

ORAGENICS, INC MANAGEMENT UPDATE CALL MAY 20TH, 2025

NYSE: OGEN



MAY 2025



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JANET HUFFMAN CHIEF EXECUTIVE OFFICER





2025

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OVERVIEW - KEY OBJECTIVES FOR THE CALL

Leadership & Strategic Vision - Introduce the team guiding Oragenics through transformation and explain our focused mission in concussion care.

Investor Alignment & Market Opportunity - Discuss how our strategy aligns with unmet market needs—and why ONP-002 represents a significant commercial opportunity.

Clinical and Regulatory Progress - Provide a transparent update on ONP-002's development, including recent approvals, trial readiness, and next-phase planning.

Milestones & Execution Timeline - Walk through key upcoming catalysts across clinical, regulatory, and business development milestones.

Next Steps & Stakeholder Engagement - Outline how we plan to strengthen communication, pursue strategic partnerships, and deliver on near-term value drivers.



MANAGEMENT OVERVIEW

Janet Huffman, Chief Executive Officer:

Bio Highlights:

- Appointed CEO of Oragenics in April 2025 after joining the company in March 2023.
- Brings over 15 years of executive leadership experience across healthcare sectors, including home health, skilled nursing, rehab, pharmacy, and health IT.

Prior Roles Include:

- CFO, TRxADE HEALTH (NASDAQ: MEDS): Led finance for a digital health services company focused on retail pharmacies.
- Founding CFO, Banyan Pediatric Care Centers: Key architect of its merger with Assisted 4 Living, later renamed Arboreta Healthcare Inc.
- Held senior financial leadership roles at Signature HomeNow, Infinity Homecare, and Family Home Health Services, overseeing growth, M&A, and operational transformation.



Our Team Board and Scientific Team

Board of Directors Highlights: Longtenured leadership with biotech, finance, and regulatory experience

Neurological Team: Experts in TBI, emergency medicine, neurosteroids (Dr. Peacock, Dr. Kelly)

THE PROBLEM - CONCUSSION IS A PUBLIC HEALTH CRISIS

Concussions Are a Subset of Traumatic Brain Injury (TBI)

- A concussion is classified as a mild traumatic brain injury (mTBI)—the most common form of TBI.
- While considered "mild," concussions can cause serious neurological symptoms that may <u>last weeks, months, or longer</u>.
- All concussions are TBIs, but not all TBIs are concussions. TBI severity ranges from mild (concussion) to severe (bleeding, coma, permanent impairment).
- Concussions often go <u>undiagnosed</u> because they may not appear on standard imaging, yet they disrupt brain function in real and lasting ways.
- Understanding concussion as part of the broader TBI spectrum highlights the critical need for early, targeted intervention.

O R A G E N I C S

Assault 9%

Traffic Incidents 14%





OUR SOLUTION - ONP-002 INTRANASAL THERAPY

ONP-002: First-in-class intranasal drug for moderate-to-severe concussion

- Rapid brain biodistribution
- Reduces swelling, inflammation, and oxidative stress
- Completed Phase 1 clinical trial: Safe and well-tolerated

Strategic Impact

- Expanded Oragenics' pipeline into neurology and intranasal drug delivery.
- Targets high-growth markets, including a concussion treatment market projected to reach \$8.9 billion by 2027.
- Addresses a significant unmet need: An estimated 5 million concussions occur annually in the U.S., including 3.8 million from sports-related injuries—up to 50% of which go unreported. ORAGENICS



RECENT CLINICAL ACCOMPLISHMENTS

Clinical Progress:

<u>Significant progress, with patient enrollment expected to begin in Q2 2025 in Level 1 trauma</u> emergency departments in Australasia.

Phase 1:

Clinical trial reports and data have been finalized and delivered. The regulatory team is organizing the data for submission in an IB and IND package.

Phase 2:

- HREC Approval Australia (April 2025): Received Human Research Ethics Committee (HREC) clearance to initiate Phase II clinical trial of ONP-002.
- HDEC Submission New Zealand (April 2025): Health and Disability Ethics Committee (HDEC) allowing trial to be conducted at Christchurch Hospital, a leading emergency and research institution serving over 83,000 patients annually.

