

Our goal is to alleviate pandemic infectious diseases that impact our global community.

Executive Summary

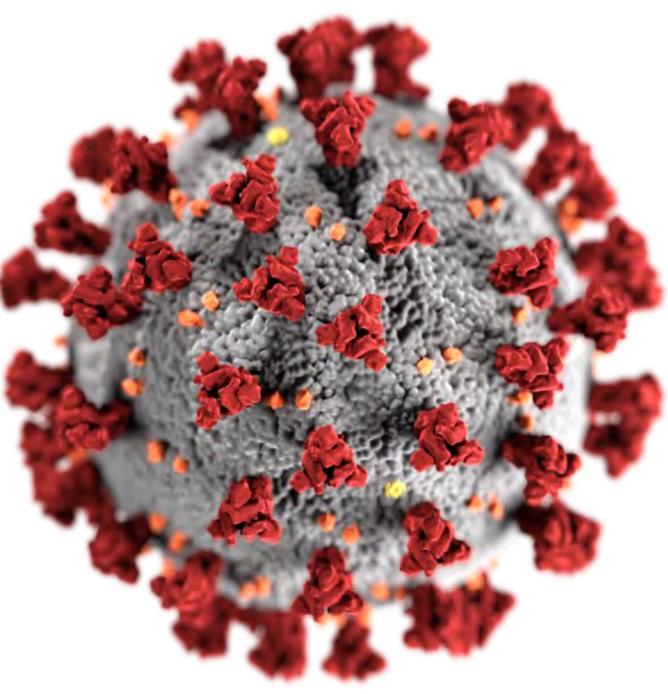
- COVI-001 has significant clinical efficacy, patent protection and first-to- market position
- COVI-001 satisfies clear unmet needs for patients and healthcare environment
- Ask: \$25M Series A fundraise

The Unmet Need

•There are no effective treatments to cure the sickest ICU patients with COVID-19

•Dexamethasone is the best available weapon.

•There are no FDA approved treatments for late-stage/severe COVID-19





The Unmet Need: Newly Reported Deaths Last 7 Days



11.05.22

A	I Europe	North Americ	a Asia	South America	Africa Ocea	ania				
#	Country, Other ↓1	Cases in the last 7 days	Cases in the preceding 7 days	Weekly Case % Change	Cases in the last 7 days/1M pop	Deaths in the last 7 days ↓	Deaths in the preceding 7 days	Weekly Death % Change	Deaths in the last 7 days/1M pop ↓↑	Population 1
	World	2,065,438	2,531,011	-18%		8,101	10,508	-23%		
1	<u>USA</u>	189,609	273,243	-31%	566	1,548	2,281	-32%	5	335,251,946
2	<u>Germany</u>	252,641	398,437	-37%	2,993	991	1,062	-7%	12	84,410,641
3	<u>UK</u>	17,660	37,167	-52%	257	737	1,003	-27%	11	68,720,327
4	<u>Russia</u>	42,362	52,659	-20%	290	497	580	-14%	3	146,080,815
5	Taiwan	204,620	243,770	-16%	8,555	455	438	+4%	19	23,917,605
6	France	153,743	224,810	-32%	2,343	404	503	-20%	6	65,611,363
7	<u>Japan</u>	375,907	272,236	+38%	2,994	387	364	+6%	3	125,571,177
8	<u>Italy</u>	110,988	208,501	-47%	1,842	335	559	-40%	6	60,253,923
9	Brazil	26,916	36,302	-26%	125	276	478	-42%	1	216,094,853
10	Philippines	6,461	9,040	-29%	57	245	253	-3%	2	112,994,709

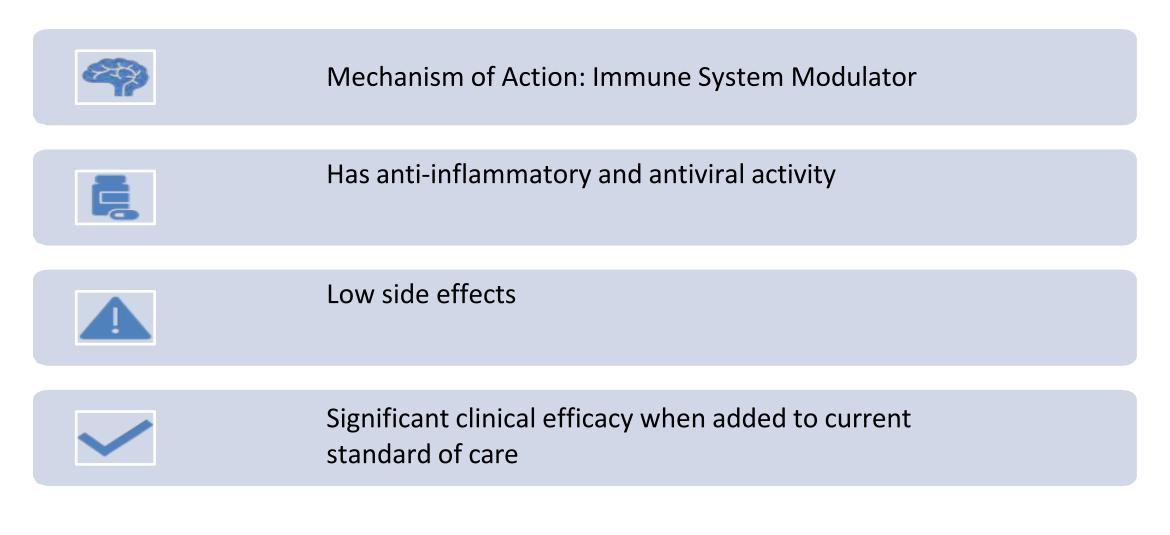


The Problem: COVID-19's Lethal Trigger

- Reduction of circulating viral load is needed
- Inflammation modulation is necessary



