



Bio₂XyTran[®]

Corporate Overview

Forward Looking Statement

This Descriptive Presentation (the “Presentation”) has been prepared by Bioxytran, Inc (the “Company”) and recipients are not entitled to rely on the accuracy or completeness of the Presentation. Statistical information contained in this Presentation is based on information available to the Company that the Company believes is accurate. It is generally based on publications that are not produced for the purposes of securities offerings or economic analysis. The Company has not reviewed or included data from all sources and cannot assure prospective parties of the accuracy or completeness of the data included in this Presentation. Forecasts and other forward-looking information obtained from these sources are subject to the same qualifications and the additional uncertainties accompanying any estimates of future market size, revenue and market acceptance of products and services. The Company undertakes no obligation to update forward looking information to reflect actual results or changes in assumptions or other factors that could affect those statements.

This Presentation has not been filed or reviewed by the Securities and Exchange Commission (“SEC”) or any securities regulatory authority of any state, nor has the SEC or any such authority passed upon the accuracy or adequacy of this Presentation. This Presentation does not constitute an offer to sell or solicitation of an offer to buy any securities. This Presentation does not purport to contain all information which may be material to a prospective party, and recipients of this Presentation should conduct their own independent evaluation and due diligence of the Company. Each recipient agrees, and the receipt of this Presentation serves as an acknowledgment thereof, that if such recipient determines to engage in a transaction with the Company, its determination will be based solely on the terms of the definitive agreement relating to such transaction and on the recipient’s own investigation, analysis and assessment of the Company and the transaction. The Company does not intend to update or otherwise revise this Presentation following its distribution.

Living with Viruses

Should we embrace



COVID-19 Infections
95.7 million



Respiratory Syncytial Virus (RSV)
2.1 million (under 5yrs)



COVID-19 Deaths
1.07 million



Viral Conjunctivitis
5.0 million



HIV Positive
1.2 million



Long Covid
Over 20% of worlds cases (1 in 5)

Statistics are unacceptable

Something needs to be done

INVESTING IN A COMPANY

THAT

Rejects the idea that Living with the virus is okay

Mission Statement

Bioxytran is a clinical stage pharmaceutical company developing platform technologies in the fields of Glycoviropology, Hypoxia and Degenerative Diseases to eliminate viruses and prolong lifespan using carbohydrate drug design.



GLYCOVIOLOGY



HYPOXIA



DEGENERATIVE DISEASES

COVID is OVER – Why Invest?



**No One
Masking**



**No One
Testing**



**No One
Distancing**



**President Says
“its over”**

Propaganda or Facts

- CDC no longer reports daily infections (only weekly)
- Averaging 400 deaths daily (20 of 50 states reporting)
- New immune evasive variants like XBB BQ.1.1
- 4 million sidelined by Long COVID (Brookings Institute)
- 1 in 5 people present the signs of Long COVID

**So is COVID really over?
What about Long COVID?**

COVID Has

- Easy indication to prove efficacy (viral elimination 3 days)
- Label expansion likely (platform technology)
- Non-toxic profile = favorable regulatory treatment outside USA

**No Relevance to Overall
Strategy**

GOAL: DRUG APPROVAL ASAP!

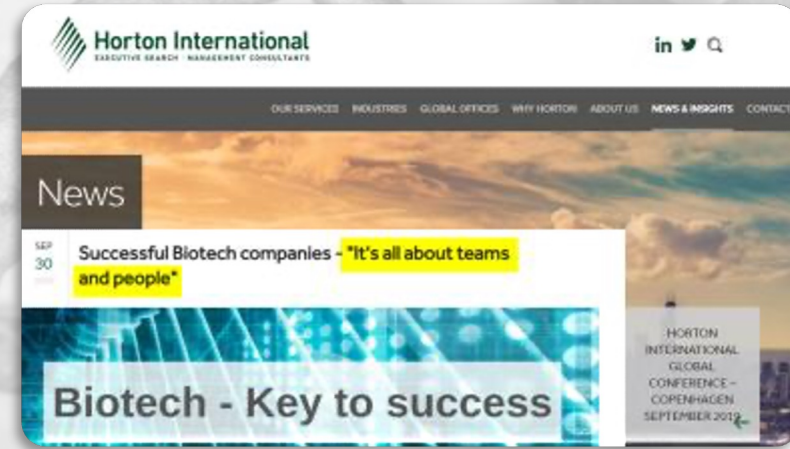
What to Look for in a Biotech

Attributes of a Successful Biotech

(The M&M's of Biotech)

Management – Regulatory Experience

Magic – Science



GRIT DAILY®
The Premier Startup News Hub.

Press Releases ↗ Enterprise ▾

Defining a Successful Biotech Entrepreneur

By Paul Brennan

So, what is more important to the livelihood of biotech entrepreneurs? Is it the scientific breakthrough? Or perhaps it is the management and finance skills needed to bring the company's innovation to commercialization? The answer is, all are equally important – good management fails because the technology could not support it, and good technology fails because of poor management decisions. **Management and technology, as well as money**, are the key components necessary for a successful biotech venture.



Life Sci VC

Recovering scientist turned early stage VC - A biotech optimist fighting gravity



Leadership At All Levels In Biotech

Posted January 23rd, 2019

By Deanna Petersen, in Corporate Culture, From The Trenches, Leadership, Talent



This blog was written by Deanna Petersen, CBO of ATROBIO, as part of the From The Trenches feature of LifeSciVC.

Biotech is an industry that calls for leaders. It requires people who are willing to take risks, conquer new science, and have endurance for the many years it takes to develop a new medicine. In addition, professionals who thrive in biotech have the know-how, confidence and guts to tackle business goals.

