

# Sorrento Potent Omicron Neutralizing Antibody (nAb) STI-9167 IV STI-9199 IN

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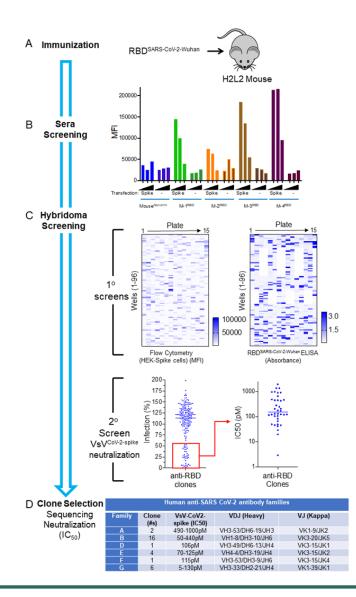
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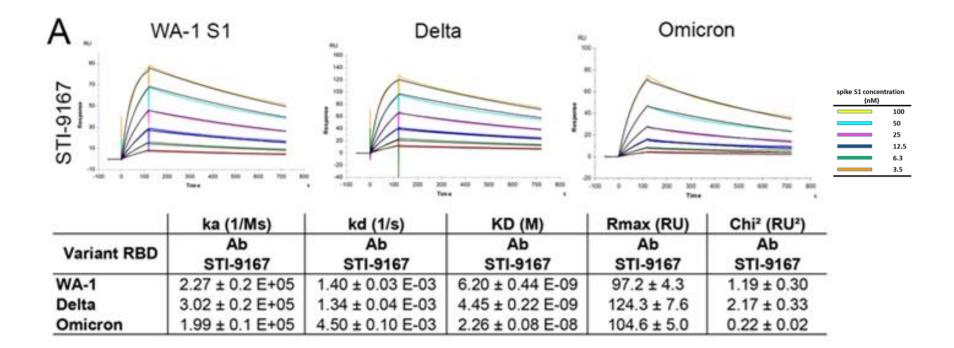


# Discovery of STI-9167





#### STI-9167 Binding to S1 Domain of SARS-CoV-2 Variants of Concern (VOCs)

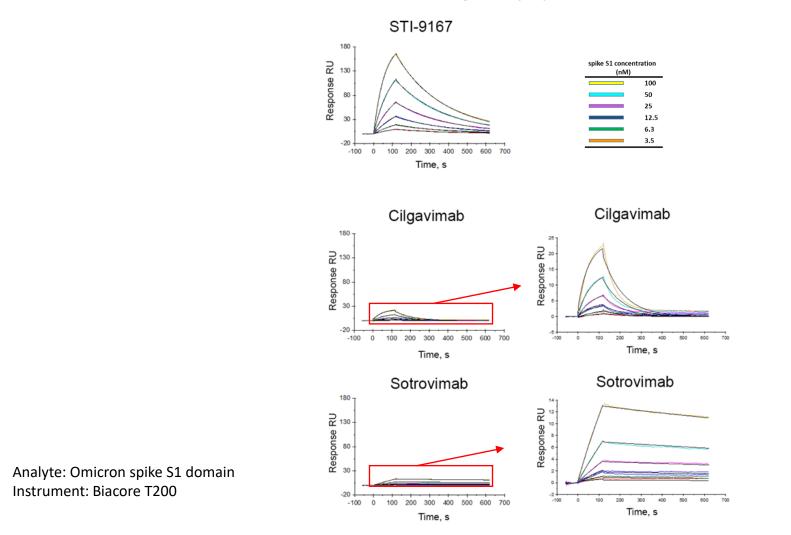


# Analyte: WA-1, Delta, or Omicron spike S1 domain Instrument: Biacore T200

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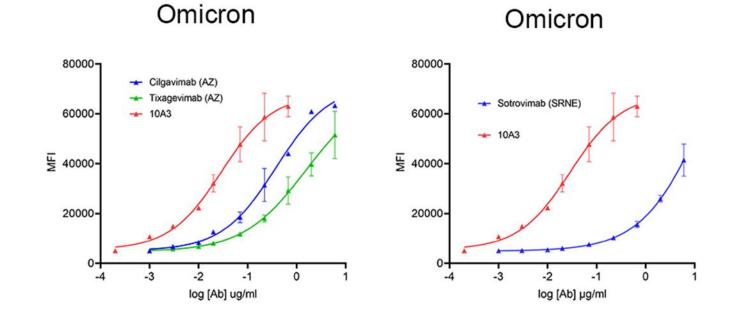
#### STI-9167 Binding Kinetics to Spike S1 Domain Compared to Those of EUA-approved nAbs

Binding Affinity Spike S1 Omicron





#### STI-9167 Binding to SARS-CoV-2 VOC Spike Proteins Expressed on HEK293 Cells



cell-expressed spike binding EC50 (μg/mL)		
Variant	Ab	
spike	STI-9167	
WA-1	0.025	
Delta	0.011	
Omicron	0.025	
Omicron +	0.023	
R346K		



