



VAXIL
VAXIL BIO THERAPEUTICS

OCTOBER 2020

**DIRECTING
IMMUNITY
DIRECTLY**

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Executive Summary

Vaxil is an Israeli biotechnology company developing novel immunotherapies including neoantigen-like signal peptides and antibodies to treat cancer and infectious diseases. Our experienced board and scientific team have access to world class medical centers and academia.

Leading platforms:

COVID- 19 Vaccine Candidate (Corvax™)

- Successful COVID-19 pre-clinical work continues after potential Vaccine discovered and patents filed (3/20)
- Successful pre-clinical completed (9/20)
 - Immune response in vitro
- Path to clinical trial within 9-12 months

Oncology

- Phase 1/2a: multiple myeloma (US/EU orphan)
- High immunogenicity
- Robust immune response
- Specific to tumor cells
- No immune tolerance
- Bypasses tumor escape
- In-licensed P-EsBP polymer

Vaxil Team



David Goren, MBA CEO and Chairman

- Head, Pfizer Worldwide Strategy and BD
- CEO, Pfizer Israel
- President, AstraZeneca Israel
- VP Digital Health Strategy



Gadi Levin, CPA, MBA CFO and Director

- CFO, LabStyle Innovations Ltd
- CFO, BriaCell Therapeutics Corp.
- VP / CFO of 2 investment houses



Yuval Avnir, PhD, MA Head of R&D

- Research Fellow, Dana-Farber Cancer Institute
- Head, Scientific Equipment Center, Bar Ilan U Faculty of Medicine (Galilee)



Dana-Farber
Cancer Institute



Riva Kovjazin, MD, PhD Senior Scientist

- Project leader, Center for Clinical Immunology
- Senior scientist at XTL Biopharmaceutical

Ari S. Kellen, MD, MBA, Director

- Strategic advisor and investor to early-stage innovative healthcare companies
- Director, Theraclion
- Senior partner, McKinsey & Co.
- EVP and GM, Bausch & Lomb US
- Advisory board, BioNanoSim (BNS)

Shawn Langer, MD, Director

- Senior partner, McKinsey & Co.
- Portfolio manager, BHMS Investments

Ronny Pinkus, PhD, CMC

- VP Mfg & Operations, Moebius Medical
- Head Analytical, InSight Biopharmaceuticals

Rivka Zaibel, Regulatory & QA

- President, ADRES, VP RA/QA, BTG

Beth Leffel, PhD, MPH, PreClinical

- PharmAthene, NBACC, USAMRIID

Financial Summary

(updated September 30, 2020)

Market Cap

- CAD\$13.6 M

Share price

- CAD\$0.13



Cap Table

- Shares: 104.5 M
 - Warrants: 21.4 M (exercise price CAD\$0.10)
 - Options: 6.6 M (avg exercise price CAD\$0.10)



Trading History

- 2016: RTO into Canadian company (TSX.V)
- 2012: RTO by Vaxil on TASE
- 2011: Listed on TASE



Funds Raised (last 4 yrs) primarily from individuals

- 2020: CAD\$1.4 million (warrant exercises)
- 2020: CAD\$0.13 million (debentures)
- 2018: CAD\$1.8 million (private placement)
- 2016: CAD\$2.6 million (private placement)



Pipeline



ID

Tuberculosis

COVID-19 Vaccine

Other candidates

Oncology

MUC1-SP

Additional SPs

P-ESBP

mAb program

In Brief

01

Targeting COVID-19
(infectious disease) and
solid tumors

02

VaxHit™ target ID for
signal peptides and mAbs

03

Clinical stage immunotherapy

04

5 Patent families

05

Successful Phase 1/2a multiple
myeloma clinical trial

06

Orphan designation for
multiple myeloma (FDA / EMA)

