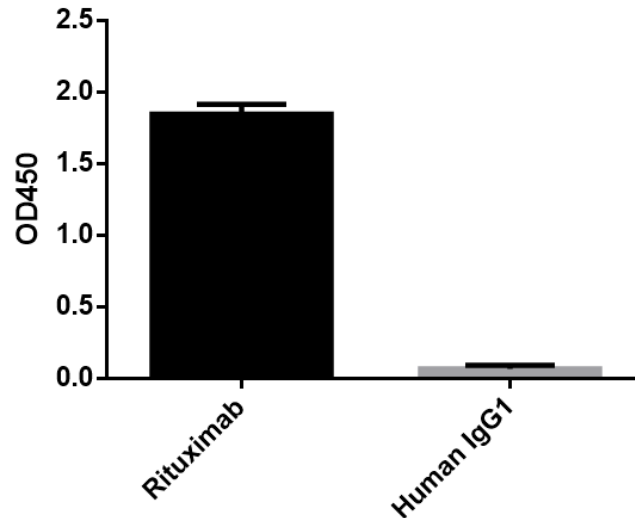
**Isotype:**

Chicken IgY

**Applications:**

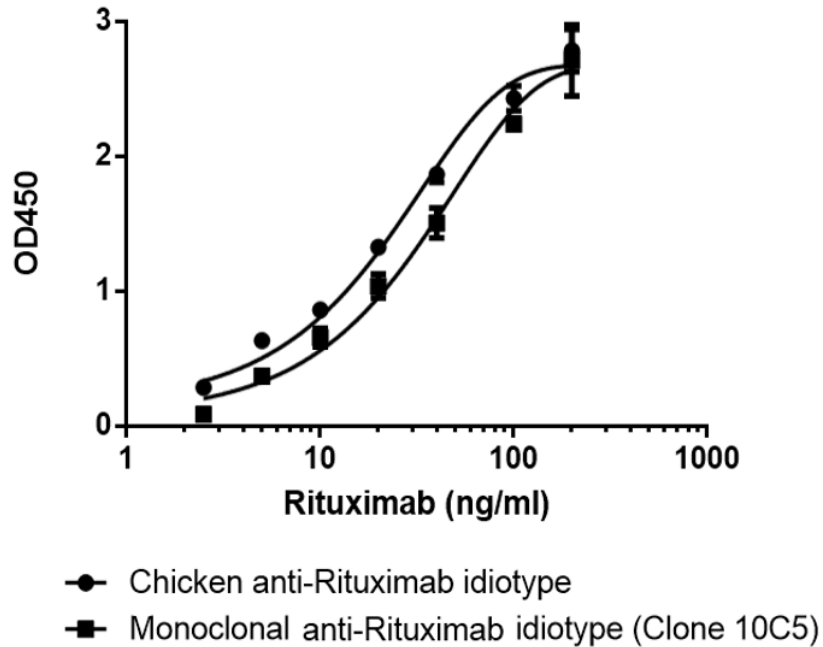
ELISA capture, blocking.

### Binding of chicken anti-Rituximab idiotype to Rituximab vs human IgG1



**Figure 1, Binding of anti-Rituximab is only to the idiotype and not the whole IgG molecule.** Anti-Rituximab idiotype was tested for binding to Rituximab and human IgG1 by indirect ELISA. Rituximab or human IgG1 was coated onto ELISA plates at equivalent concentrations and blocked using 2% BSA in PBS. Chicken anti-Rituximab idiotype was diluted to 0.25ug/ml and applied in triplicate to the coated wells and incubated for 1 hour at room temperature. After washing the wells, peroxidase conjugated detection antibody (Rabbit anti chicken IgY) was applied to the wells at a concentration of 5ng/ml and incubated for 1 hour at room temperature. After a final wash, the wells were developed with TMB and absorbance at 450nm was measured. These results show that the anti-Rituximab idiotype antibody is strongly specific to the Rituximab binding site (idiotype), and has negligible binding to the conserved regions of the IgG1 protein.

## Comparing chicken anti-Rituximab idiotypic to monoclonal anti-Rituximab idiotypic in Rituximab sandwich ELISA.



**Figure 2. Comparison of a chicken anti-Rituximab idiotypic and monoclonal anti-Rituximab idiotypic (Clone 10C5) for use in Rituximab capture ELISA.** Each antibody (chicken anti-Rituximab idiotypic vs. monoclonal antibody 10C5) was coated onto polystyrene ELISA plates at equivalent concentrations for comparison in a Rituximab capture ELISA. Rituximab was spiked into human serum at concentrations from 200ng/ml down to 2.5ng/ml and applied to each ELISA plate in duplicate and incubated for 1 hour at room temperature. After washing, peroxidase conjugated detection antibody (anti-human IgG1) was applied to the wells at a concentration of 650pg/ml and incubated for 1 hour at room temperature. After a final wash, the wells were developed with TMB and absorbance at 450nm was measured. These results show that the chicken anti-Rituximab idiotypic antibody performs better as a capture antibody than the competing monoclonal antibody (10C5) and provides a lower limit of detection.