Key Leadership in Galectin Science

David Platt PhD, CEO, CSO, Chairman
Carbohydrate chemistry expert, founded
four publicly traded companies, raised
\$150m in public markets, created \$1B in
shareholder value, and led development
of two drugs.

4,000+

Journal Articles on
Target Receptors

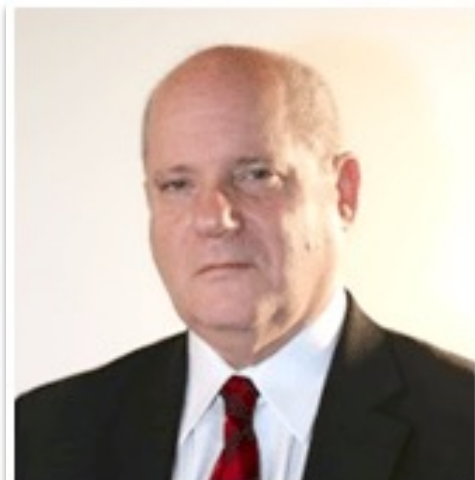
Galectin Science



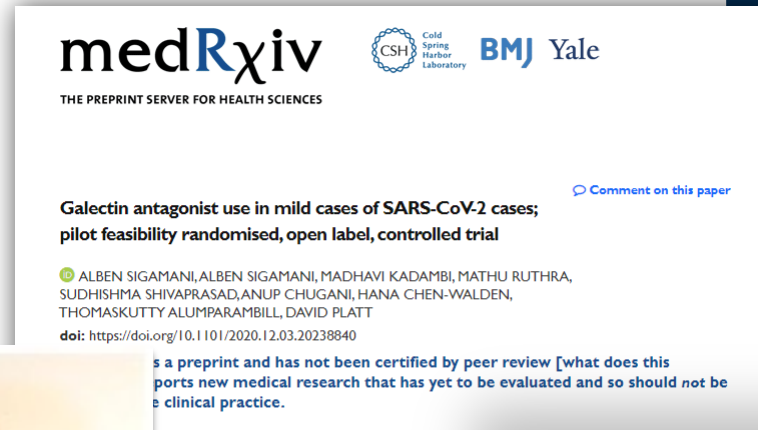
Proven Safety Profile
in Drug Class



Efficiency in many
etiologies



- 2 Textbooks
- 5 Public Companies
- 10 Journal Articles
- 30 Patents
- 30 Clinical Trials
- 200 Animal Experiments



is a preprint and has not been certified by peer review [what does this
means?]. It reports new medical research that has yet to be evaluated and so should not be
used to guide clinical practice.



Journal of Vaccines & Vaccination

Galectin Antagonist use in Mild Cases of SARS-CoV-2: Pilot Feasibility Randomised, Open Label, Controlled Trial

Alben Sigamani¹, Madhu Raghav², Sudhishma³, Samudh Shetty⁴, Madhu⁵, Anup Chugani⁶, Hana Chen-Walden⁷, David
Platt⁸, Thomas Kott⁹

¹Department of Clinical Research, Nanyang Health, Bangalore, India; ²Mayo Clinic Medical Center, Nanyang Health, Bangalore, India;
³Department of Clinical Research, Nanyang Health, Bangalore, India; ⁴Pharmaceuticals, Inc., Boston, USA; ⁵Bydure Score-B White

30+ years

of research in Galectins,
carbohydrate-binding
proteins

The Team

Management

David Platt PhD, CEO, CSO, Chairman

Carbohydrate chemistry expert, founded four publicly traded companies, raised \$150m in public markets, created \$1B in shareholder value, and led development of two drugs.

Ola Soderquist CPA, MSA, MBA, CFO

>30 years multi-industry financial experience.

Mike Sheikh, CCO

>10 years of business development in life sciences. Broker and Research Analyst.

Hana Chen-Walden MD, CMO, Board Member

>30 years experience in pharmaceutical regulatory affairs in US and Europe.

Board of Directors

Anders Utter MBA, Director

Audit Committee Chair, >25 years of managerial finance and accounting in medical devices and manufacturing.

Dale Conaway DVM, Director

Veterinary Medical Officer, Federal Research.

Alan Hoberman PhD, Director

Executive Director of Site Operations and Toxicology at Charles River Laboratories.

Advisory Board

Avraham Mayevsky PhD, Professor Emeritus

Worldwide authority in the field of minimal invasive monitoring of tissue and organ physiology; and professor at the Faculty of Life Sciences, Bar-Ilan University, Israel.

Kevin H Mayo, Ph.D.

Professor of Biochemistry, Molecular Biology & Biophysics at the University of Minnesota (UMN). Known authority in the field of structural biology and structure-based drug design

Alben Sigamani, MD

Professor and Head of Clinical Research Narayan Health, Bangalore. >17 years of experience in clinical research

Technology Overview

ProLectin Rx – Glycoviropology

Virology:

- Covid-19
- Influenza
- Conjunctivitis
- RSV
- Other virologic diseases

Long term symptoms resulting from viral infections (long-hauler):

- ARDS
- Pulmonary Fibrosis

ProLectin-M is a licensed technology that targets COVID-19 mild to moderate cases

BXT-25 – Hypoxia & Degenerative Diseases

Ischemia:

