

27 July 2022

Euroz Hartleys Healthcare Conference Presentation

Key highlights

- Release of updated corporate presentation to be delivered at Euroz Hartleys Healthcare
 Conference being held today in Perth including a number of other leading ASX listed companies
- The presentation includes:
 - an update on the accelerated FDA submission timetable of the New Drug Approval (NDA) for Sofpironium Bromide which is now planned for this quarter;
 - o insights into the market opportunity for Sofpironium Bromide; and
 - confirmation of timetable for completion of rosacea (BTX 1702) Phase 1/2 clinical study and the canine dermatitis pilot study (BTX 1204A in Q3 CY2022

Philadelphia PA and Phoenix USA, 27 July 2022: Clinical dermatology company, Botanix Pharmaceuticals Limited (ASX: BOT, "Botanix" or "the Company"), is pleased to release a corporate presentation being used for the Euroz Hartleys Healthcare Conference being held today in Perth.

Botanix's Chief Commercial Officer, Howie McKibbon, will be presenting an update on Sofpironium Bromide for which the FDA filing has been accelerated and is now planned for this quarter, further insights into the market opportunity for Sofpironium Bromide, as well as an update on the Company's BTX 1702 Phase 1b/2 rosacea study and the BTX 1204A Phase 1b canine dermatitis study, which are on target for completion in Q3 CY2022.

Executive Director Matt Callahan will also be presenting some thoughts on the FDA drug development and approval processes and how companies (and investors) can mitigate risk in development.

Release authorised by

Vince Ippolito

President and Executive Chairman

About Botanix Pharmaceuticals

Botanix Pharmaceuticals Limited (ASX:BOT) is a dermatology company based in Philadelphia and Phoenix (USA) which is committed to the development of novel treatments for a range of common skin diseases. The Company has a mature dermatology pipeline with its first product, Sofpironium Bromide, for the treatment of primary axillary hyperhidrosis, planned to be filed for FDA in Q3 CY2022. The Company also has a pipeline of other products in late-stage clinical studies for the treatment of moderate to severe rosacea, dermatitis and acne respectively. Botanix is also developing a topical



antimicrobial product for the eradication of bacteria on the skin surface, initially in patients who are undergoing hemodialysis.

Botanix leverages its proprietary drug delivery system (PermetrexTM) for direct skin delivery of active pharmaceuticals in all skin diseases, which is utilised in its existing development programs and is being explored with a view to being utilized in a number of other product opportunities. To learn more please visit: http://www.botanixpharma.com/

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Any statements in this press release about future expectations, plans and prospects for the Company, the Company's strategy, future operations, the expected timing and/or results of regulatory approvals and prospects of commercialising product candidates or research collaborations with its partners, including in Japan, the outcome and effects of Sofpironium Bromide and the market for Sofpironium Bromide and other statements containing the words "anticipate," "believe," "estimate," "expect," "intend," "may," "plan," "predict," "project," "target," "potential," "will," "would," "could," "should," "continue," and similar expressions, constitute forward-looking statements. Actual results may differ materially from those indicated by such forward-looking statements as a result of various important factors, including: the Company's or its partners' ability to successfully develop its product candidates and timely complete its planned clinical programs and the Company's or its partners' ability to obtain marketing approvals for is product candidates. In addition, the forward-looking statements included in this press release represent the Company's views as of the date hereof. The Company anticipates that subsequent events and developments will cause the Company's views to change. However, while the Company may elect to update these forward-looking statements at some point in the future, the Company specifically disclaims any obligation to do so. These forward-looking statements should not be relied upon as representing the Company's views as of any date subsequent to the date hereof.