The Solution: COVI-001 Oral Fixed Dose Combination Therapy for Severe COVID-19



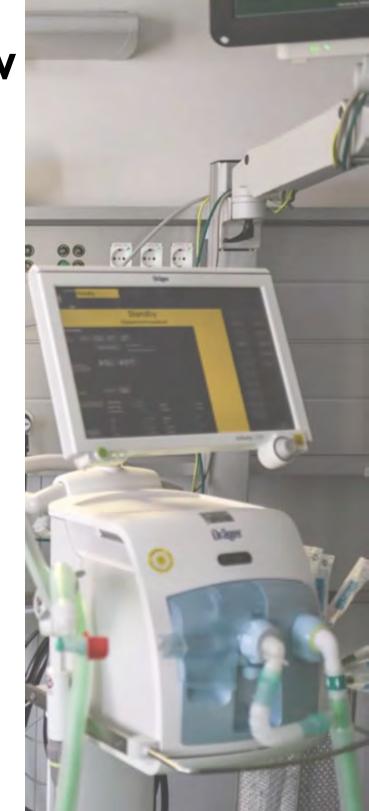


COVI-001: Retrospective Chart Review

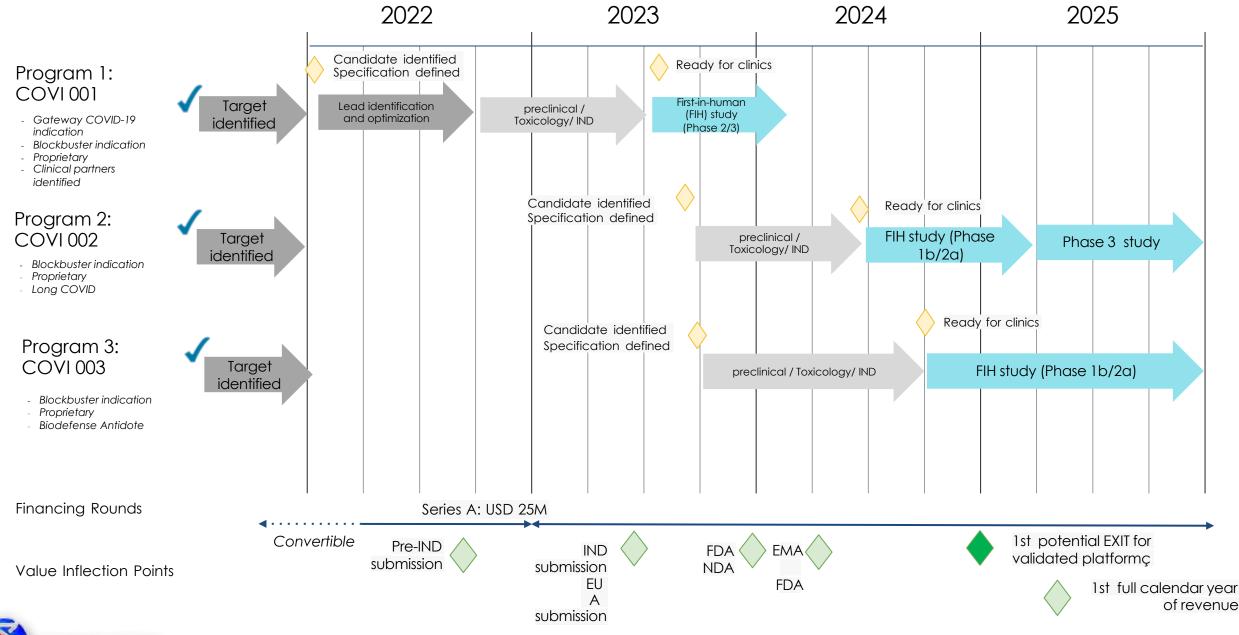
In a 2020 clinical study of 121 patients hospitalized for COVID-19- induced oxygen-hunger. 77 patients on one drug component of COVI-001 demonstrated:

- 68% reduction in mortality (p<0.02)
- 37% reduction in length of hospital stay (p<0.01)
- 93.5% survival rate (p<0.02)





Drug Development Program Timelines





Pipeline

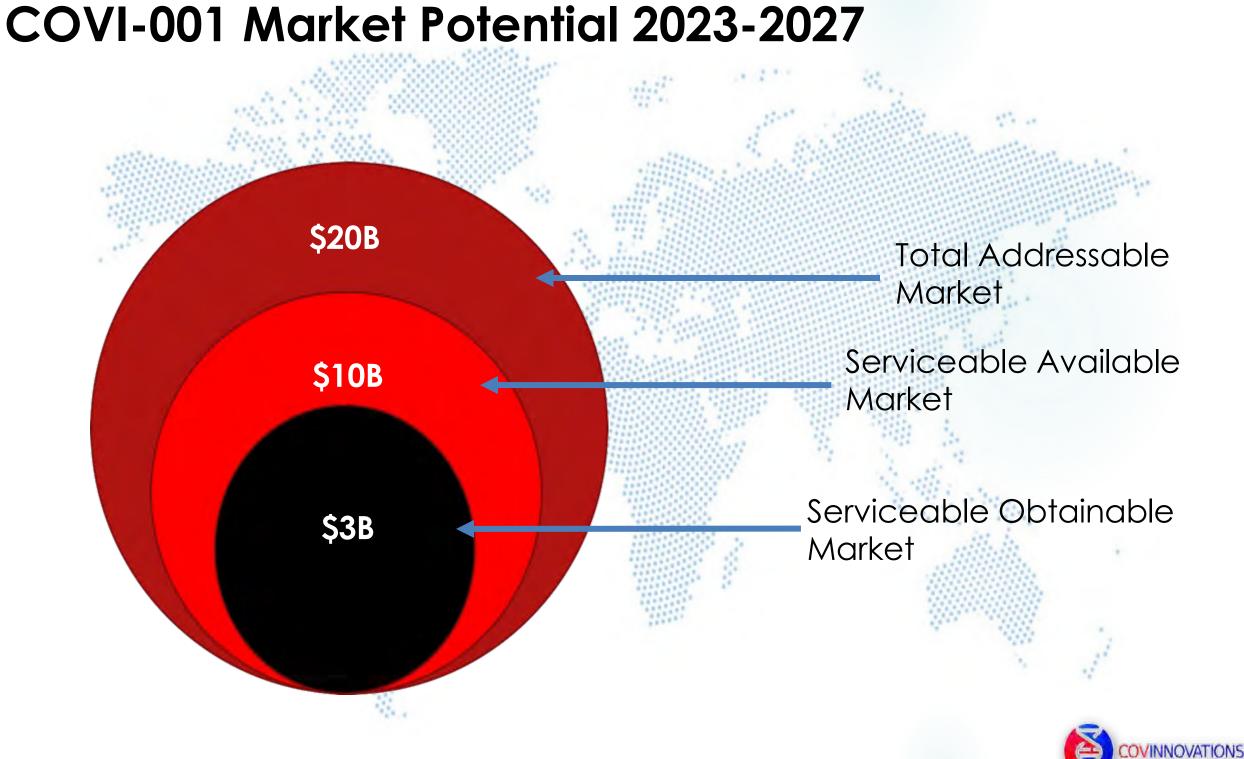
Strong IP Estate and First-to-Market

- Our unpublished human data, is the only human data using this API combination in COVID-19 patients, giving us first-tomarket advantage
- Our IP estate has multiple strong patent applications for use, • Mine (9) provisional utility patent applications;

 - One (1) US utility application; and
 - Two (2) International Patent Cooperation Treaty (PCT) applications
- Early and broad adoption as the standard of care is predicted