- Stroke
- Alzheimer
- Dementia
- Traumatic Brain Injury

Anemia

Wound healing

Oncology and Fibrosis

- Cancer Metastasis
- NASH
- Other Fibrotic condition

**Platform
Overlap**

Galectin-3: One Molecule for an Alphabet of Diseases, from A to Z

Conditions and diseases in which a role for Gal-3 has been postulated

Aa	Bb	Cc	Dd	Ee	Ff	Gg	Hh	Ii
Asthma	Blood test	Cancer	Degenerative Aortic Stenosis	Endometriosis	Fibrosis	Gastritis	Heart	Inflammation
Atherosclerosis		Cerebral infarction	Diabetes Mellitus	Enteric nervous system			HIV Infection	Interstitial lung disease
Atopic Dermatitis		COPD		Encephalitis				
aaa	bbb	ccc	ddd	eee	fff	ggg	hhh	iii
Jj	Kk	Ll	Mm	Nn	Oo	Pp	Qq	
Juvenile Idiopathic Arthritis	Kidney	Liver Fibrosis	Mortality	NASH	Obesity	Pneumonia	Q Fever	
						Pulmonary hypertension		
jjj	kkk	lll	mmm	nnn	ooo	ppp	qqq	
Rr	Ss	Tt	Uu	Vv	Ww	Xx	Yy	Zz
Rheumatoid Arthritis	Sepsis	Target therapy	Urinary tract infections	Venous Thrombosis	Wound Healing	X syndrome of the heart	Yeast infection – Candidiasis	Zoster-related pain
	Systemic Sclerosis							
rrr	sss	ttt	uuu	vvv	www	xxx	yyy	zzz

ProLectin Rx Glycovirolology

ProLectin-Rx Galectin Antagonist Platform



Versatile

Mutation agnostic
therapeutic



Tested (Phase 2*)

No toxicity
Reduction of viral load to
undetectable levels 7
days



No Expected Limitations



Efficient

Eliminate contagion

**First line of defense
against all mutations
of Coronaviruses**

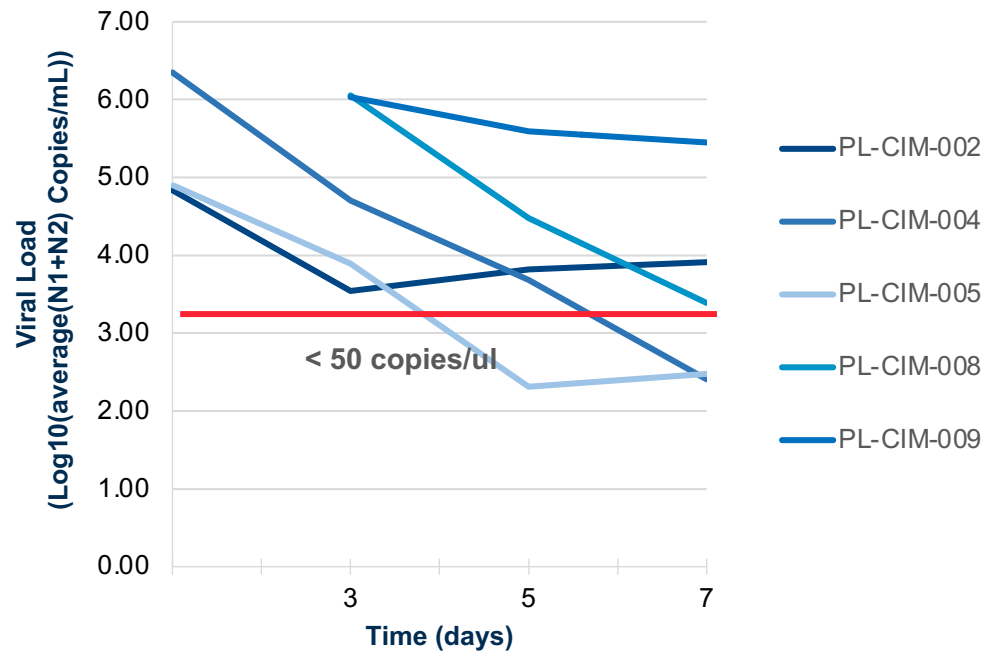
STATUS:

Clinical Trial Stage – **Phase II**

*Galectin approach to lower covid transmission - Drug Development for clinical use (medRxiv.org)

Patients Treated With Galectin Antagonist Experienced Reductions In Viral Load

Viral Load Vs. Time in Treated Patients



Day	N1+N2 Copies/mL			
	1	3	5	7
PL-CIM-002	137080	6970	13180	16340
PL-CIM-004	4468590	101170	9660	510
PL-CIM-005	159730	15630	410	600
PL-CIM-008	N/A	2268630	60180	4890
PL-CIM-009	N/A	2154530	783750	563430

Patient 9 appeared to be an anomaly. Additionally, Patient 8 had no PCR measurement of viral load on day 1, so day 1's PCR measurement was assumed to equal day 3's measurement.