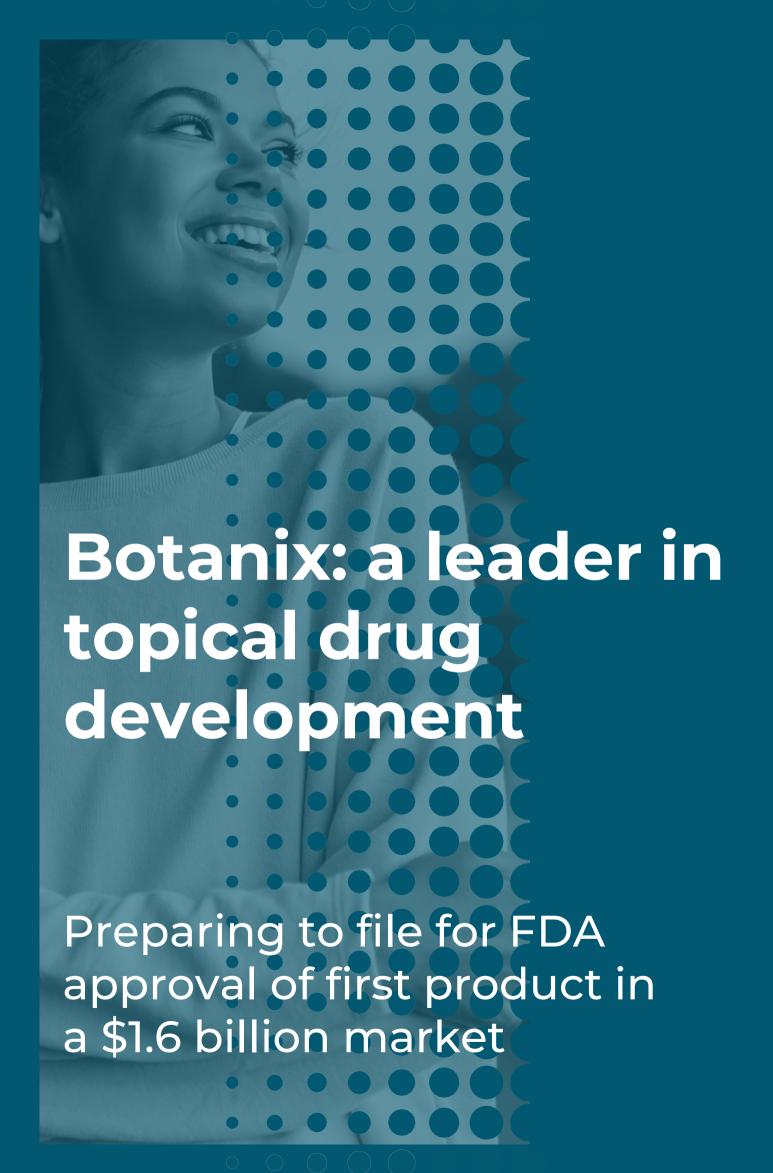
Euroz Hartleys Healthcare Conference

July 2022

Preparing to file first dermatology product for FDA approval in Q3 2022









Pharmaceutical focus

New treatments for common skin diseases - such as excessive sweating (hyperhidrosis), rosacea and acne



Topically driven

Targeting key indications with topical treatments that are safe, well tolerated and validated with clinical efficacy



World class team

US based team that have been responsible for more than 30 dermatology drug developments and launches



Sofpironium Bromide

First and only new drug for "primary axillary hyperhidrosis" (excessive underarm sweating) already approved in Japan¹



Near-term catalysts

Upcoming filing for FDA approval for Sofpironium Bromide and data readouts from other pipeline products in 3Q CY2022