07

P-ESBP In-licensing

08

VXL traded on TSXV

Signal Peptide Characteristics

Characteristics



**Conserved
short sequences**



**Dense number
of epitopes**

- Wide HLA-subclasses
- Cellular / humoral response



**Specific to tumor
cells**



**HLA-independent
presentation**

Vaxil Advantage

**No antigen loss
Broad population**



**Very high
immunogenicity**

- Universal – wide population
- Robust immune response



Neoantigen

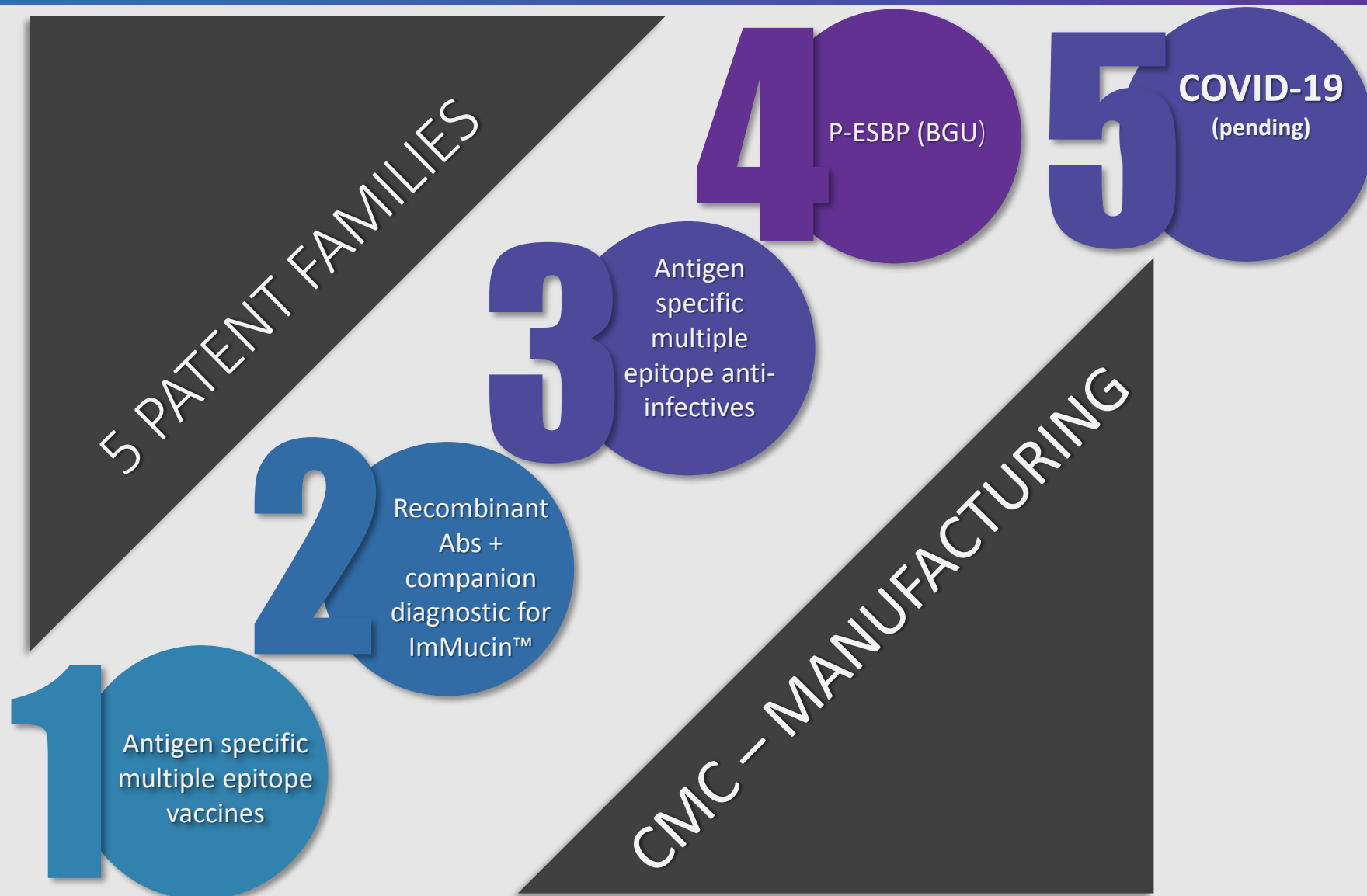
- No immune tolerance
- Safety profile



**Bypasses
tumor escape**



Universal Neoantigen



Applying Signal Peptide Platform to COVID-19

- Proprietary VaxHit™ tool identified signal peptides in SARS-CoV-2 (COVID-19) genome with high immunogenic potential
- Unique approach: Complete immune response
 - Cellular (T cell) AND humoral (antibodies)
 - Targeting infected cells, not virus
- Not susceptible to viral escape due to mutations
- Not prone to resurge due to protein mutations
- Potential for immunity preventing other outbreaks and therapeutic effects
- Efficiently manufactured and scaled (scale-up cost, simplicity)