RECENT STRATEGIC ACCOMPLISHMENTS

Department of Defense (DoD):

Submitted a grant application to support ONP-002 development for concussion care.

Strategic partnership with BRAINBox Solutions:

Entered into a partnership with BRIANBox to create a first-of-its-kind test-to-treat platform for concussion. Combines ONP-002 with BRAINBox's blood biomarker diagnostic tools.

- Leader in multi-modality diagnostics for TBI, to co-develop the first comprehensive "trigger-to-treat" platform for concussion. This collaboration represents a transformative growth step.
- Enhancing patient selection and precision enrollment for the upcoming Phase IIa trial—improving data quality, reducing variability, and enabling a potentially faster path to clinical proof-of-concept.
- Establishing a differentiated commercial model that positions Oragenics not just as a drug developer, but as part of a next-generation care pathway for mTBI—delivering both the diagnosis and the treatment.





DR. JIM KELLY CHIEF MEDICAL OFFICER

ONE OF THE NATION'S FOREMOST AUTHORITIES ON CONCUSSION AND TRAUMATIC BRAIN INJURY, LEADING CLINICAL STRATEGY FOR ONP-002.

- Former Director, NICoE at Walter Reed, advancing military TBI care
- Longtime neurological consultant to the NFL; helped shape modern concussion protocols
- Co-author of the American Academy of Neurology's concussion guidelines
- Former Professor of Neurology, University of Colorado School of Medicine
- Renowned for bridging clinical research, neurorehabilitation, and real-world sports medicine

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WHAT HAPPENS IN THE BRAIN **AFTER A CONCUSSION?**

Head Trauma - Ionic Shift - Inflammation - Oxidative Stress - Brain Swelling - Synaptic Dysfunction

- There are approximately 5M documented concussions in the US a year which exceeds the combined incidence of Stroke, Alzheimer's Disease, Parkinson's Disease, Multiple Sclerosis, and ALS aka Lou Gehrig's Disease.
- Annual worldwide cost for managing TBI to healthcare system is over \$400B
- Core groups driving the need for a concussion treatment are athletics, military, pediatrics and the rapidly increasing elderly population at risk for falls.
- Repetitive concussions may be linked to other chronic neurodegenerative conditions.

Concussions initiate a chain reaction of biological damage.

Secondary effects worsen over hours or days if untreated.

Synaptic disruption leads to memory, mood, and motor deficits.

The first 24 hours post-injury are the key therapeutic window.



MILD VS SEVERE TBI ACUTE BRAIN RESPONSE





CONCUSSION CLINICAL ASSESSMENT

<u>SAC</u> Standardized Assessment of Concussion FORM A Name:						
Agan G	Sex:Ex:	minar				
Age:	Sel:EX					
Nature of Injury: Date of Exam:Time:Exam No						
1) ORIENT	ATION:					
Month:			0 1			
Date:			A T			
Day of weel	k:		0 1			
Year:			0 1			
Time (withi	n 1 hr.):		0 1			
	Total Score		/5			
			completed regardless of			
	2; score equals sum					
List	Trial L	Trial 2	Trial 3			
Blbow	0 1	0 1				
Apple	0 1	0				
Carpet Saddie	0 1	0	0 1			
Bubble		0				
Total		V				
	Monto Tot	AL FARMA	/ 15			
	Memory Tot form the subject that					
	DGICAL SCI sciousness (p		1			
Recollection	<u>t of injury</u> (pr	e- or post-tra	aumatic amnesia)			
<u>Strength:</u>						
Sensation:						
<u>Coordinatio</u>	<u>m:</u>					