Source 1: ASX release May 4 2022

Corporate Overview

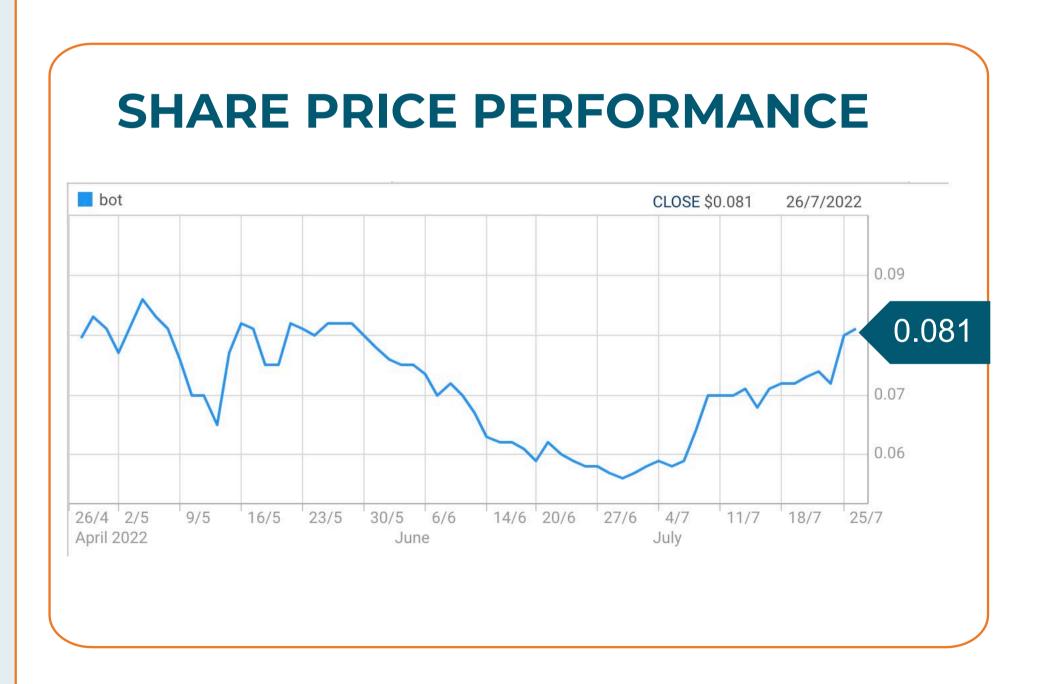
Positioned to transition to commercial dermatology company

ASX: BOT TRADING INFORMATION

Share price	A\$0.081
6-month low / high	A\$0.052/0.095
Shares outstanding	979,233,384
Market Capitalisation	A\$79.3m
Cash (31 Mar 2022)	A\$ 16.4m
Debt (31 Mar 2022)	Nil

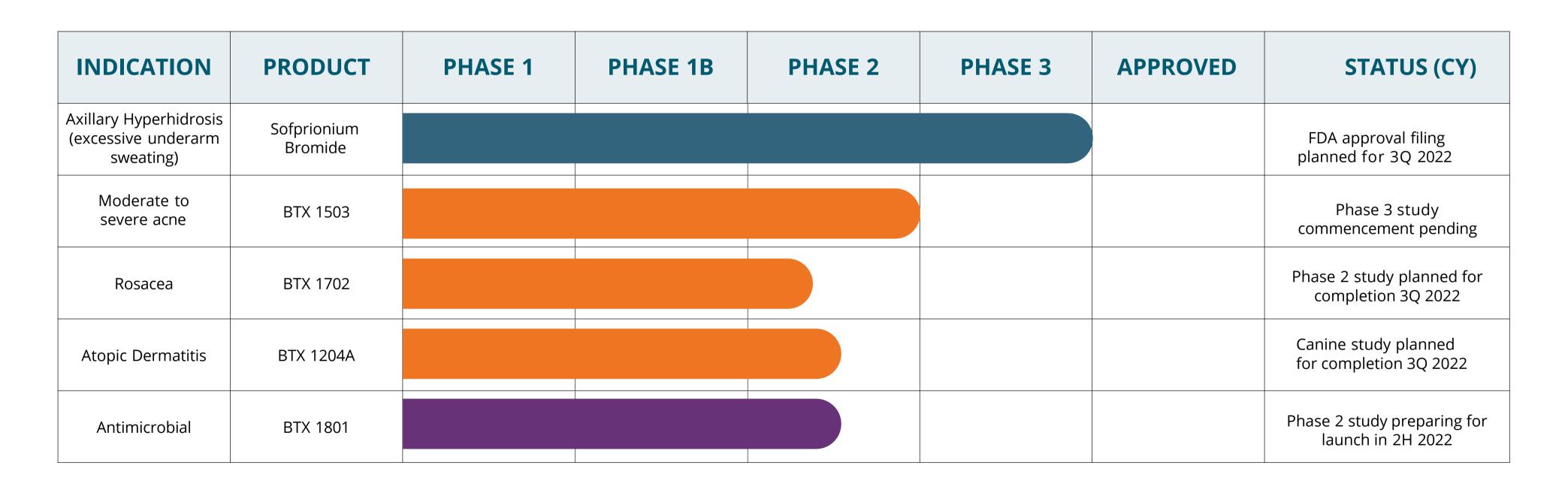
SUBSTANTIAL SHAREHOLDERS

Shareholder	%
Board and Management	8.29%
Caperi Pty Ltd, Co-Founder	5.4%



Sofpironium Bromide leads late-stage pipeline

Planned for FDA filing in 3Q 2022 with 12-month review period



Sofpironium Bromide is a significant opportunity in its own right, but also fits well alongside acne, rosacea and dermatitis



