



Immunic
THERAPEUTICS

Immunic Therapeutics Targeting COVID-19 with IMU-838

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→ A further list and descriptions of these risks, uncertainties and other factors can be found in the section captioned “Risk Factors,” in the Company’s Annual Report on Form 10-K for the fiscal year ended December 31, 2019, filed with the Securities and Exchange Commission (“SEC”) on March 16, 2020, the Company’s Quarterly Report on Form 10-Q for the quarter ended March 31, 2020, filed with the SEC on May 8, 2020, and in the Company’s subsequent filings with the Securities and Exchange Commission. Copies of these filings are available online at www.sec.gov or ir.immunic-therapeutics.com/sec-filings and on request from Immunic.

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Development Pipeline

Program	Indication	Target	Preclinical	Phase 1	Phase 2	Phase 3
IMU-838	Multiple Sclerosis	DHODH	Completed or ongoing	Completed or ongoing	Completed or ongoing	In preparation or planned
	Ulcerative Colitis	DHODH	Completed or ongoing	Completed or ongoing	Completed or ongoing	
	Crohn's Disease	DHODH	Completed or ongoing	Completed or ongoing		
	PSC	DHODH	Completed or ongoing	Completed or ongoing	Completed or ongoing	Investigator-Sponsored Trial performed at Mayo Clinic / NIH
	COVID-19	DHODH	Completed or ongoing	Completed or ongoing	Completed or ongoing	In preparation or planned
IMU-935	Psoriasis	ROR γ t	Completed or ongoing	Completed or ongoing		
	Orphan AI Diseases	ROR γ t	Completed or ongoing	In preparation or planned		
IMU-856	GI	Intestinal Barrier Function	Completed or ongoing	In preparation or planned		

■ Completed or ongoing

■ In preparation or planned

◆ Top-line data from the phase 2 trial of IMU-838 in relapsing-remitting multiple sclerosis is expected to be available in the first half of August 2020

IMU-838: Triple Attack on COVID-19

IMU-838 is an advanced clinical drug candidate with attractive pharmacokinetic, safety and tolerability profile with more than **650 individuals exposed to date**



IMU-838 attacks COVID-19 disease by three complementary mechanisms:

- 1** Inhibition of **virus replication** by depletion of nucleotide pool
- 2** Insufficient first immune response due to SARS-CoV-2 encoded interferon antagonists
Induction of **innate immune response** by DHODH inhibition independent of interferon signaling
- 3** Excessive activation of adaptive immune response – “cytokine storm”
Inhibition of “overreacting”, **cytokine high** producing immune cells



Thank you!

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