



Immunic
THERAPEUTICS

Immunic Therapeutics

Developing Selective Oral Drugs in Immunology

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BMO 2020 Prescriptions for Success Healthcare Conference

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Summary and Highlights



Advanced and well-balanced pipeline:
Three products in development



Shares outstanding: 12.8 million
(as of May 1, 2020)



Phase 2 data read-outs ahead:
Several clinical phase 2 trials with IMU-838 expected to read-out in the next couple of months



Cash position of **USD 18.6 million**
(as of March 31, 2020)
USD 40 million ATM in place



Phase 2 trial in COVID-19 ongoing
IMU-838 is a potential COVID-19 solution
Broad-spectrum antiviral activity



Raised approximately **USD 42 million** in April and June 2020, substantially extending **cash runway beyond important value inflection points**

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	
	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

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

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