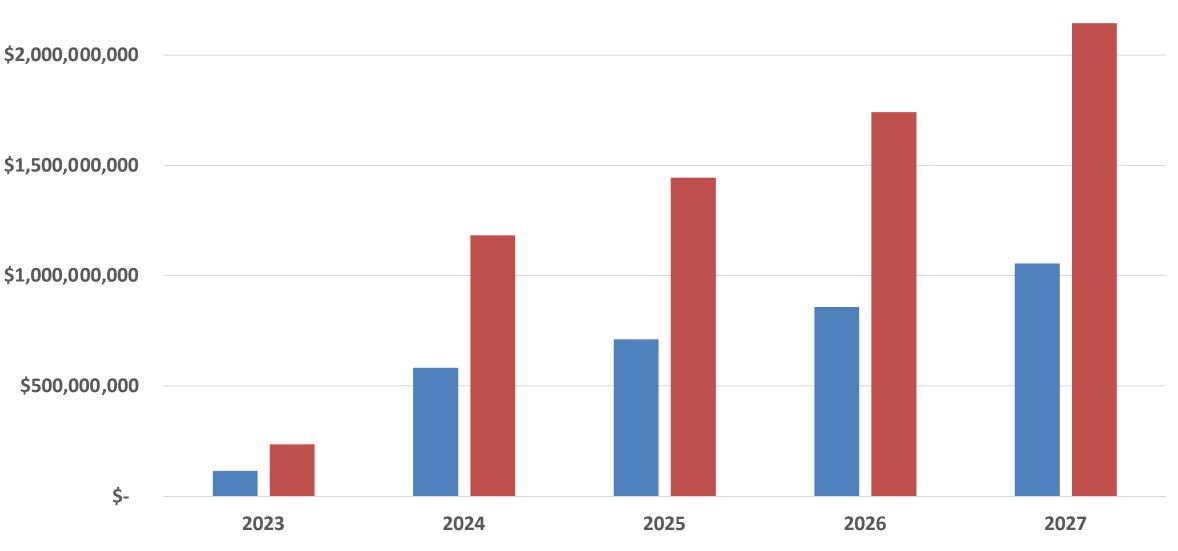




Global Hospital Revenue Forecast

US = Domestic ROW= Rest of World

\$2,500,000,000





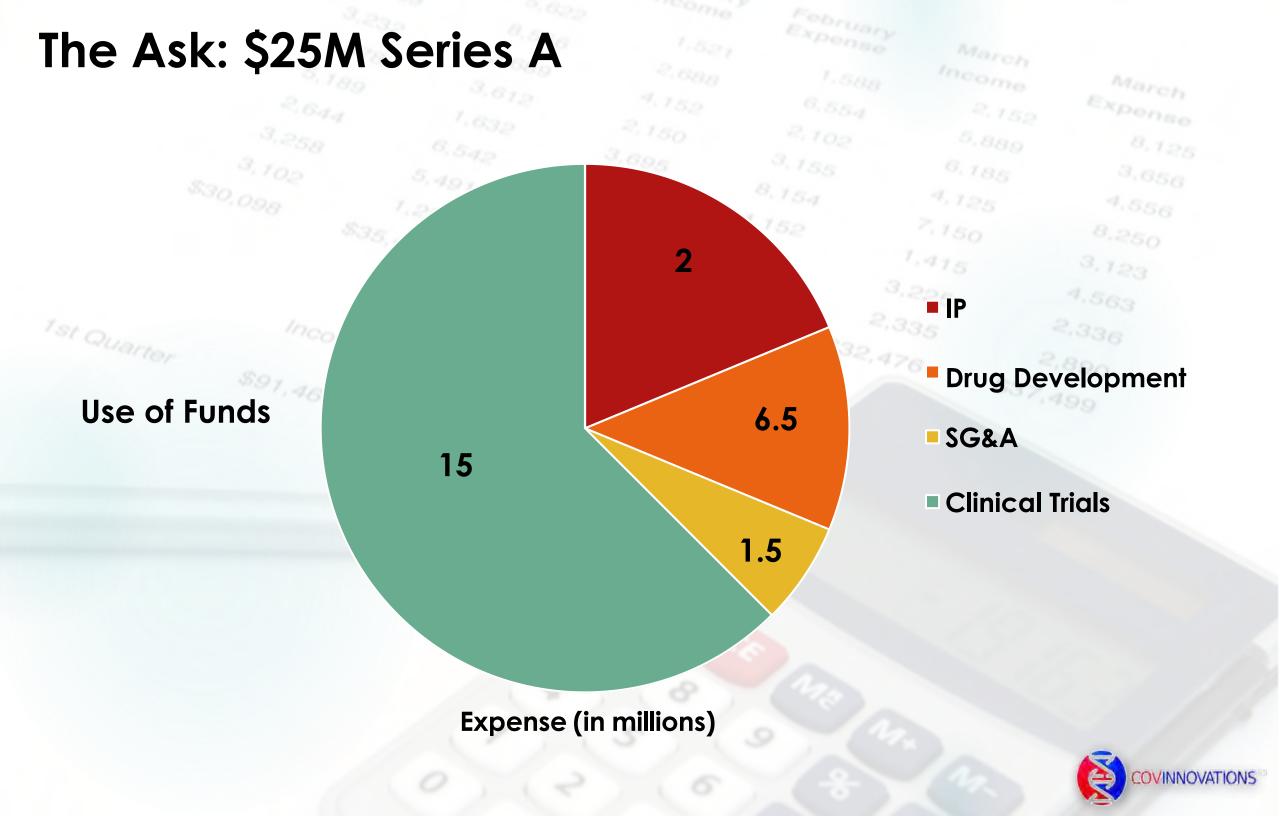
\$70B COVID-19 Therapeutic Global Market by 2030

2022 trending towards \$35B with 10% CAGR

Drug	Company	MOA	Indication	Admin	DDI	\$\$\$	'22 Annual Revenue
COV-001	CovInnovations	ISM	Severe	Oral	Low- Medium	\$500	\$1.9B***
Paxlovid	Pfizer	Antiviral	Mild to Moderate	Oral	High	\$529	\$22B
Molnupiravir	Merck	Antiviral	Mild to Moderate	Oral	Low	\$707	\$6B
Remdesivir	Gilead	Antiviral	Mild to Moderate	IV	Low	\$4,680	\$5B
Bebtelovimab	Eli Lilly	mAbs	Mild to Moderate	IV	Low	\$2,200	\$2.2B

Source: <u>https://aspr.hhs.gov/COVID-19/Therapeutics/Documents/side-by-side-overview.pdf</u> ***Note: first full calendar year revenue 2024



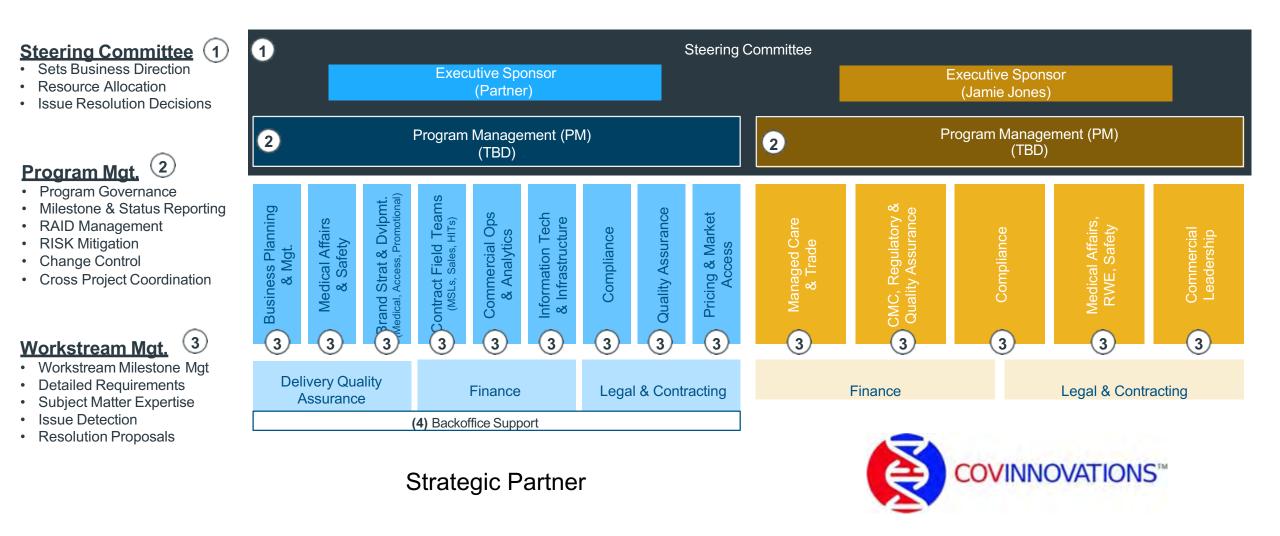


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COVID-19 EUA / BLA Go-to-Market Strategy