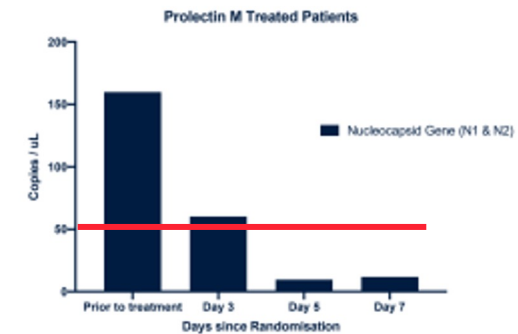
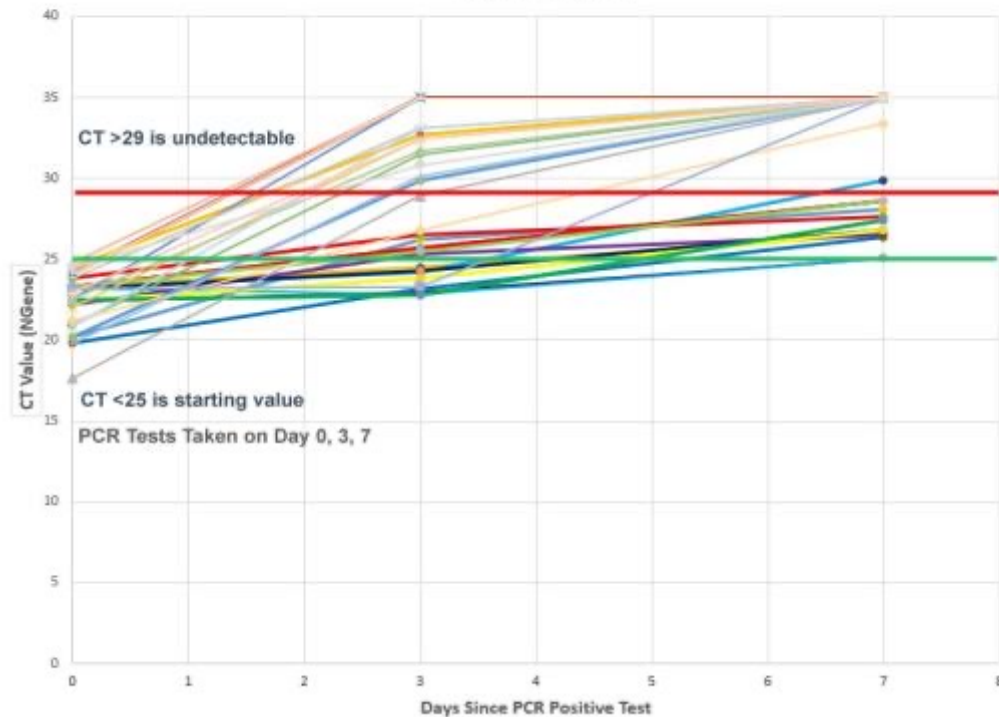


Figure 3 – drop in absolute copy numbers of nucleocapsid gene over time – Treated group

PCR Test Data (Blinded)

Prolectin-M Clinical Trial Results

Blinded Data n=34



Cycle Threshold (Ct) values are used to assess infectivity of the patient using a nasal pharyngeal test. Lower values are a proxy for higher viral loads and increased infectivity. Values over 29 are considered PCR negative. Starting Ct values of patients were under 25.

Day 3 – 15 out of 34 were PCR negative (44.1%)
Day 7 – 18 out of 34 were PCR negative (52.9%)
n = 34

No toxicity signals

Randomized 1:1

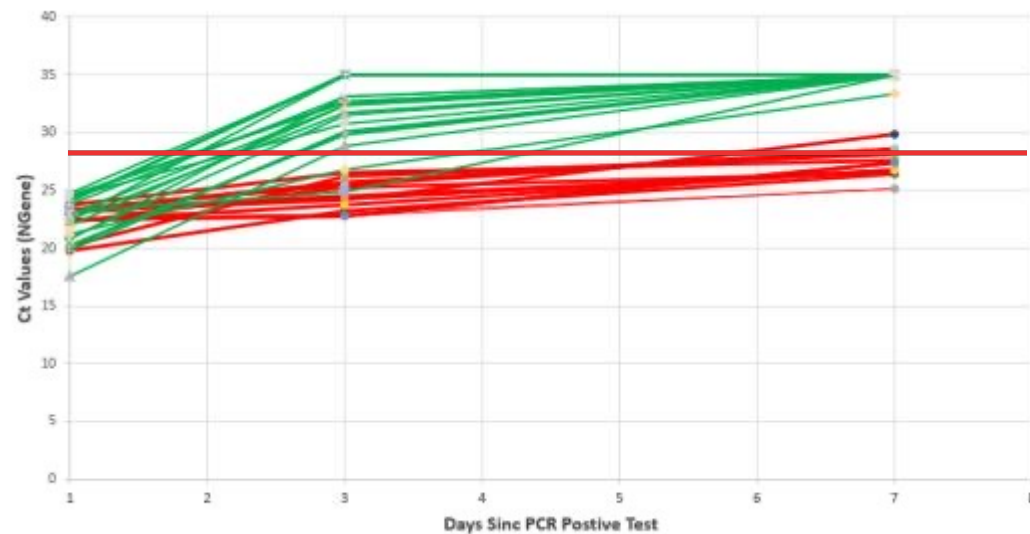
Double Blind Placebo Controlled Trial

Very Encouraging Data Grouping and indications of efficacy with no safety signals