Accelerating Botanix towards revenue generation

Newly acquired asset Sofpironium Bromide being prepared for FDA approval filing in 3Q 2022



Addressing unmet needs

First and only new chemical entity for "primary axillary hyperhidrosis"



Positive Phase 3 Data

All co-primary and secondary endpoints were statistically significant and side effects were mild to moderate with no treatment-related serious adverse events



Attractive Terms

Minimal upfront payment and back-ended deal decreases risk and allows Botanix to share success when it's achieved (based on commercial success)



Significant Market

More than 16 million people suffer from hyperhidrosis in the US alone and market for treatments is ~\$US1.6B per annum which is projected to grow to \$US2.8B by 2030^{1,2}



De-risked Asset

Molecule already approved by Japanese equivalent of the FDA with partner Kaken Pharmaceuticals and recently launched in Japan

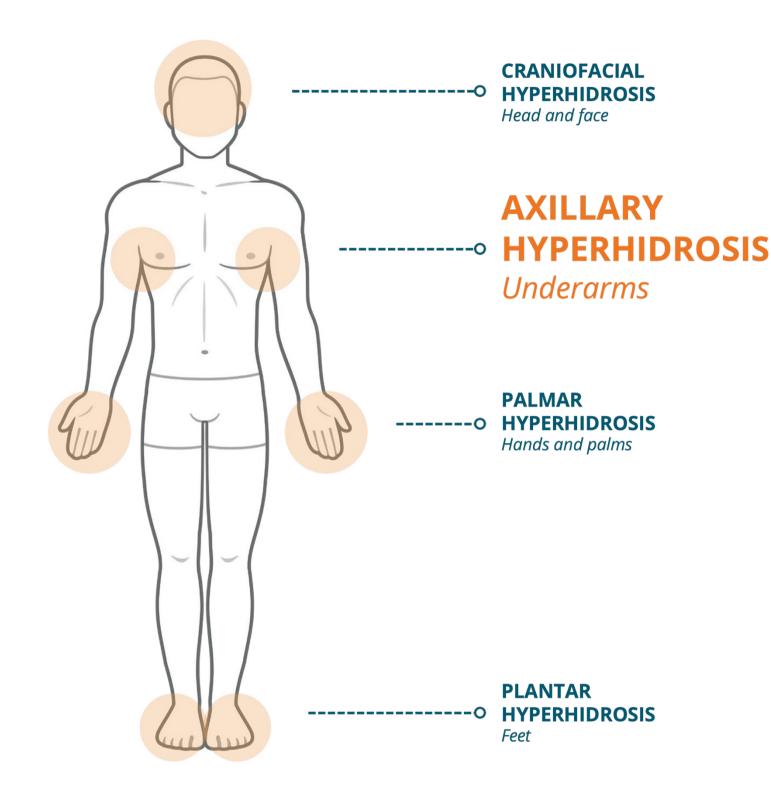
Hyperhidrosis

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

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 2030





FREQUENTLY CHANGE CLOTHES



FRESHEN UP BY WIPING OR **BATHING**



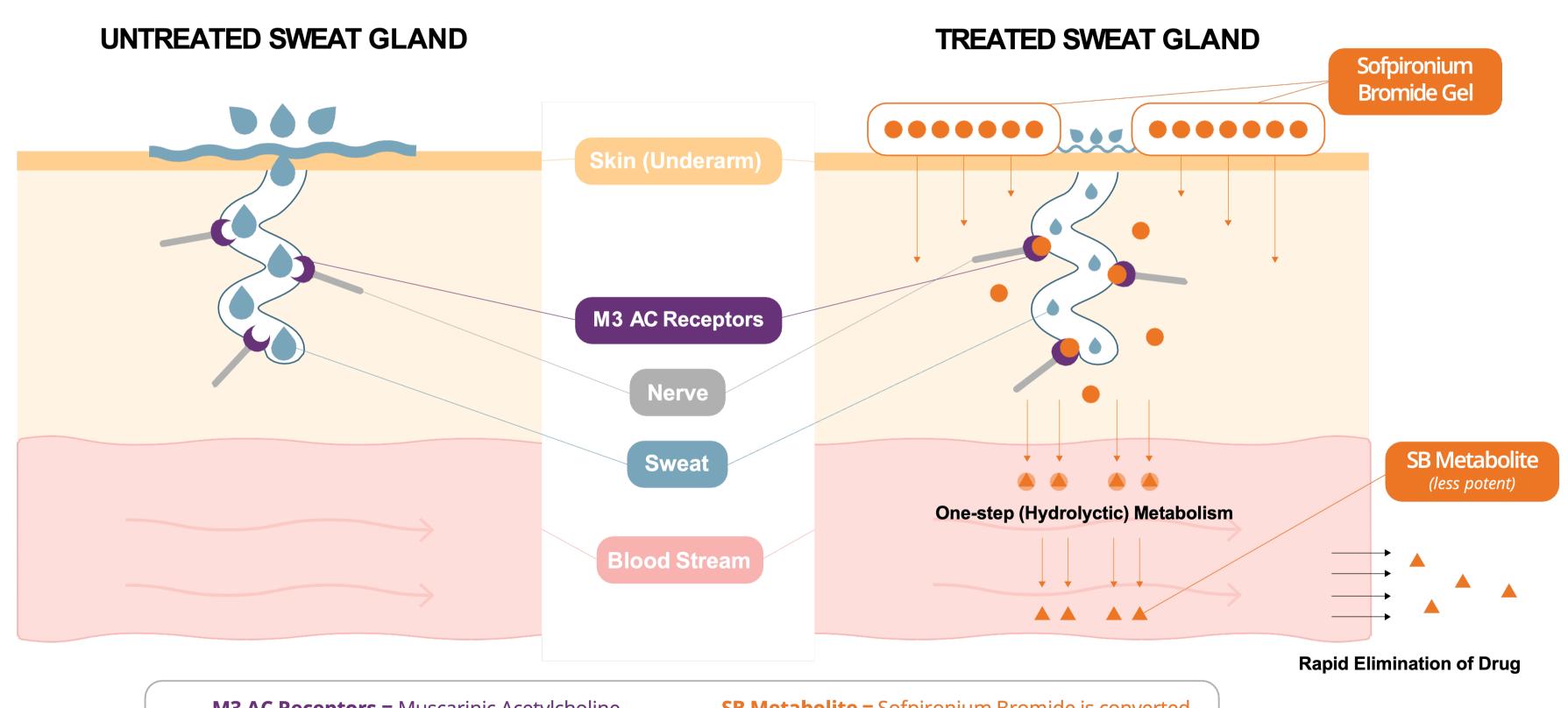


PLACE NAPKINS OR PADS UNDER THEIR **ARMS OR THEIR POCKETS**

HIDE UNDER DARK-COLOURED **BULKY CLOTHES**