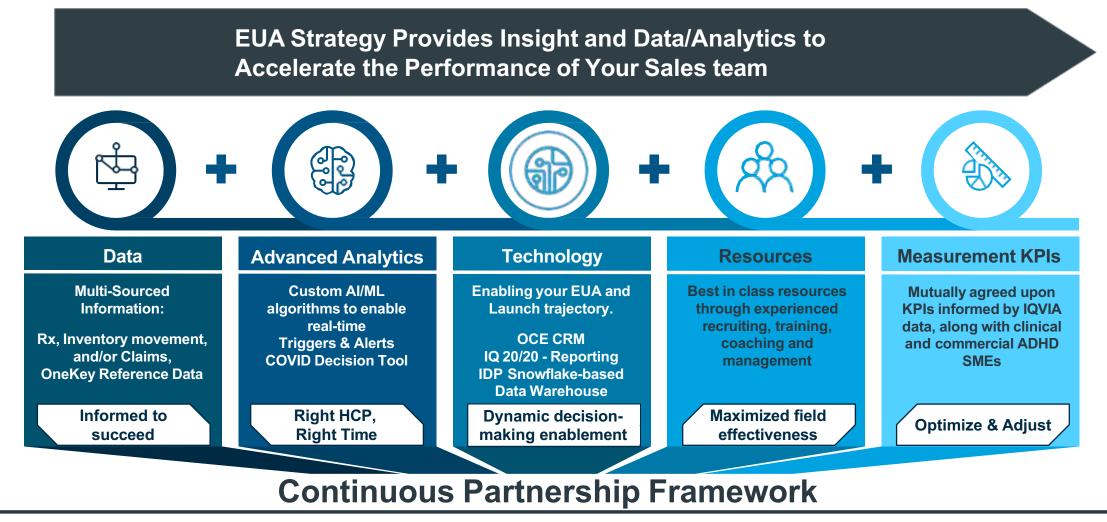
Engagement model ensures operational excellence to optimize EUA success

Centralized engagement model via PM Office ensures efficient execution of strategic vision



Create unique ability to bridge the EUA period to the commercial continuum brings inherent value

Partnership model fueled by Data / Analytics / Technology to help optimize Critical Success Factors



Strategy for EUA Activation helps deliver information & education to treaters so can be used with the patients who need it

EUA limits the deployed activities and establishes a leverageable foundation to support commercial launch.

Business Planning & Mgt.	Medical Affairs & Safety	Compliance	Pricing & Market Access	
 EUA Model Planning & CSFs Go-To-Market (Launch) Model Planning & CSFs PMO (Business & Technology) GTM HCP/HCO Engagement Strategy & Sourcing Model Awareness, Trial, and Usage Studies 	 KOL profiling & mapping Scientific comms. platform design Medical Information Call Center Product Quality Complaint Processing Pharmacovigilance & Safety Reporting Real World Evidence generation (post marketing study) 	 MLR Governance & Support Compliance SOP Assessment Transparency SOP Dev Transparency Reporting Fair Market Value HCP & Funding Process Design 	 Pricing & Contracting Strategy (& Targeting) Distribution Strategy Payer Positioning & messaging, Content Dev 3PL Data Aggregation 	

Commercial Operations & Analytics

Incentive Compensation Design &

EUA Data Analysis & KPI selection

(Virtual Ad-Boards KOLs, HCPs, Payers)

Data Strategy & Procurement

Promotional Segmentation,

Targeting & Alignments

Speaker Bureau & Events

Administration

Data Stewardship

•

Contract Field Teams & Support	Brand Strategy & Development				
MSL Team Field Sizing, Recruiting, Training	 Positioning, Messaging & Content Dev (MSLs, MI FAQs, HCP Web Site) 				
Ongoing Resource Mgt & Support	Product Website with HCP login if needed				
Med. Science Liaisons	Positioning, Messaging & Content Dev				
Access Advisor	(Product Web Site)				
Key Account Managers	Positioning, Messaging & Content				
Hospital Sales Reps	Dev (Promotional HCP & Payer)				
HCP Sales Reps	Digital Engagement Strategy (Medical)				
	Digital Engagement Strategy (Promo)				

Multi-channel mktg. design & execution

Information Technology & Infrastructure Support

- IT Roadmap
- CRM Implementation & Support
- Bl/Analytics, Field Reporting
 Implementation/Support
- MLR Content Management

Quality Assurance

- Quality SOPs
- Quality Management Automation

EUA Scope
 Launch Scope

Highlights of an EUA Activation Program

EUA priorities drive plans and preparations

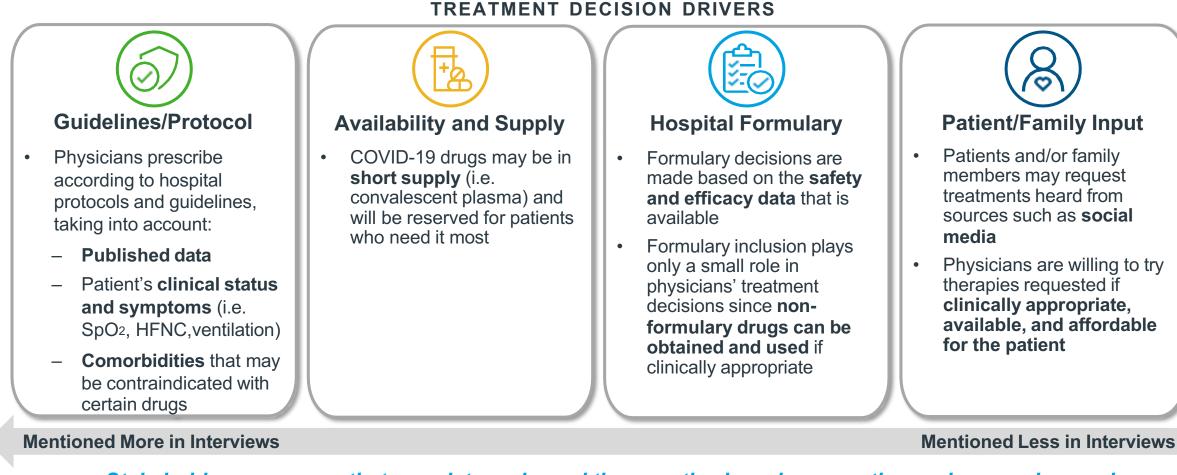
Program Component	Output or Description
1. Program Management & EUA activation plan	Overall Program management of the EUA program and delivery
2. KOL & Hospital Identification & Mapping	MSL call and contact targets
3. Pricing study and finalization	Pricing study, Product dossier, HEOR
4. EUA Product Website	Product Website
5. EUA HCP Registration Website	EUA program registration website
6. Medical Information Content and Call Center	Call Center, Collect product quality, AEs event details. Develop FAQ, Standard Responses, Approved Email content, etc.
7. Pharmacovigilance	Monitoring and reporting of PV events per regulatory requirements
8. CRM Systems setup	CRM to support MSL and KAM persona interactions
9. MLR Content and Processes	Develop Med-Legal-Regulatory Process and Service to support execution
10. Training Materials development/delivery	Product and Medical training for MSL, Hospital, & Home Healthcare Partner
11. Field Teams: MSL, KAM Recruiting and Hiring	Hire and rollout MSL and KAM teams
12. Trade Readiness & NAM Team Onboarding	Market Access Program Plan and implementation of syndicated NAM Team
13. EUA Analytics, MDM, and Reporting	Abbreviated Reporting and MDM stewardship to support EUA
14. Compliance Risk Assessment & Policies	Develop risk assessment program and standard operating procedures
15. Hospital to Home Healthcare Continuity of Care Strategy	Develop overall hospital engagement / HHC strategy to support continuity of care