PCR Test Phase 2 Data

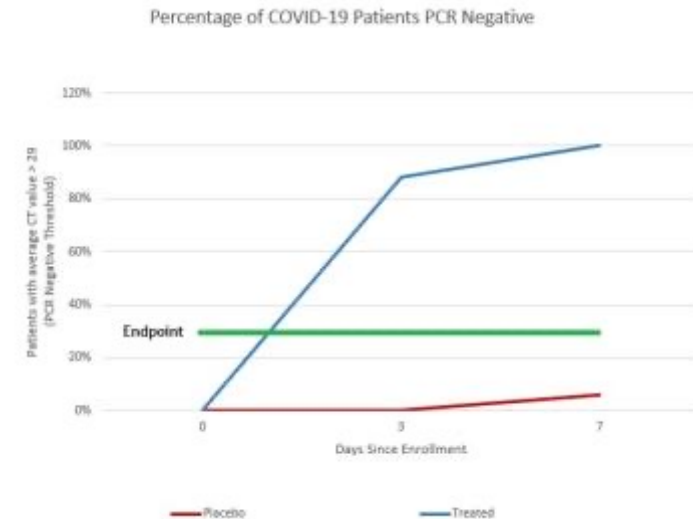
ProLectin-M Ct Values

Unblinded Data n=34



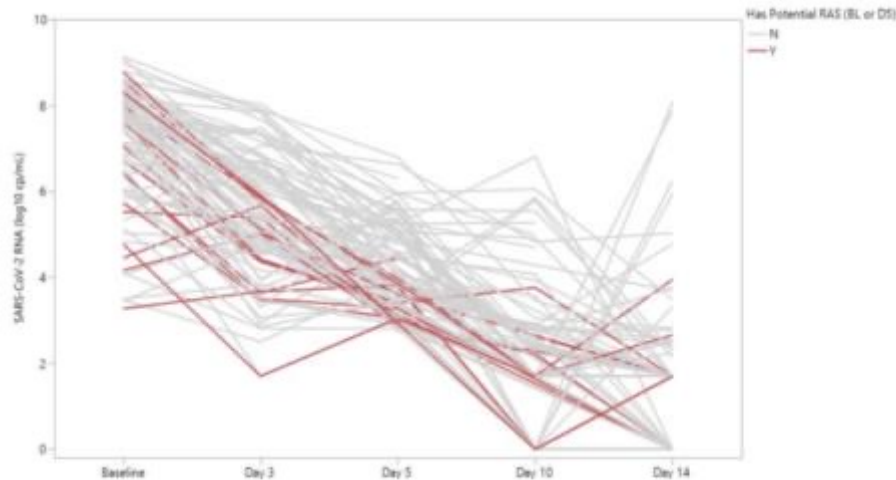
Day 3 – 14 out of 17 were PCR negative (**88%**)
 Day 7 – 17 out of 17 were PCR negative (**100%**)
 n = 34
 No toxicity signals
 Randomized 1:1
 Double Blind Placebo Controlled Trial

Actual results show tight grouping btw treated arm and placebo arm



Elimination of Viral Rebound

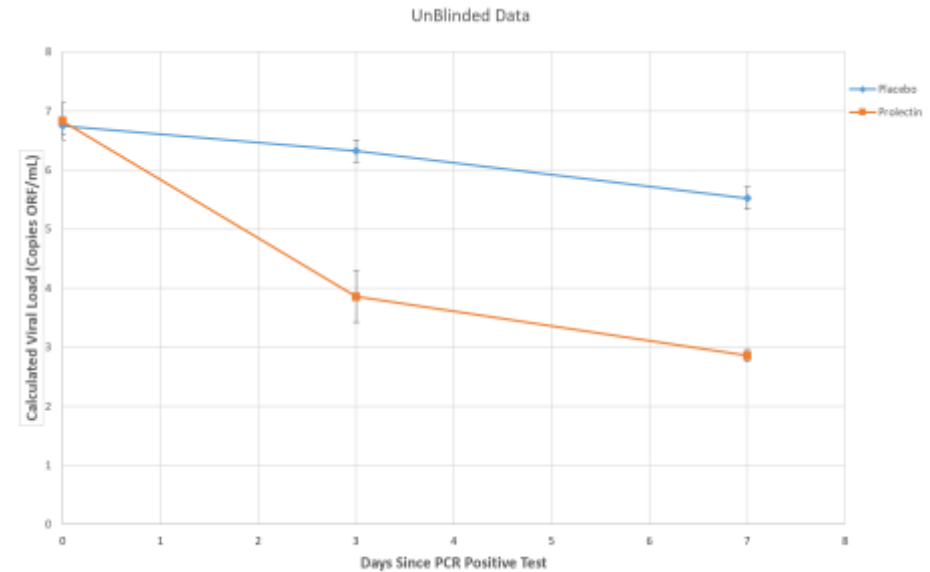
Paxlovid rebound was common after day 10



Source: FDA analysis.

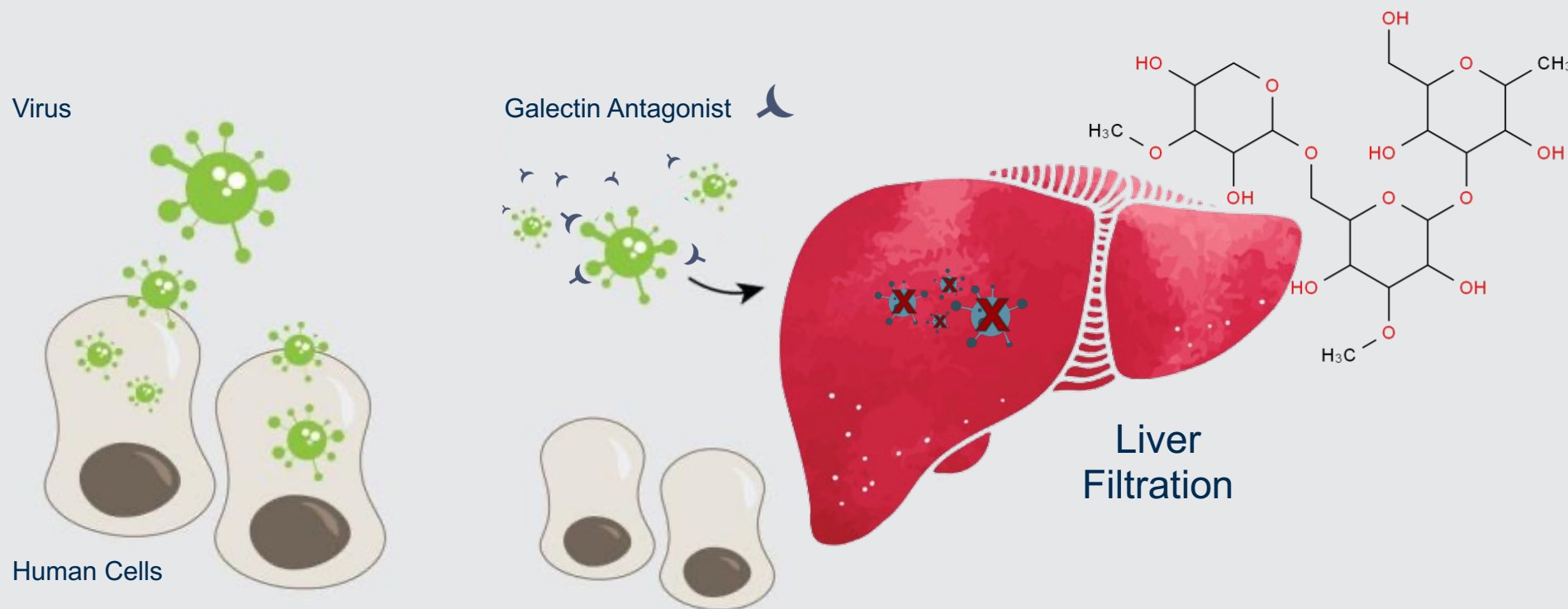
Figure 2. SARS-CoV-2 RNA levels in NP swabs among Paxlovid treated subjects with or without SARS-CoV-2 amino acid substitutions detected in Mpro or cleavage site positions potentially associated with resistance.