3) CONCENT	BATION:			
	<u>ard:</u> (If context, go b	o next swing to	ദ്യഥ	IE incorrect, read
4-9-3	6-2-9	0	1	
	3-2-7-9	ő	-	
6-2-9-7-1		ő	-	
7-1-8-4-6-2		Ŏ	-	
	<u>verse Order</u> : (em -Scp-Aug-Jul	ี่เฒ่ พระการคายด	uenca	e correct for 1 pt.)
Jun-May-Apr	-Mar-Fcb-Jan	0	1	
Concentratio	na Total Score		5	

EXERTIONAL M				
5 jumping jacks	5 push-ups 5 kncc-bends			
5 sit-ops				
4) DELAYED RECALL				
Elbow	0	1		
Apple	0	1		
Carpet	0	1		
Saddle	0	1		
Bubble	0	1		
Delayed Recall Total Score	1	5		

SUMMARY OF TOTAL SCOR	ES:
Orientation	/ 5
Immediate Memory	/ 15
Concentration	/ 5
Delayed Recall	/ 5
Overall Total Score	/ 30
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CONCUSSION CLINICAL SYMPTOMS

		None		oderate Seve	re			
Score According to Severity		0 1	2 34	5		6		40 -
Symptom	Preseason Baseline	Time of Injury	24 Hours Post- Injury	Day 3 Post- Injury	Day 4 Post- Injury	Day 5 Post- Injury		
Blurred Vision								
Dizziness								
Drowsiness								
Sleeping More than Usual								30 -
Easily Distracted								
Fatigue							_	
Feeling "In a Fog"							9	D
Feeling "Slowed Down"							CC Total Coore	,
Headache							ũ	5
Unusually Emotional							2	20
Irritability							3	20 -
Loss of Consciousness)
Loss of Orientation							<i>u</i>	2
Memory Problems							S	,
Nauseous								
Nervousness								
Personality Changes								10 -
Poor Balance/Coordination								
Ringing in the Ears								
Sadness								
Seeing Stars								
Sensitivity to Light								
Sensitivity to Noise								0 -
Sleep Disturbances								0
Vacant Stares/Glassy Eyes								
Vomiting								в
TOTAL SYMPTOM SCORE:								







ONP-002 - PRECLINICAL EFFICACY

ONP-002 – Molecular Studies:

- Rodent Reduces inflammation, oxidative stress and swelling in the injured brain
- Neuronal Culture Enhances brain cell survival and growth when challenged with low oxygen and glucose
- ONP-002 Behavioral Studies
 - Rodent Improves sensory and motor performance, anxiety and depression-like behaviors, and short-term memory following brain injury



ONP-002 - PRECLINICAL SAFETY AND BIODISTRIBUTION

ONP-002 – Toxicology Studies:

- Rodent, Canine, and Monkey Well-tolerated 2X-daily/14-days intranasally
- High safety margin (>20X) between animal dose and human dose (Phase I and upcoming Phase II trial) ONP-002 Brain Biodistribution
- Canine Intranasal delivery provides for a 4X higher level of drug in brain compared to plasma
- Similar levels are seen throughout brain regions within 30 min of administration

All IND-enabling studies have been completed for cardiac and genotoxicity, and drug: drug interactions



ONP-002 – CLINICAL PRODUCT MANUFACTURING

- ONP-002 is a novel, neuro-steroid created through a synthetic process that results in a pure, well defined compound.
- ONP-002 is formulated as a intranasal powder formulation manufactured into a nanoparticle size improving uptake from nose to brain.
- ONP-002 has been shown to be stable up to 104 degrees for 18-months and -20 degrees for 30-days, preventing need for temperature-chain protocols in the field.



A WINNING COMBINATION: ONP-002 + INTRANASAL DELIVERY



Intranasal Delivery Advantages

- Rapid and direct access to the brain
- Swift therapeutic effects
- Quicker onset compared to oral medications



- Innovative Design
 Patients blow into the device, elevating the soft palate
- Minimizes systemic exposure and side effects



User-Friendly and Portable

Compact and lightweight design ensures ease of use, making it accessible for patients in acute brain injury scenarios





• Innovative Double-Tube Airflow System: Optimizes drug dispersion to the nasal roof.

