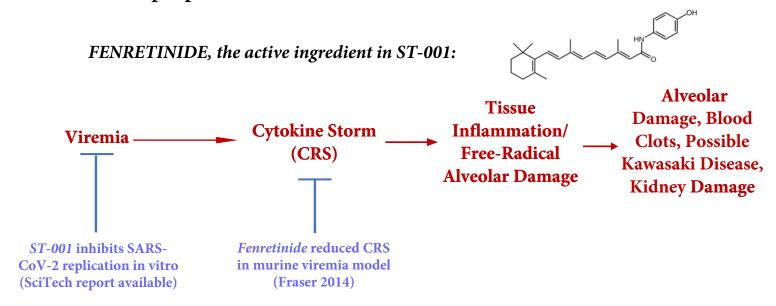


## ST-001 (nanoFENRETINIDE IV): Repurposed Retinoid Treatment for COVID-19 ARDS



## ST-001 (nanoFENRETINIDE IV) Resumé

- Repurposing ST-001 from oncology to COVID-19
- Cross-filing on SciTech's current oncology IND #135475 to expedite clinical entry
- Utilize ST-001 manufacturing process from IND #135475
- ST-001 is SciTech's patented (expires 2030) formulation to administer high strength fenretinide via 4 hr infusion
- Only triglyceride-free IV formulation of fenretinide, avoiding other product's hypertriglyceridemia toxicity
- Phase 1 starting dose level targets a blood level above the  $IC_{50}$  of 2  $\mu$ M for SARS-CoV-2
- 20 μM blood levels of fenretinide have demonstrated safety in prior trials of other formulations
- Fenretinide is a safer analog of Vitamin A
- Compatible with monoclonal antibody therapies in published preclinical studies

## **CLINICAL DEVELOPMENT OF ST-001**

IND Filed/ST-001 Manufacturing	Phase 1a	Phase 1b	Phase 2
	Rapid Dose-Finding Design:  Establish Safety  Identify Treatment Dose  Proof of Concept  Reduced IL-6  Increased IL-22BP  Explore Efficacy  Reduced D-dimer  Negative COVID-19 test	Phase 1a Treatment Dose:  Confirm Safety  Explore Efficacy  Respiratory status  Reduced mortality  Reduced ICU LOS  Improved lab values  Confirm POC  IL-6  IL-22BP	Treatment Dose:  • Efficacy/outcome endpoints after FDA consultation
		<b>↓</b>	
		Phase 1	Phase 2
ST-001 COMBINATION THERAPIES:		ST-001 Treatment Dose with Remdesivir, Kineret, Sylvant, or Actemra: Confirm Safety Explore Efficacy Select best combination	ST-001 Combination Treatment:  • Efficacy/outcome endpoints after FDA consultation