EUA Activation Timeline & Milestones for COVI-001

purposes. Co	ntracting	Activities	Start			EUA			MSLs	Product La	unc
	Month 1	Month 2	Month 3	Month 4	Month 5	Month 6	Month 7	Month 8	Month 9		
. EUA activation plan & Program Management											
2. KOL & Hospital Identification & Mapping			٠								
 Pricing study and finalization 				٠							
4. EUA Product Website					•						
5. EUA HCP Registration Website					•						
6. Medical Information Content and Call Center						•	.			Commercial	
7. Pharmacovigilance						•	٠			Preparation	
3. CRM Systems setup				:				•			
). MLR Content and Processes							•				
0. Training Materials development/delivery							•				
1. Field Teams: MSL, KAM Recruiting and Hiri							•				
2. Trade Readiness & NAM Team Onboarding			•	•							
3. EUA Analytics, MDM, and Reporting								•	:		
4. Compliance Risk Assessment & Policies				•		:					
 Hospital to Home Healthcare Continuity of C Strategy 	Care					•	•				

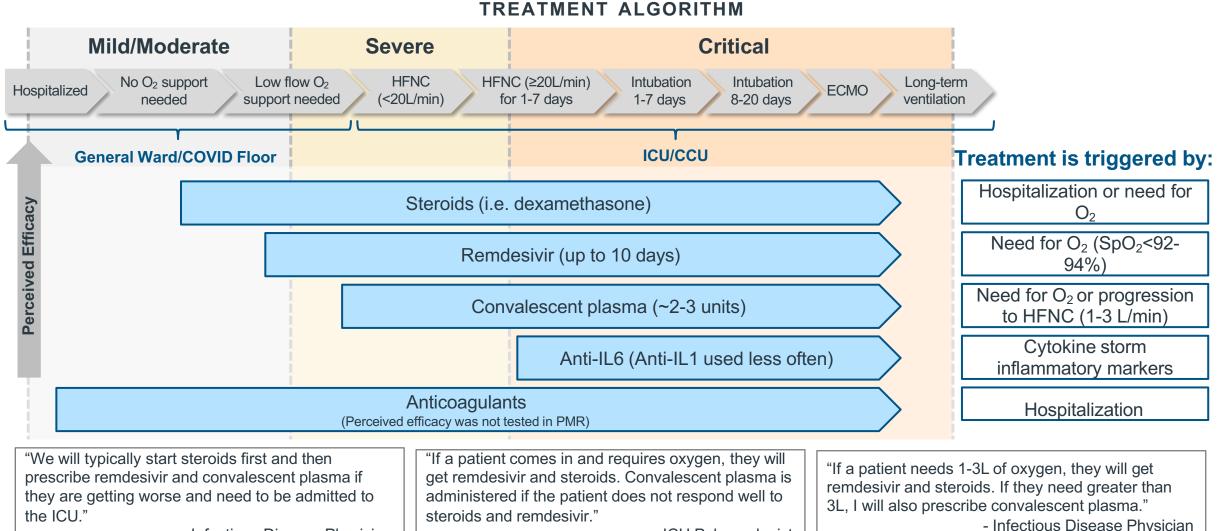
COVID-19 EUA / BLA Go-to-Market Strategy Capabilities and Experience for CovInnovations

Treatment decisions for critical COVID-19 is mainly driven by guidelines / protocols (based on recent literature) and treatment availability/supply



Stakeholders are aware that new data and novel therapeutics have been continuously emerging, and treatment protocols and available options are changing on an extremely frequent basis

Patients are initially treated with steroids & anticoagulants; if disease progresses, remdesivir, plasma, & anti-IL6s may be used



-ICU Pulmonologist

-Infectious Disease Physician

COVID-19 EUA / BLA Go-to-Market Strategy Capabilities and Experience for CovInnovations

Most therapies for COVID-19 are on hospital formulary and available for ICU specialists with limited restrictions

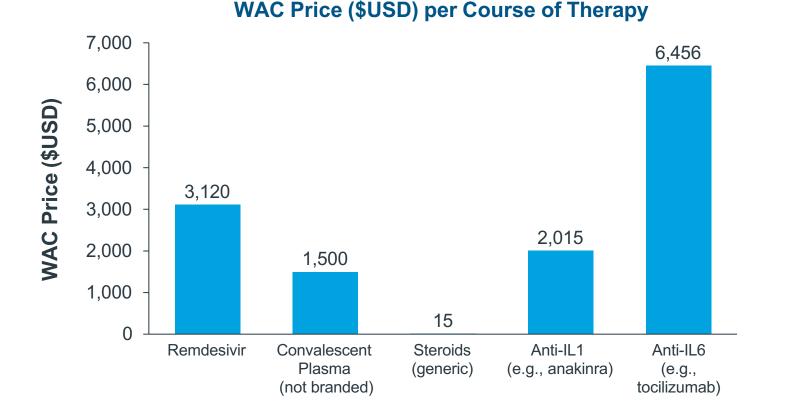
CURRENT FORMULARY INCLUSION

	Formulary Inclusion	Restrictions	 Formulary Inclusion COVID physicians have broad access to therapies, 			
Remdesivir	\checkmark	Therapies have similar	 Hospitals often have protocols to drive appropriate use, however these are not typically strictly enforced 			
Convalescent Plasma	\checkmark	 restrictions: Specialist approval (most often ID, sometimes pulmonologist and intensivist) Minimal restrictions by patient type, but use must be supported by some level of clinical evidence 	 Specialist approval (most often ID, sometimes pulmonologist and intensivist) Minimal restrictions by patient type, but use must be supported by some level of clinical 	 Protocols and restrictions are implemented via EHR, or if stricter, with sign-off of pharmacist / ICU head Use of high-cost products may be more closely monitored, such as requiring sign-off by department head 		
Steroids (e.g. dexamethasone)	\checkmark			intensivist)Minimal restrictions by	intensivist)Minimal restrictions by	 Protocols are evolving as data becomes available Hospitals have previously controlled use of therapies, however this is related to supply issues, particularly with
Anti-IL1 (e.g. anakinra/Kineret)	√ /~			 plasma and the early days with remdesivir, rather than cost Supply and federal requirements (i.e., documentation needed for EUA products) are more typical barriers 		
Anti-IL6 (e.g. tocilizumab/Actemra)	√ /~		 New products can be added relatively quickly (as fast as 1 week for review and to become available in some cases) Federal requirements and delay in receiving supply can add a few weeks or months to the timeline 			

✓ Available ~ Available, but not often in protocols

Available branded therapies for COVID-19 are priced between ~\$2,000-\$6,500 per course of therapy

