

Euroz Hartleys Rottnest Island Institutional Conference

March 2024

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Botanix – accelerating towards commercialization of *Sofdra*™

| DERMATOLOGY FOCUS | New treatments for underserved common skin diseases, with an initial focus on excessive sweating ("primary axillary hyperhidrosis") |
|-------------------------------|---|
| TOPICALLY DRIVEN | Targeting key indications with topical (gel) treatments that are designed for safety, tolerability, and clinical efficacy |
| EXPERIENCED TEAM | US-based team that has been responsible for successful development and commercial launches of more than 30 dermatology drugs |
| NEW PRODUCT "SOFDRA" | Sofpironium Bromide <i>(Sofdra)</i> ¹ is the first and only new chemical entity developed for primary axillary hyperhidrosis (5% strength approved in Japan with solid sales) ² |
| TARGETING MID-24 FDA APPROVAL | Resubmission of NDA for approval was completed in late December 2023; targeting FDA approval in late June 2024 |



1. Sofdra (sofpironium bromide gel, 15%) is an investigational drug and is not FDA approved. The Sofdra brand name is under review by FDA.

2. ASX release May 4, 2022.

Corporate Overview

Well-funded to FDA approval, supported by leading life science institutional investors

| ASX: BOT TRADING INFORMATION | | | | | |
|------------------------------|---------------|--|--|--|--|
| Share price | A\$0.185 | | | | |
| 6-month low / high | A\$0.12/0.20 | | | | |
| Shares outstanding | 1,563,437,373 | | | | |
| Market Capitalization | A\$275m | | | | |
| Cash | A\$ 18.3m | | | | |
| Debt | Nil | | | | |
| | | | | | |

SUBSTANTIAL SHAREHOLDERS

| | Shareholder | % | | |
|--|----------------------|------|--|--|
| | Antares Capital | 9.0% | | |
| | Board and Management | 7.0% | | |
| | Тор 20 | 33% | | |

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Hyperhidrosis

A medical condition where excessive sweating occurs beyond what is needed to maintain normal body temperature



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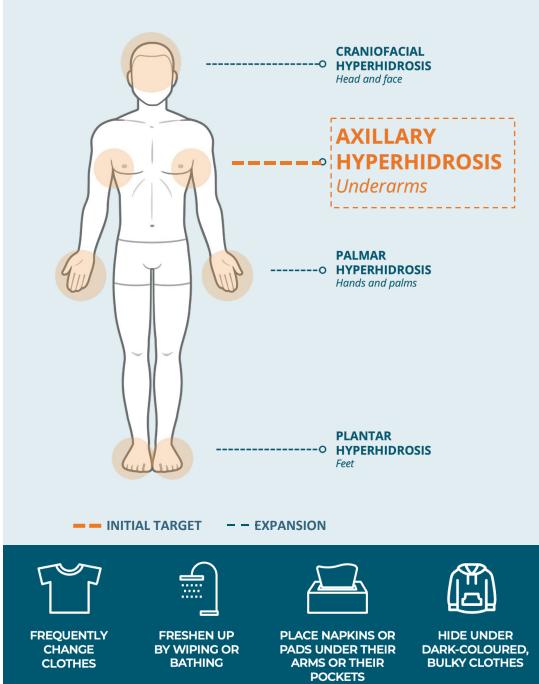
Hyperhidrosis affects ~16M people in the US¹

Results from overstimulation of the nervous system (a physiological not psychological condition)¹

90% of axillary (underarm) patients also have it in a second region¹

The most common age of onset for axillary hyperhidrosis patients is 12–17²

Market for treatments is ~\$US1.6B per annum projected to grow to \$US2.8B by 2030²



Our lead asset: Sofpironium Bromide (Sofdra)¹

The only new chemical entity developed specifically for the treatment of primary axillary hyperhidrosis

Met both co-primary endpoints in two Phase 3 trials²

- 60% of subjects had ≥2-point improvement in HDSM-Ax
- 65% had a significant reduction in GSP sweat production
- Met all secondary endpoints including clinically meaningful effect on 85% of patients
 - ≥1-point improvement in HDSM-Ax
 - Statistically significant improvement

Favorable tolerability and safety profile³

Well-tolerated with adverse events that were mostly mild or moderate, and events were transient



Sofdra (sofpironium bromide gel, 15%) is an investigational drug and is not FDA approved. The Sofdra brand name is under review by FDA

Two identical randomized, double-blinded, vehicle-controlled Phase 3 trials for primary axillary hyperhidrosis (pooled; sofpironium bromide gel, 15% n=353; vehicle n=348)
Dry mouth and blurred vision were the predominant treatment-emergent adverse events at 14.4% and 8.5%, respectively, and are common among anticholinergic drugs

Innovative launch strategy to accelerate adoption following approval

Rapidly establish *Sofdra* as a safe and effective first line treatment of primary axillary hyperhidrosis in adults and pediatric patients 9 years of age and older



Drive Derm adoption through comprehensive engagement around a compelling clinical story



Engage and motivate patients to take control of their hyperhidrosis by visiting a telemedicine doctor for a diagnosis and prescription