There were no rebounds within 14 days.



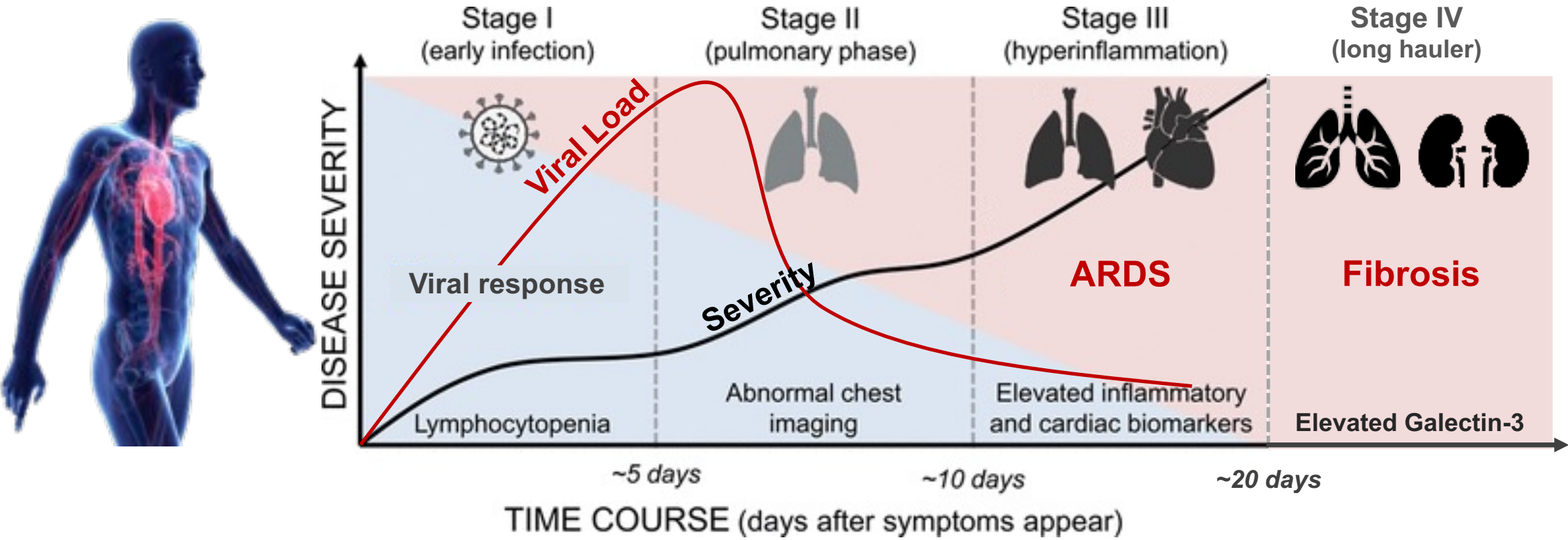
Galectin Antagonists Tags Virus For Elimination

Theoretical Mechanism of Action



End-to-End Solution

Treatment	ProLectin-M Oral	ProLectin-I Intravenous	ProLectin-A Intravenous	ProLectin-F Intravenous
Combination				



Glycovirology Development Pipeline

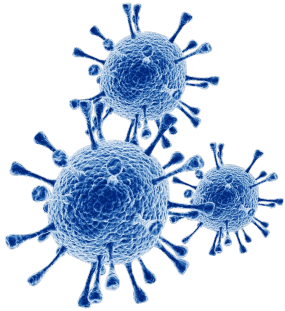
Drug & Process Development Cleared

→ Completed → Planned

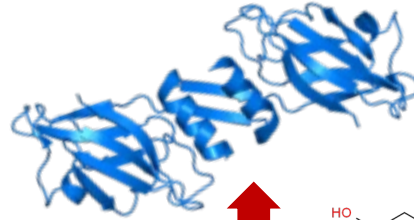
Product	Indication	Preclinical	IND Submission	Phase I	Phase II	Phase III	Phase IV
ProLectin-M <i>Oral</i>	Virology – Mild to Moderate • Covid-19	→					
ProLectin-I <i>Intravenous</i>	Virology – Severe cases • Covid-19 • Long Covid	→					
ProLectin-F <i>Intravenous</i>	Fibrosis: • Pulmonary Fibrosis • Other Fibrotic Conditions	→					
ProLectin-X <i>Miscellaneous</i>	Other Viral Indications: • Conjunctivitis (in progress) • Influenza (evaluation) • RSV (evaluation)	→					
ProLectin-A + Oxsense* <i>Intravenous combination treatment</i>	ARDS as result from viral infection	→					