THERAPY COSTS



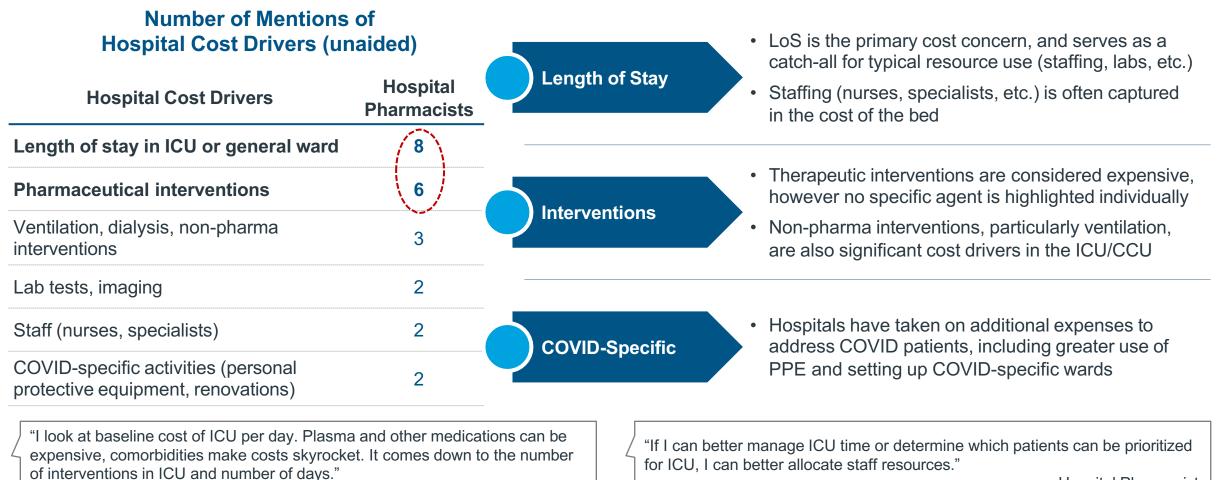
Current Therapy Costs

- Given the unmet need in critical patients and efficacy, stakeholders do not see existing therapies as price benchmarks
 - Available therapies are being used broadly, rather than only in critical patients
- Costs per course of therapy may differ by patient: patients may receive a different total dose based on physician prescribing
- Dexamethasone is considered an affordable, highly-effective option, however pharmacists and MCOs recognize it is not a price benchmark for novel branded therapies
 - Cost is not a concern when considering adding-on steroids to branded therapy

Pipeline competitors may impact the price benchmark (e.g. Regeneron contracted 70-300K courses of treatment with HHS for ~\$450M, equivalent to ~**\$1,500 - \$6,430 per course**)

In addition to cost of pharmacologic therapies, length of stay is a more critical driver of hospital costs for COVID patient management

HOSPITAL COST STRUCTURE

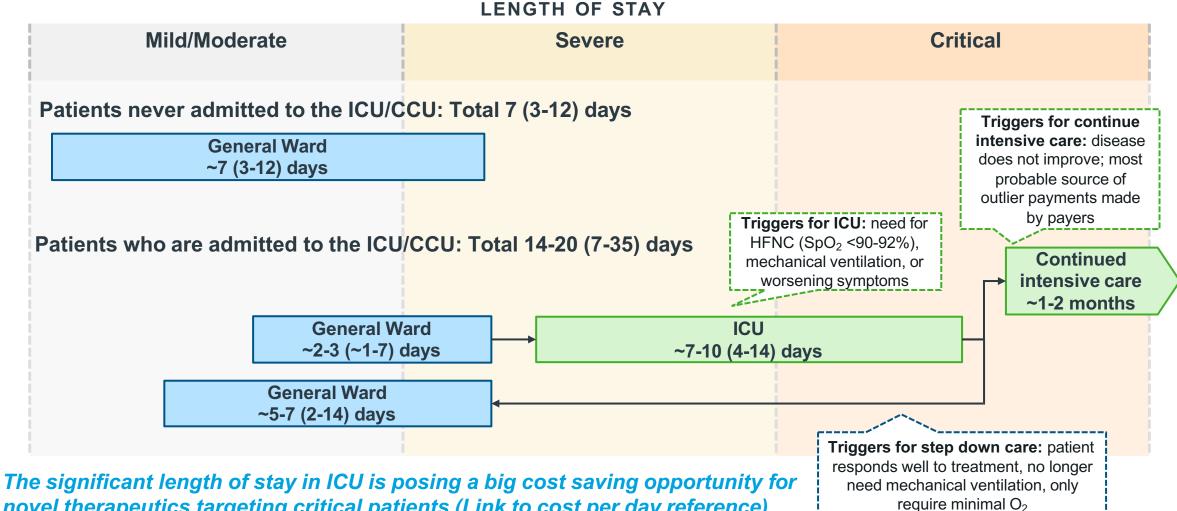


- Hospital Pharmacist

Hospital Pharmacist

COVID-19 EUA / BLA Go-to-Market Strategy Capabilities and Experience for CovInnovations

Patients admitted to the ICU spend 14-20 days total in the hospital, with 7-10 of those days in the ICU



novel therapeutics targeting critical patients (Link to cost per day reference)

Hospitals are reimbursed for COVID-19 patients primarily through Diagnosis **Related Groups (DRGs) and case rates**

Diagnosis Related Group (DRG)* Case Rate Per Diem Percent of Charges (Commonly Used) (Rarely Used) (Commonly Used) (Rarely Used) Hospital is paid flat rate • Fixed amount per day DRG determined by diagnoses/procedures regardless of charges Length of stay matters to Hospital is paid based on codes from ICD-10-CM services provided Hospitals and MCOs MCOs, but charges do not Pay per admission negotiate rate Common for national More common for For cases that are above a cost threshold, PPOs and financially Charges/length of stay specialized hospitals may receive supplemental challenged rural hospitals only used for threshold academic/tertiarv payments known as outlier payment determination hospitals For **commercial coverage**, rates are negotiated between hospitals and commercial payers (MCOs) - case rate (similar to DRG), % of charges (fee for service/FFS), or per diem Inpatient reimbursement for Medicare and Medicaid coverage is almost exclusively DRGrates for in-patient reimbursement (but may be $\sim 20\%$ higher) based

COVID-19 FUNDING & REIMBURSEMENT

Commercial insurers frequently follow CMS's lead when it comes to DRG/case

Due to the case rate reimbursement model, MCOs have minimal risk and do not manage hospital-based diseases

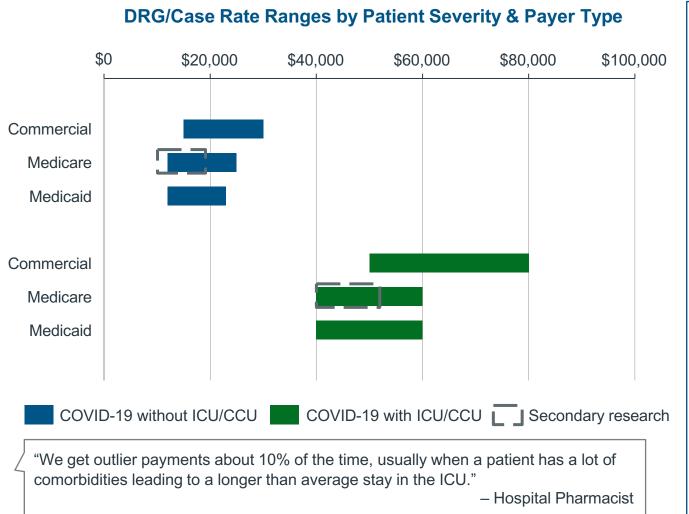
Carve-outs

Funding Mechanism

Reimbursemen

- High-cost therapeutics can be reimbursed separately ("carved out") of a DRG if it would otherwise be too expensive • to be used as part of the DRG
- Products can be carved out for CMS through the NTAP system, however this often takes 2-3 years ٠

Hospitals are reimbursed for COVID-19 patients primarily through DRGs / case rates, which range from ~\$40K-\$80K for ICU patients