Maximize favorable coverage



Provide patient access and immediate fulfillment through telemedicine and pharmacy network with mail-order fulfillment to drive trial while ensuring compliance



Hire and train a highly effective Sales Force

Planned launch activities targeting high prescribers of HH products

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In-office rep activities will include video, animation, and printed leave behinds

Digital advertising to drive targeted prescribers to SofdraHCP.com



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Patient launch activities to target active HH information seekers

Planned search engine optimization/marketing and all materials will drive patients to Sofdra.com Planned social media and digital advertising will drive patients to quiz on Sofdra.com



Images of marketing materials are for representative purposes only

Proactive, pre-approval engagement with Payors with >200K lives

Optimize access ahead of planned launch

| | Rx Con PBM | Account Lives | | Lives | Rnk | Clin Pro | es | |
|------------|------------------|--------------------|--|-------------------|------------|----------|-----|-----------|
| 6 | CVS | CVS Caremarl | CVS Caremark - Advanced Control, Performance Standard Control, Value 1,8 | | 1 | Yes | | |
| Ē | EXPRESS | | Express Scripts - High Performance, Basic 1,718,678 | | | Yes | | |
| | EMISAR ASCENT | Rx Con PBM Account | | | Lives | 5 | Rnk | Clin Pres |
| | ASCENT | ZINC | CVS Caremark - Advanced Control, Performance Standard | 30,650,000 | | 1 | Yes | |
| 6 | N/A | ASCENT | Express Scripts - National Preferred Formulary | | 26,709,534 | | 1 | Yes |
| Q | ZINC | EMISAR | OptumRX Premium Standard, Value, Select Standard | | 15,435,000 | | 1 | Yes |
| P | PROCARE | ZINC | Anthem Essential HMO, PPO, National, Traditi | 12,833,835 | | 2 | Yes | |
| U | PRIME | EMISAR | United Healthcare- Access, Advantage, Choice, Esse | 12,658,000 | | 2 | Yes | |
| | | ASCENT | Cigna- Advantage, National Preferred, Perform | 8,760,900 | | 2 | No | |
| | ASCENT | KAISER | Kaiser Permanente | 8,303,484 | | 1 | Yes | |
| | EMISAR | TRICARE | TriCare | | 7,214,213 | | 2 | Yes |
| | EMISAR | ZINC | AETNA- Open, Standard, Fully Insured | | 5,958,336 | | 2 | Yes |
| $(\square$ | DIVIDEND | CVS | (FEHBP)- Basic, Focus, Standard | | 5,330 | ,051 | 1 | Yes |
| 9 | | DoD | DEPARTMENT OF VETERANS AFFAIRS | | 4,701 | ,838 | 2 | Yes |
| | NAVITUS | PRIME | BCBS IL/ Tx/NM/MT (HCSC)- HMO or PPO Enhanced, Perfor | mance, Multi Tier | 4,575 | ,000 | 2 | No |
| | | ASCENT | Prime Therapeutics | | 2,460 | ,000 | 2 | Yes |
| | \supset) | PRIME | BCBS FL- HMO, PPO Multi Tier | | 2,125 | ,000 | 2 | No |

- Completed Payor profiles and engagement plan
- Engaged target Payors around unmet need in primary axillary hyperhidrosis and *Sofdra* value proposition
- Confirmed hyperhidrosis reimbursement status as medical condition
- Commenced initial discussions with target Payors responsible for 80% of covered lives

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Docs will e-prescribe directly to our national pharmacy network

Instructions are provided to patient in doctor's office when prescription is written

Strong value and convenience messaging includes capping patient's out-of-pocket cost

QR code to enter instantly into digital space and begin interaction with our pharmacy network

Pharmacy mails Sofdra the same day that the patient completes their intake form



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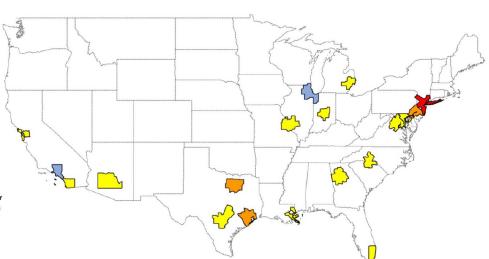
Territories created based on prescriptions and HH diagnosis data

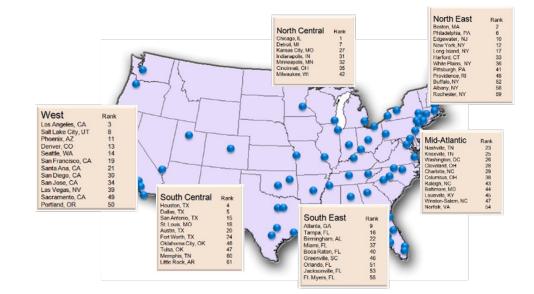
National programs focused on educating physicians and office staff

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Territories aligned with prescriber and HH Data

Targeted cities based on prescriber and HH data





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Focused pre-launch period ahead

- FDA approval targeted for late June 2024
- The issue being considered by the FDA is related to patient Instructions For Use—no efficacy, safety or manufacturing issues remain
- Commercial preparation is accelerating in anticipation of FDA approval
- Company is funded through approval and has multiple commercialization options



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