* FDA 510(k) Clearance

Therapeutic Approaches for COVID-19



Immunomodulatory



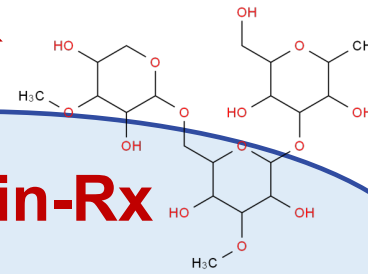
Antiretrovirals

(Compete for polymorphism)



ProLectin-Rx

- Unique Galectin Antagonist – Oral/IV Polysaccharide
- It's not a vaccine, nor an antiretroviral drug

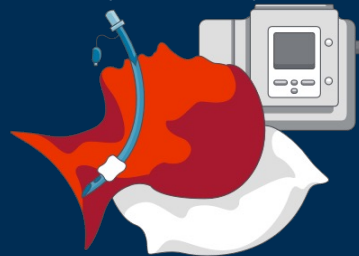


Vaccines



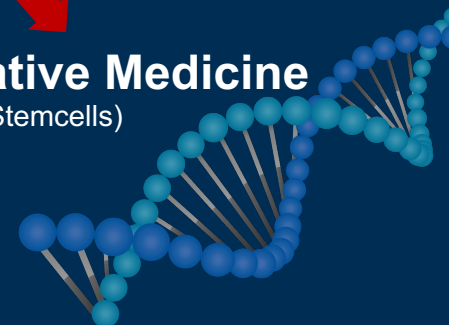
Oxygenation

(Ventilators)



Regenerative Medicine

(Stemcells)



Competitive Landscape

Oral COVID-19 Therapeutics

Drug	Company	Description	Worldwide Sales	Clinical stage
Molnupiravir	Merck	Mutagenesis via RdRp – forced mutations induced apoptosis	\$5.5 billion	EUA
Paxlovid	Pfizer	3CL protease inhibitor – Antiviral & Immune sensitization; Ritonovir – inhibitor enhancer	\$20 billion	EUA
Tollovir	Todos Medical	3CL protease inhibitor – Antiviral & Anti-Cytokine activity	n/a	Phase 2/3
Tempol	Adamis	RNA-dependent RNA Polymerase (RdRp) via antioxidant & Anti-Cytokine activity	n/a	Phase 2/3
Ensovibep (intravenous)	Molecular Parnters/ Novartis	DARPin domains attach to the viral spike protein	n/a	Phase 2/3

BXT-25 – Hypoxia & Degenerative Diseases



Solution: BXT-25

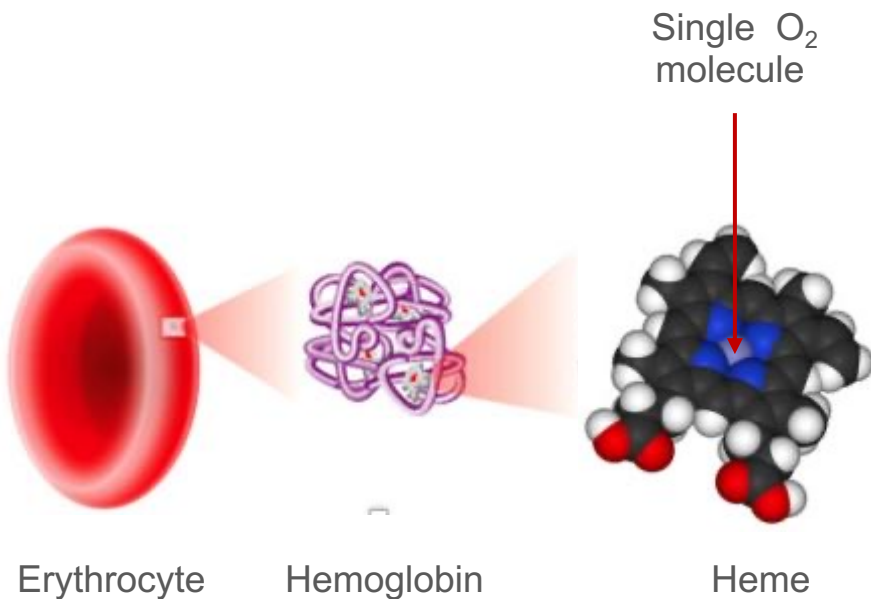
an Oxygen Bridge

- BXT-25 is a hemoglobin-based polymer; 5,000 times smaller than a red blood cell
- It can be used both in Ischemic and Hemorrhagic Stroke
- It can penetrate a blood-clot and reach the brain within 3 minutes
- Reduction of average Time-to-Needle by 90%



BXT-25
polymer

How It Works? BXT-25 – Stabilized Oxygen-carrying Protein



- Delivered as an IV solution
- Universally compatible with all blood types
- Non-immunogenic
- Low viscosity
- Stable at room temperature
- 3-year shelf-life in liquid formulation
- Extended shelf-life in dry formulation

Laboratory production-line developed & 1st batch GLP material manufactured
Developing the scale up methods & Pre-clinical trials pending

Degenerative Disease/Hypoxia Development Pipeline

➡ Completed ➡ Planned

Treatment/Device	Indication	Init Drug & Process Dvlp	Preclinical	Phase I	Phase II	Phase III	Phase IV
BXT-101	Cancer Metastasis	➡	➡				
BXT-102	NASH <ul style="list-style-type: none"> Cirrhosis Fibrosis 	➡	➡				
BXT-251+ Oxsense	Organ Transplantation <ul style="list-style-type: none"> Preservation agent Organ monitoring 	➡	➡				
BXT-25	Stroke <ul style="list-style-type: none"> Ischemic Hemorrhagic 	➡	➡				
BXT-252	Wound Healing	➡	➡				
BXT-253	Anemia	➡	➡				
BXT-255	Traumatic Brain Injury	➡	➡				