Sofpironium Bromide mechanism of action

Blocks sweat gland receptors and rapidly degrades for excretion



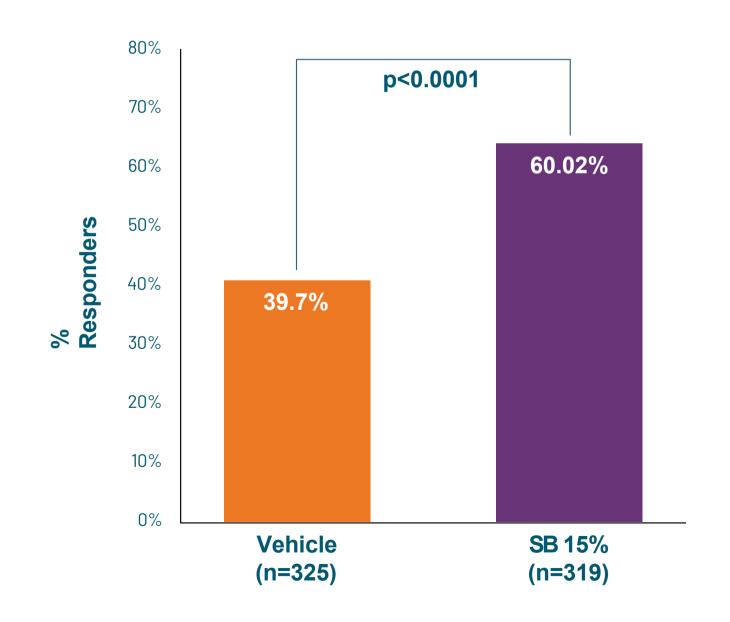
M3 AC Receptors = Muscarinic Acetylcholine Receptors which regulate the function of sweat glands **SB Metabolite** = Sofpironium Bromide is converted into a less active form to help minimize side effects

Phase 3 co-primary endpoints - highly statistically significant

Measured reduction in Gravimetric Sweat Production (GSP) and HDSM-Ax-7 scale responses

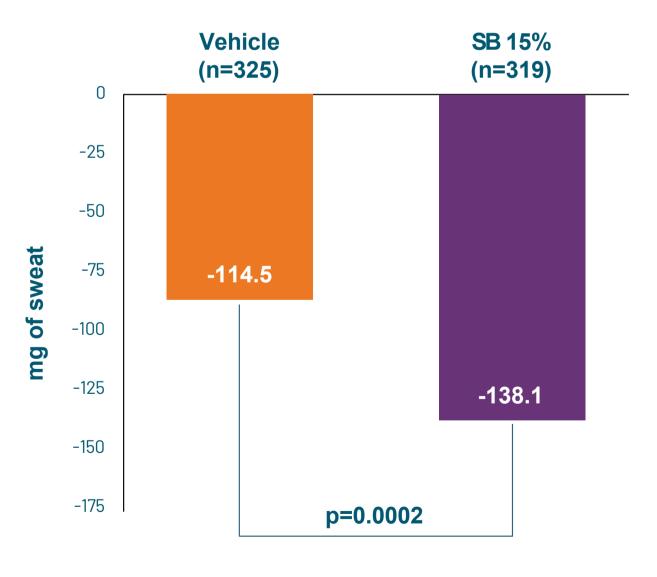
Pooled Data (Cardigan I and II)

≥2-point improvement in HDSM-Ax-7 from baseline to end of treatment¹



Pooled Data (Cardigan I and II)

