# Monoclonal Anti-SN38 Antibody (MALS verified)

Catalog # SN8-S223



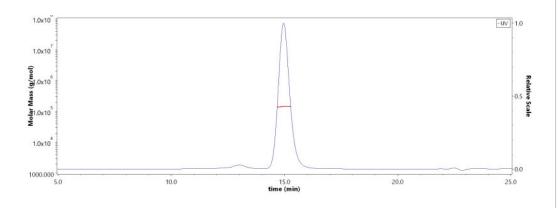
Source	Formulation
Monoclonal Anti-SN38 Antibody, is produced from a hybridoma resulting from fusion of SP2/0 myeloma and B-lymphocytes obtained from a mouse immunized with SN38.	Lyophilized from 0.22 $\mu$ m filtered solution in PBS, pH7.4 with trehalose as protectant.
Isotype	Contact us for customized product form or formulation.
Mouse IgG1/kappa	Reconstitution
Specificity	Please see Certificate of Analysis for specific instructions.
Specifically recognizes the target-SN38.	For best performance, we strongly recommend you to follow the reconstitution
Purity	protocol provided in the CoA.
	Storage
>95% as determined by SDS-PAGE.	For long term storage, the product should be stored at lyophilized state at -20°C
>95% as determined by SEC-MALS.	
Endotoxin	Please avoid repeated freeze-thaw cycles.
Less than 1.0 EU per $\mu$ g by the LAL method.	This product is stable after storage at:
	• -20°C to -70°C for 12 months in lyophilized state;
	• -70°C for 3 months under sterile conditions after reconstitution.

#### **SDS-PAGE**

kDa	М	R
116.0		
66.2		
45.0	-	
35.0		
25.0		-
18.4	-	
14.4	_	

Monoclonal Anti-SN38 Antibody on SDS-PAGE under reducing (R) condition. The gel was stained with Coomassie Blue. The purity of the protein is greater than 95%.

## SEC-MALS



The purity of Monoclonal Anti-SN38 Antibody (Cat. No. SN8-S223) is more than 95% and the molecular weight of this protein is around 135-160 kDa verified by SEC-MALS. <u>Report</u>

### **Bioactivity-Elisa**



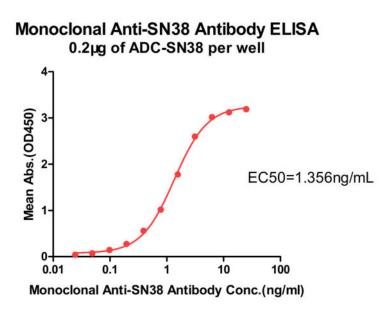
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Immobilized ADC-SN38 at 2  $\mu$ g/mL (100  $\mu$ L/well) can bind Monoclonal Anti-SN38 Antibody (Cat. No. SN8-S223) with a linear range of 0.05-1.56 ng/mL (QC tested).

#### Background

SN-38 is an antineoplastic drug. It is the active metabolite of irinotecan (an analog of camptothecin - a topoisomerase I inhibitor) but has 1000 times more activity than irinotecan itself. In vitro cytotoxicity assays show that the potency of SN-38 relative to irinotecan varies from 2- to 2000-fold. SN38 is formed via hydrolysis of irinotecan by carboxylesterases and metabolized via glucuronidation by UGT1A1. The variant of UGT1A1 in  $\sim$ 10% of Caucasians which leads to poor metabolism of SN-38 predicts irinotecan toxicity, as it is then less easily excreted from the body in its SN-38 glucuronide form. SN-38 and its glucuronide are lost into the bile and intestines. It can cause the symptoms of diarrhoea and myelosuppression experienced by  $\sim$ 25% of the patients administered irinotecan.

#### **Clinical and Translational Updates**

Please contact us via <u>TechSupport@acrobiosystems.com</u> if you have any question on this product.



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