• <u>Direct Delivery</u>: Targets the olfactory nerve for brain delivery which should improve clinical outcomes.





DR. FRANK PEACOCK CHIEF CLINICAL OFFICER

A NATIONALLY RECOGNIZED EXPERT IN EMERGENCY MEDICINE AND BRAIN INJURY **DIAGNOSTICS, LEADING REAL-WORLD CLINICAL IMPLEMENTATION FOR ONP-002.**

- Vice Chair of Emergency Medicine Research, Baylor College of Medicine
- Former Professor at the Cleveland Clinic
- Principal Investigator on concussion biomarker trials in emergency settings
- Advanced the use of high-sensitivity blood troponins for cardiac diagnostics (featured in JAMA Cardiology)
- Editor of Biomarkers of Traumatic Brain Injury, a leading clinical reference in TBI research

ONP-002 – CLINICAL R&D

PHASE I CLINICAL STUDY REPORT (CSR) AND TRIAL MASTER FILE (TMF) COMPLETED – **ONP-002 WAS SAFE AND WELL TOLERATED IN 40 HEALTHY HUMAN VOLUNTEERS**

Lead site for Phase IIa – Christchurch Hospital, New Zealand (NZ) -CRO- Comprehensive Research Associates -Clinical Protocol Submitted to HDEC in NZ

The drug is loaded in intranasal devices for 40 patient Phase IIa trial (safety and feasibility) – 9-month stability complete

FDA-IND application submission planned for Q3 – Drug: Device Combination to CDER

Phase IIb plan for initiation - 2026 in the US – 12 initial sites



PHASE IIA - ENROLLMENT CRITERIA

Enrolment criteria designed to identify most at risk for developing Persistent **Concussion Symptoms beyond 30-days of injury**

First treatment within 12-hrs of injury

- Glasgow Coma Score of 13-15 lacksquare
- Negative CT Scan
- Positive GFAP score \bullet
- Report of Headache \bullet
- History of one or more of the following: Loss of Consciousness, short-term lacksquareamnesia, Altered Mental Status



PHASE IIA TRIAL - OUTCOME AND SAFETY MEASURES

Safety

- Monitoring for AEs/SAEs Patient reported and Physical Examination
- EKG
- Drug PK levels
- Intranasal evaluation for irritation

Clinical Outcomes

- Patient reported symptoms Rivermead
- Neurocognitive performance DANA and Braincheck
- Visual-Vestibular analysis King-Devick Testing
- Blood Biomarkers levels 1st clinical analysis for surrogacy of recovery
- Patient reported function GOS-E
- Incidence of Persistent Concussive Symptoms at 30-days post injury



BRAINBOX PARTNERSHIP PHASE IIA

Trigger to Treat Model

Kit provisions for blood draws, strips for blood biomarker analysis, and tabletop readers

Blood draw timepoints, per-dose (within 12 hours of injury), 24 hours, 72 hours, 5-days, 10days, and 30-days

GFAP – elevation required for study admission with continued monitoring ST2

GFAP elevation required for study inclusion to assist in identifying the most severe concussions with poorer outcomes

Neurogranin

A brief neurocognitive test will be administered at the time of each blood biomarker analysis for correlation



y), 24 hours, 72 hours, 5-days, 10-

ONP-002: CLINICAL MILESTONES



ORAGENICS

WHY ONP-002? WHY ORAGENICS? WHY NOW?

<u>The Opportunity Is Clear</u>

- Large Market Opportunity: Concussions affect 5M+ Americans annually
- First of its kind: No FDA-approved treatment
- ONP-002 targets the root causes of concussion damage: inflammation, oxidative stress, and brain swelling
- Intranasal delivery platform: enables fast, non-invasive, field-ready use
- Strong Phase I safety data and Phase IIa trial launching now in real-world trauma settings
- Strategic partnerships and regulatory momentum position us for value inflection
- Led by a team of clinical and operational experts in TBI, neurology, and emergency medicine





Q&A SESSION THANK YOU

ORAGENICS

MAY 2025