#### **Omicron Variant and Omicron + R346K Mutation Infection Statistics**

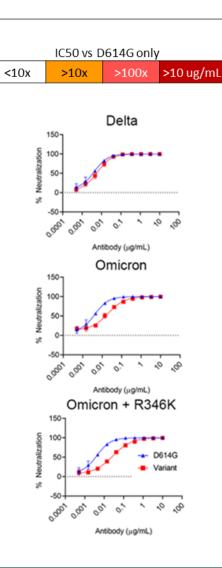
outbreak.info Cases & Deaths Variants Research Library API About	
	У f 🥶 in 🖾 📌 🔗
SARS-CoV-2 (hCoV-19) Mutation Reports Lineage Comparison C CHANGE LINEAGES O Updated 21h ago with 6,748,983 sequences from GISAID	Enabled by data from <b>GISAID</b> How to interpret these reports
dark mode include genes: ORF1a ORF1b S ORF3a E M ORF6 ORF7a ORF7b ORF7b ORF8 N ORF10	Min. mutation     Min. sequences       prevalence     per lineage       0.2 %     1
As a newly designated Variant of Concern, Omicron / B.1.1.529 is highly in flux. Expect the characteristic mutations associated with Omicron and its prevalence across locations to change as more s daily with new data from GISAID.	equences are reported. outbreak.info updates
Mutation prevalence across lineages Mutations with > 0.2% prevalence in at least one lineage. S	Page Variant / Variant / Variant / Mutation of Mutation of I00% Interest
Most Omicron sequences also contain a 3 amino acid insertion (EPE) at position 214 in the Spike protein.	data processing pipeline, and we encourage you
$ \begin{array}{c} \begin{array}{c} \begin{array}{c} \begin{array}{c} \begin{array}{c} \begin{array}{c} \end{array}{} \end{array}{} \end{array}{} \end{array}{} \end{array}{} \end{array}{} \end{array}{} \end{array}{} \end{array}{} \end{array}{$	(249,096 seqs)

# To date: No SARS-CoV-2 nAb approved or in development has been shown to have potent neutralization activity (≤ 100 ng/ml) against the Omicron + R346K variant



### SARS-CoV-2 Spike-Pseudotyped VSV Neutralization Assay

Variant lineage	nAb IC <sub>50</sub> (μg/ml)	
	STI-9167	
D614G	0.0036	
Alpha (B.1.1.7)	0.0029	
Beta (B.1.351)	0.0195	
Gamma (P.1)	0.0063	
Delta (B.1.617.2)	0.0054	
Delta Plus (B.1.617.2.1)	0.0033	
Epsilon (B.1.429)	0.0040	
Zeta (P.2)	0.0034	
lota (B.1.526)	0.0194	
lota (B.1.526.2)	0.0024	
Kappa (B.1.617.1)	0.0090	
Lambda (C.37)	0.0027	
Mu (B.1.621)	0.0186	
Omicron (B.1.1.529)	0.0148	
Omicron+R346K (B.1.1.529)	0.0239	

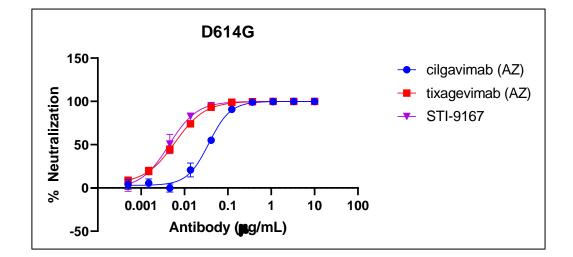


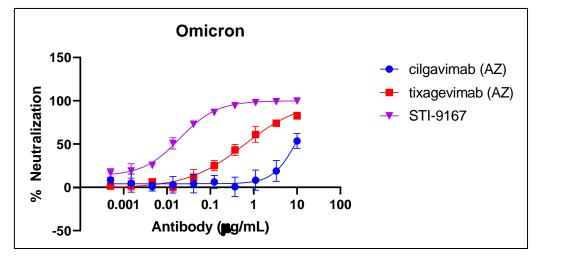


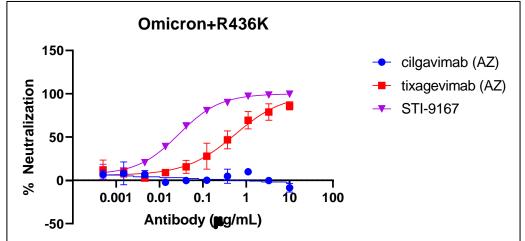
### STI-9167 Superior Neutralization Activities Against Omicron and Omicron+R346K Variants in SARS-CoV-2 Spike-Pseudotyped VSV Neutralization Assays

Variant Lineage	STI-9167	Cilgavimab (AZ)	Tixagevimab (AZ)
D614G	0.0036	0.0353	0.0056
Omicron (B.1.1.529)	0.0148	9.105	0.6386
Omicron+R346K (B.1.1.529)	0.0239	>10	0.4696