COVID-19 FUNDING & REIMBURSEMENT

DRG reimbursements vary based on patient severity and comorbidities; negotiated case rates between payers and hospitals also differ significantly between payers/hospitals DRGs are not specific to COVID Remdesivir ~\$3,120 was not viewed as a significant burden, but more than normal No COVID therapies are currently carved out of DRG \rightarrow a • possibility of New Technology can be explored • Outlier payments are made in 5-15% of cases and on average are ~20-30% of the total DRG value - Outlier payments help recoup cost overages, however the hospital does not recover the full cost of service Revisions to DRGs or creation of COVID-specific DRGs are ٠ unlikely in the near future: DRGs are updated every 2-3 years based on retrospective analysis of costs Revisions to DRG would require deliberate action by CMS in response to hospital requests, which is unlikely MCOs will follow CMS's lead regarding changes (or lack) thereof) to DRGs * Respiratory infections and inflammation with major comorbidities or complications, respiratory system diagnoses with ventilator support

COVID-19 Reimbursement

Hospital administrators recognize significant potential cost offsets with efficacious treatments, assuming demonstrated LoS reduction

Treatment Setting		Cost to Hospital per Day	Reduction in Lo	oS	Cost Offsets
In ICU/CCU with ventilation		000 - \$5,700 ICU can be ~\$10K)	-5 days		~\$25,000 - \$28,500
In general ward	~\$1,700 - \$2,200		-5 days		~\$8,500 - \$11,000
Estimated cost per day is well-ali with pharmacist and MCO expecta	benefit, howeve	feasible given clinical er it needs to be ated in trial	red	s sufficient: other benefits such as luced need for ventilators will be otured by stepping down patient	

COST OFFSETS

"These cost savings seem reasonable and fall in line with expectations. It could allow for premium pricing, but higher price would eat into LoS offsets."

-Hospital Administrator

"I want to see the actual data but if this could reduce mechanical ventilation and length of stay."

-Hospital Administrator

While reimbursement for COVID-19 has evolved, additional funding mechanisms can take ~2-3 years

FUTURE REIMBURSEMENT DYNAMICS

Expansion of DRG

- DRGs may take ~2-3 years to update; policy stakeholders do not expect revisions in the near future without legislative intervention
- While MCOs negotiate rates with hospitals in an ongoing basis, commercial/Medicare Advantage follow CMS trends

Alternative Sources

Carve Out

- Pharmacists and MCOs consider current funding sufficient for COVID: alternative funding (e.g., through legislation) is unlikely to be specifically for COVID-19
- Carve out for CMS requires NTAP program, which can take ~2-3 years; commercial is unlikely to carve out without NTAP
- Some academic and tertiary hospitals have carve out clauses for high-cost treatment, but these are for specialized facilities and are not product-specific

"We are currently looking at the effectiveness of the 20% add-on payment. Hospitals initially conducted less procedures and they used it to make up for lost income. I think the DRG will go down, not up in future."

Policy Stakeholder

"Since COVID is a unique cost, I think Medicare could create a new code for it. That way they ensure hospitals get reimbursed and they don't overpay for other respiratory related diseases."

- MCO Payer

Available funds to pay for are unlikely to change in the next few years

Proposed Go-To-Market Partnership Model

	EUA Fees	Launch Start Up	Year 1 Ops	Grand Total
Business Planning & Mgt.	\$139,000	\$496,034	\$58,333	\$693,367
Medical Affairs & Safety	\$736,825	\$680,000		\$1,416,825
Pricing & Market Access		\$375,000	\$42,000	\$417,000
Contract Field Teams*	\$2,747,435	\$239,728	\$8,387,654	\$11,374,817
Brand Strategy & Development	\$100,000	\$868,000	\$75,000	\$1,043,000
Comm Ops & Analytics	\$188,000	\$330,638	\$537,258	\$1,055,896
IT & Support	\$412,400	\$479,345	\$298,338	\$1,190,083
Quality Assurance		\$140,000	\$72,000	\$212,000
Compliance	\$135,000	\$278,000	\$62,173	\$475, 173
Total	\$4,458,660	\$3,886,745	\$9,532,756	\$17,878,161
RWE	N/A	TBD, Scoping Rqd.	TBD, Scoping Rqd.	
MICC, PQC, AE/SRL (Med Affairs)	Included Above	Excluded Above	Excluded Above	

*Contract Field Teams Deployment/Salary: 10 MSLs: \$180K 4 CNEs: \$100K 10 Sales Reps: \$140K 2 Key Account Managers: \$150K Access Advisor: \$10K per month

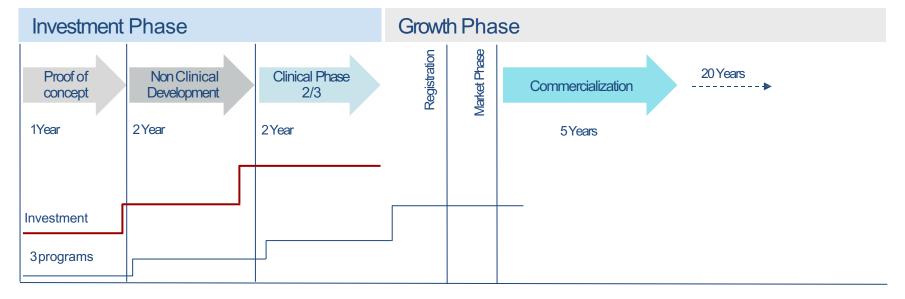
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Financing Strategy

Financing plan and investment case (incl. key assumptions and valuation sensitivities)

Business Model and Value Creation

First Exit opportunity in 2024



Growth Enabling Revenues:	Upfront Payments	Milestone Payments	Royalties
Indication specific licencing of assets			

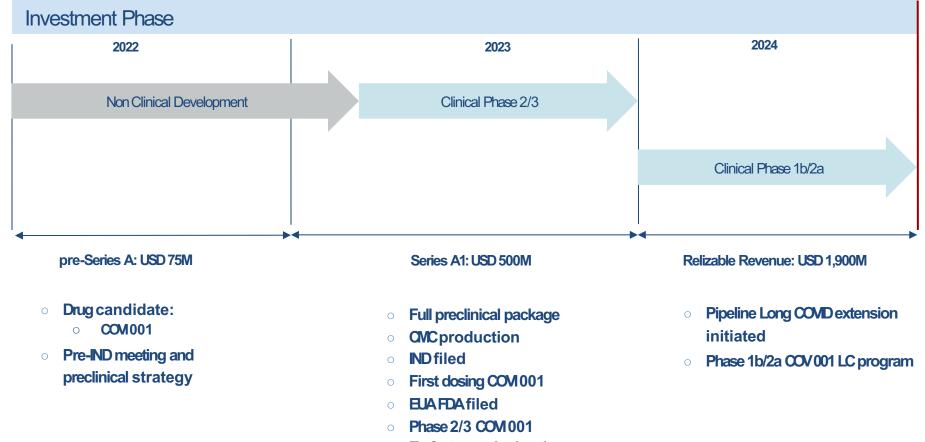
Potential Revenue Streams:

Research collaborations on core technologies

Project based payments	
	\rightarrow

Key Value Inflection Points





• Ex-factory sales begin

Strategic Roadmap

	Short Term 2022 - 2025	Long Term 2025 - Market
Strategy	 Develop 3 indications to proof of efficacy in human for licensing by end of 2024 Maximise the monetization of platform development through collaborations and partnerships Generate recurring income through global sales, licensing and co- developments 	 Develop compelling indications in the COVID-19 space Establish key partnerships based on established differentiation Generate growth capital through licensing of clinical, device and diagnostic assets
Focus	 Human proof of concept Pipeline development and platform validation Monetization of know how and technology Company established for growth 	 Pipeline development Growth of the company to multiply the platform value Business development and strategic collaborations
Investment Areas	 Clinical programs Commercial operations Platform technologies enabling clinical and pipeline development 	Clinical programs and strategic exploratory programs
Value Generated	 3 drug candidates validated in human (Phase 2) Drug candidates synergize with further indications Revenues and validation through commercialization and technology licensing deals 	Increasing pipeline