Commercialization & Targets

Clinical Trial Strategy

➡ Completed ➡ Planned

Product	2022				2023			
	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4
ProLectin-M	Drug & Process Development		Pre-clinical Studies	Phase II	Phase III			
ProLectin-I	Drug & Process Development		Pre-clinical Studies	Phase I	Phase II	Phase III		
ProLectin-F	Drug & Process Development		Pre-clinical Studies	Phase I	Phase II	Phase III		
ProLectin-A / BXT-25	Drug & Process Development				Pre-clinical Studies			Phase I/II

Use of Proceeds

Current Round	ProLectin-M	ProLectin-I	ProLectin-F	ProLectin-Rx*
Estimated Project Cost in thousands USD*	\$ 2,700	\$ 1,650	\$ 1,000	\$ 5,350
Development & GMP	-	-	-	-
Pre-Clinical	100	150	150	400
IND Submission	150	200	200	550
Clinical Trials	2,000	1,000	500	3,500
G&A	450	300	150	900
End Point	Phase III	Phase II	Phase II	Total
* \$5.1 million have previously been spent on proof-of-concept and GMP manufacturing of ProLectin-M, -I, and -F				

Future Round	ProLectin-A	BXT-25	Total Upcoming
Estimated Project Cost in thousands USD*	\$ 10,000	\$ 10,000	\$ 20,000
Development & GMP	3,150	3,150	6,300
Pre-Clinical	1,200	1,200	2,400
IND Submission	300	300	600
Clinical Trials	4,000	4,000	8,000
G&A	1,350	1,350	2,700
End Point	Phase II	Phase II	Total

Commercialization & IP Strategy

Intellectual Property (IP)

- Three (3) issued international patents
- One (1) issued US patent
- One (1) provisional US patent
- Additional applications to strengthen our IP position are ongoing*

* Future patents to be filed at commercialization stage



Licensing



Partnership



Divesting
product line

Comparables

COVID-19 Antivirals


 **MERCK**
\$425 million

BIOTECH

Merck inks \$425M Oncolmmune buyout to bag COVID-19 drug

By Nick Paul Taylor • Nov 23, 2020 08:35am

COVID-19 • mergers and acquisitions • coronavirus • Merck





\$350 million

Roche to pay Atea \$350M for ex-U.S. rights to COVID-19 antiviral

By Nick Paul Taylor • Oct 22, 2020 08:40am

antiviral • COVID-19 • licensing • Atea Pharmaceuticals



 **NOVARTIS**
\$580 million & 22% royalties

Novartis Signs an Option and License Agreement with Molecular Partners to Develop Two DARPin Therapies for COVID-19

Tuba Khan Oct 28, 2020 COVID-19



Novartis Signs an Option and License Agreement with Molecular Partners to Develop Two DARPin Therapies for COVID-19

Shots:

- Molecular Partners to receive \$65.8M as up front- including equity- and will receive \$164.7M as an option payment for both MP0420 and MP0423 along with royalties on sales of therapies. Novartis to get an option to in-license global rights of MP0420 and MP0423
- During the option period- Molecular Partners will conduct a P-I study for MP0420- expected to initiate in Nov'2020- and perform all remaining preclinical work for MP0423 while Novartis will lead P-II & P-III study

Big Pharma Licensing Targets

THERAPEUTIC AREAS

Virology

Oncology

Immunology

Respiratory

Inflammation

Neuroscience

Cardiovascular

Hematology

Stroke

IDEAL PLATFORM DRUG
FOR MANY BIG PHARMAS

- **AbbVie** – Oncology, Neuroscience, Immunology, Virology
- **Allergan** – Neuroscience, Ophthalmology, Gastroenterology
- **Amgen** – Oncology, Cardiovascular, Hematology, Inflammation
- **AstraZeneca** – Oncology, Cardiovascular, Respiratory
- **Biogen** – Oncology, Neuroscience, Inflammation, Stroke, Pain
- **Boehringer Ingelheim** – Oncology, Neuroscience, Immunology, Ophthalmology
- **Bristol Myers Squibb** – Oncology, Immunology, Cardiovascular, Hematology, Inflammation
- **Eli Lilly** – Oncology, Neuroscience, Immunology, Diabetes, Pain
- **Gilead** – Oncology, Respiratory, Hematology, Inflammation, HIV
- **GlaxoSmithKline** – Oncology, Immunology, Respiratory, HIV, Vaccines
- **Johnson & Johnson** – Oncology, Neuroscience, Immunology, Cardiovascular, Vaccines, HIV
- **Merck** – Oncology, Neuroscience, Immunology, Cardiovascular, Respiratory, Diabetes, Vaccines
- **Novartis** – Oncology, Neuroscience, Immunology, Cardiovascular, Respiratory, Ophthalmology
- **Pfizer** – Oncology, Neuroscience, Cardiovascular, Diabetes
- **Roche Holdings** – Oncology, Neuroscience, Immunology, Hematology, Ophthalmology
- **Sanofi Aventis** – Oncology, Neuroscience, Immunology, Inflammation, Diabetes, Vaccines



75 2nd Ave., Suite 605
Needham MA, 02494
(617) 454-1199
www.bioxytraninc.com
info@bioxytraninc.com

Mike Sheikh

mike.sheikh@bioxytraninc.com
(509) 991-0245