Achievement to Date

- Candidate/s discovered using proprietary VaxHit™ tool
- IP secured
- Preclinical efficacy experiments successfully demonstrated immunogenicity
 - ex vivo and in vivo immune response and functional assays
- Successfully established that CorVax™
 - demonstrates a favorable immune response in healthy blood cells
 - demonstrates a favorable immune response in animals

Next Steps

- Pre-clinical "challenge" in vivo experiment*
- Initiate GMP production, formulation including stability and solubility testing
- Secure IND regulatory pathway with gap analysis
- Tox, pre-IND and IND submission
- Phase I/IIa study

* In a "challenge" experiment animals are injected with the experimental vaccine followed by exposure to COVID-19. The experiment attempts to demonstrate that the animals who were vaccinated have a better result (i.e., are less sick).

CorVax™ Pre Clinical Experiments

In Vitro Immunogenicity:

Aim: Establish *CorVax*™ immunogenicity by assessing T cell proliferation.

Method: Healthy donor peripheral blood mononuclear cells (PBMCs) stimulated by peptide-loaded autologous dendritic cells.

Results: T cell proliferation was observed by flow cytometry for all candidate signal peptides, to varying degrees, compared to unstimulated cells.

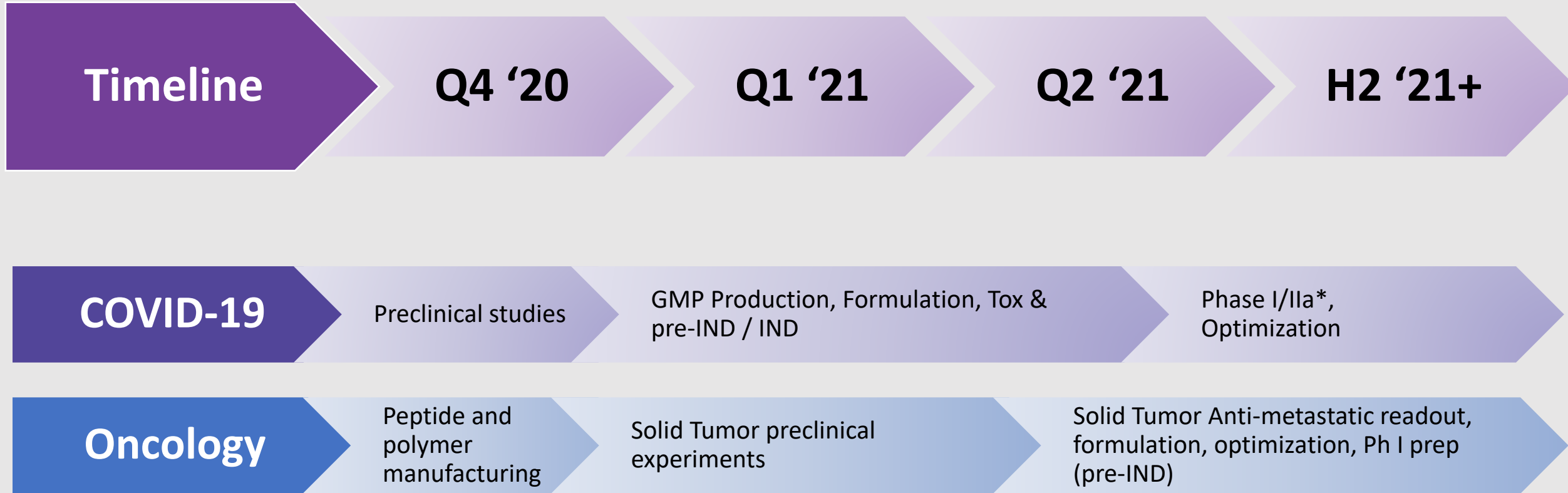
In Vivo Immunogenicity:

Aim: Establish *CorVax*™ immunogenicity in an animal model by assessing antibody and T cell response.

Method: Two mice strains vaccinated with 3 weekly *CorVax*™ injections.

Results: Immune response was observed when analyzing results from ELISA, ELISpot, and restimulation assays.

VAXIL Planned Milestones



* subject to regulatory considerations due to COVID situation

Thank You

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