Overview key assumption

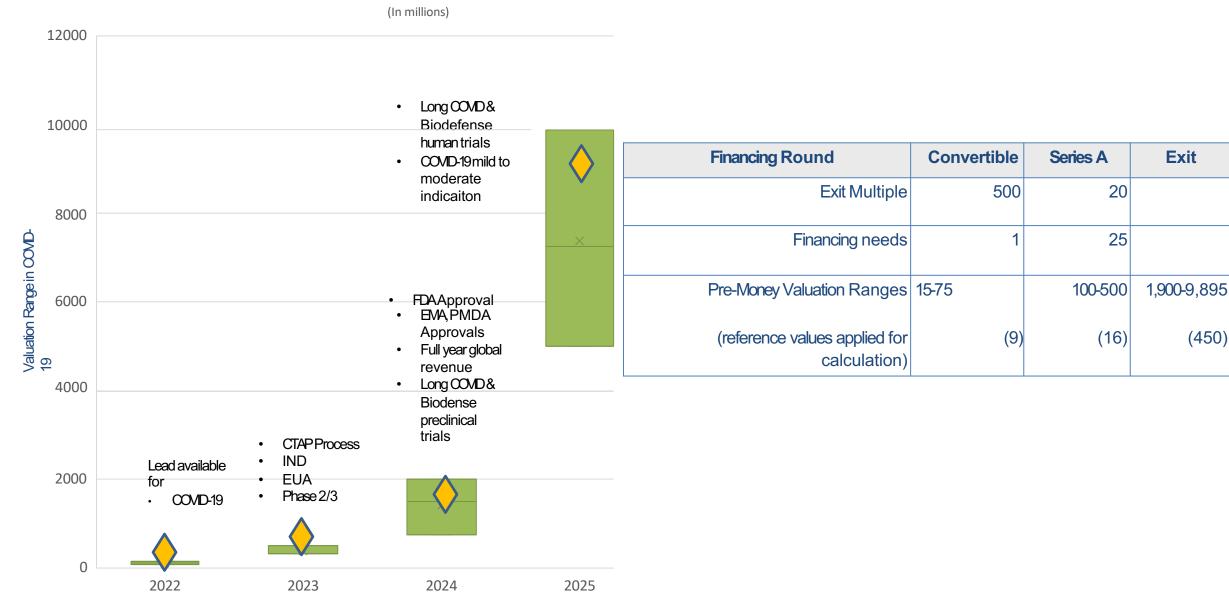
	Asset COVI 0001	Asset COVI 002	Asset COVI 003	Comments/Source
Valuation Approach	EV/Revenue Multiple	EV/Revenue Multiple	EV/Revenue Multiple	Enterprise Value to Revenue Multiple is a valuation metric used to value a business by dividing its enterprise value (equity plus debt minus cash) by its annual revenue
Discount Rate	15%	15%	15%	15%
Revenue Starting Year	2023	2026	2026	
Probability to market	62%	59%	62%	Sources: FDA, Booz Allan
# of years included in forecast	5	0	0	
Geographical region used	US, EU, Japan, WOW	US, 2027 for EU, Japan, WOW	US	
Patient segmentation	1st line treatment	1st line treatment	1st line treatment	
Market share	10% in 2023 to peak to 60% in 2027			
Sales Price	Ex-factory price:	Ex-factory price:	Ex-factory price:	
	USA: USD 633	TBD	TBD	
	WOW: USD 500			

Value inflection and Return on Investment



- Valuations for Series Ais based on benchmark cases and expected multiple assumptions
- Valuation at exit is based on 5.6x median EV/Revenue Multiple calculation
- IRR's and multiples represent an attractive investment opportunity
- Value is mainly driven by COVI-001 asset
- Early licencing deals are not reflected in the current business case

Value inflection and Return on Investment



Ownership over time

Expected ownership at pre-Series A and post-series A based on the following assumptions:

• Series A raise only capital raise until relizable revenue in 2H of 2023

	Pre-Series A	Series A
Existing shareholders	100%	60%
Convertible loan investors	0%	5%
Series A	0%	25%
ESOP (incl. exercised)	0%	10%
Total	100.00%	100.00%

CovInnovations Leadership Team



Aury Nagy, M.D.

Board Chairman, Founder and Chief Science Officer

- Neurosurgeon and board member of the Nevada State Medical Board and former Chief of Neurosurgery at University Medical Center
- Graduated from Yale University, Baylor College of Medicine, and George Washington School of Medicine Department of
 Neurological Surgery
- Studied at Stanford, Harvard, Duke, Cal Western, LSU, University of Arizona, and the University of Pennsylvania

Jamie Jones

CEO

- Senior Executive with 30 years experience at Pfizer, Gilead Sciences, Alnylam Pharmaceuticals, and Double Rainbow Biosciences
- Track record of ten drug launches, engineered commercial startups on three continents, and an award-winning industry executive for innovation, groundbreaking marketing, and sales strategies
- MBA in Marketing and a BS in Finance from the University of Central Florida

Kristine Leavitt, MSN, FNP-BC

Chief Operating Officer

- Board-certified nurse practitioner in family medicine, hospice, and palliative care
- Former program coordinator for the University Medical Center of Southern Nevada's transplant program Master of Science in Nursing (MSN) from the University of Nevada – Las Vegas

Amy Stone, Ph.D.

Lead Scientist

- Assistant Professor of Microbiology and Immunology, Touro University Nevada, Henderson, NV
- Graduated with Ph.D. in Immunology from University of Colorado Denver, CO and B.A. in Biochemistry and Molecular, Cellular, and Developmental Biology from University of Colorado Boulder

Richard Renfrow

CMC-Lead

- Assistant Professor of Microbiology and Immunology, Touro University Nevada, Henderson, NV
- Graduated with Ph.D. in Immunology from University of Colorado Denver, CO and B.A. in Biochemistry and Molecular, Cellular, and
 Developmental Biology from University of Colorado Boulder

CovInnovations Board of Directors

Aury Nagy, M.D.

Board Chairman, Founder and Chief Science Officer

Elliot Goldstein, MD

Founder and former CEO of ProMIS Neurosciences, Inc. (Toronto: PMN.TO, CAD 5 million market cap). He has 30 years of experience bringing products through the FDA to market, including cyclosporine

Eric Henderson

Founder and CEO of CBG consulting. He has contributed to US companies securing approximately \$25 billion in projects financed annually and assisting US and global companies with participating in World Bank-financed procurement opportunities in more than 150 countries. Mr. Henderson earned an MA in international relations and affairs from American University.

Ibrahim Pataudi

Vice President of Business Development for NuID, a blockchain company with National Security Agency (NSA) contracts. Mr. Pataudi earned a bachelor's degree in International Relations and Affairs from Claremont McKenna College.

Mansoor Ijaz

Founder and CEO of Crescent Global Partners. Mr. Ijaz is a proprietary trader and hedge-fund manager who founded Crescent Investment Management in New York in 1990. Crescent, and its successor companies, operate CARAT, a proprietary trading system developed by Ijaz in the late 1980s during his graduate research studies at MIT.



Contact Us

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