IC50 (ug/mL)

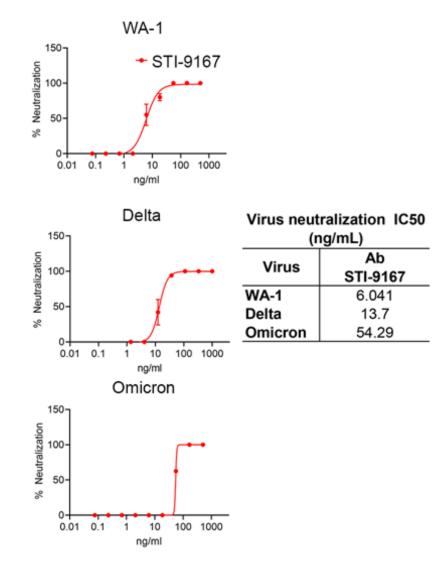








### *Neutralization Activity of STI-9167 in the SARS-CoV-2 Live Virus Neutralization Assays*





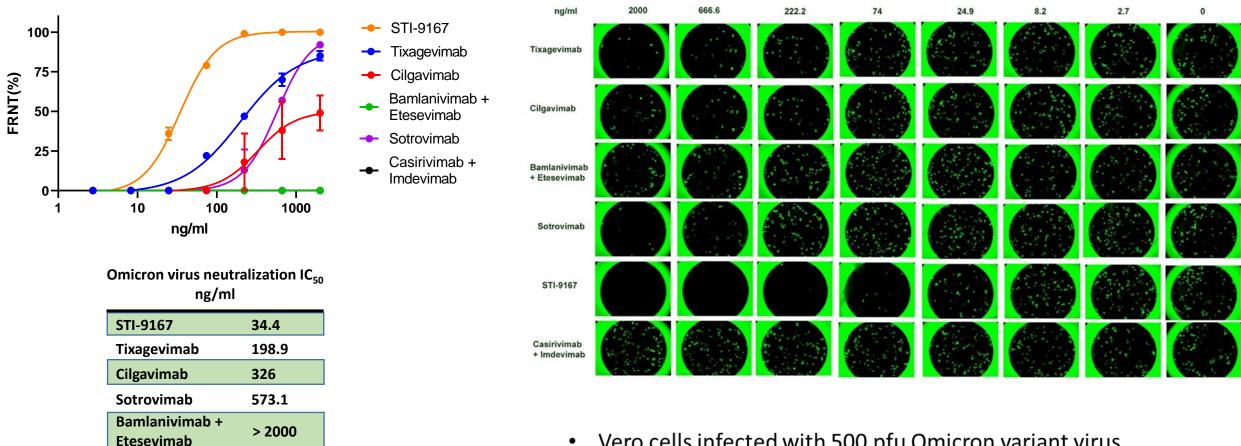
# Superior Neutralization Activity of STI-9167 as Compared to EUA-approved Neutralization Antibodies (nAbs) in the SARS-CoV-2 Live Omicron Virus Neutralization Assays

Omicron

Casirivimab +

Imdevimab

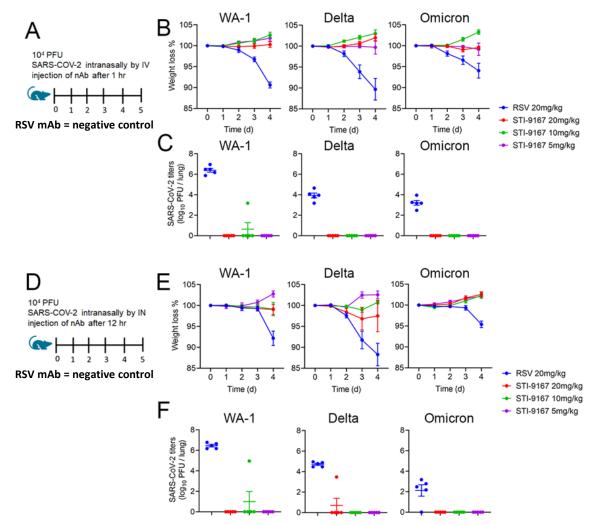
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- Vero cells infected with 500 pfu Omicron variant virus
- Infection visualized with anti-SARS-CoV-2 nucleoprotein antibody



### STI-9167 Neutralizing Activity Following IN or IV Administration in the K18-hACE2 Transgenic Mouse Model of COVID-19





# STI-9167 GMP Manufacturing and Clinical Developmental Status

- ✓ STI-9167 GMP Master Cell Bank generation completed
- STI-9167 GMP Drug Product to supply Phase 1/2 studies formulated for intranasal and intravenous dosing has been F/F
   Already manufactured enough drug substance for 100,000's of intranasal doses
- $\checkmark$  IND-enabling preclinical safety and toxicology studies completed
- IND submission in early February
- Phase 1 clinical trials planned in healthy individuals and infected patients for both intranasal formulation (COVIDROPS) and intravenous formulation (COVISHIELD)