GSP change from baseline to end of treatment¹

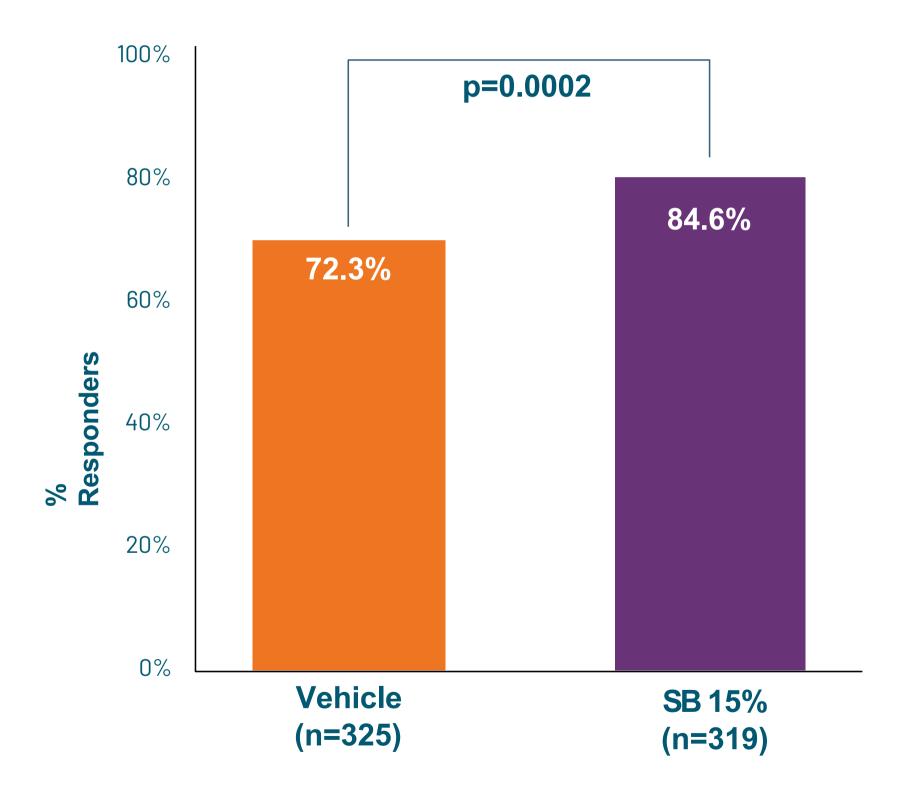


SB = Sofpironium Bromide

Secondary **Efficacy Endpoint:** Almost 85% of patients experienced a statistically significant and clinically meaningful response

Pooled Data (Cardigan I and II)

HDSM-Ax-7 reduction (≥1-point improvement) from baseline to end of treatment¹

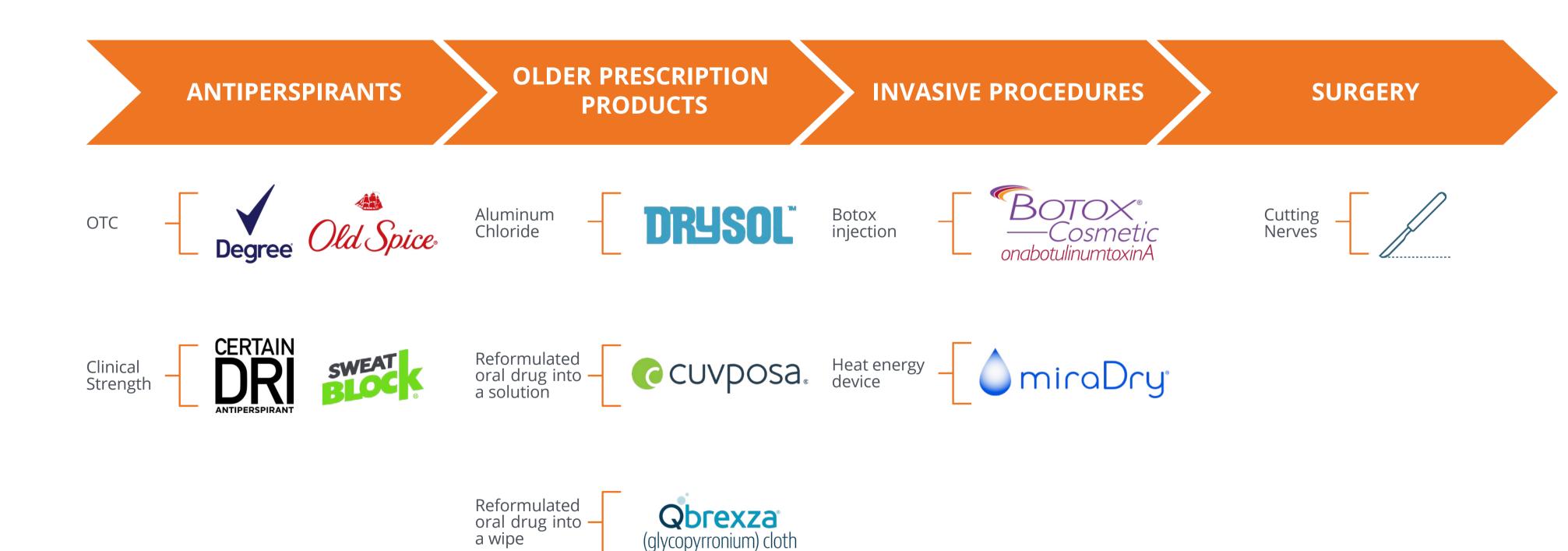


SB = Sofpironium Bromide

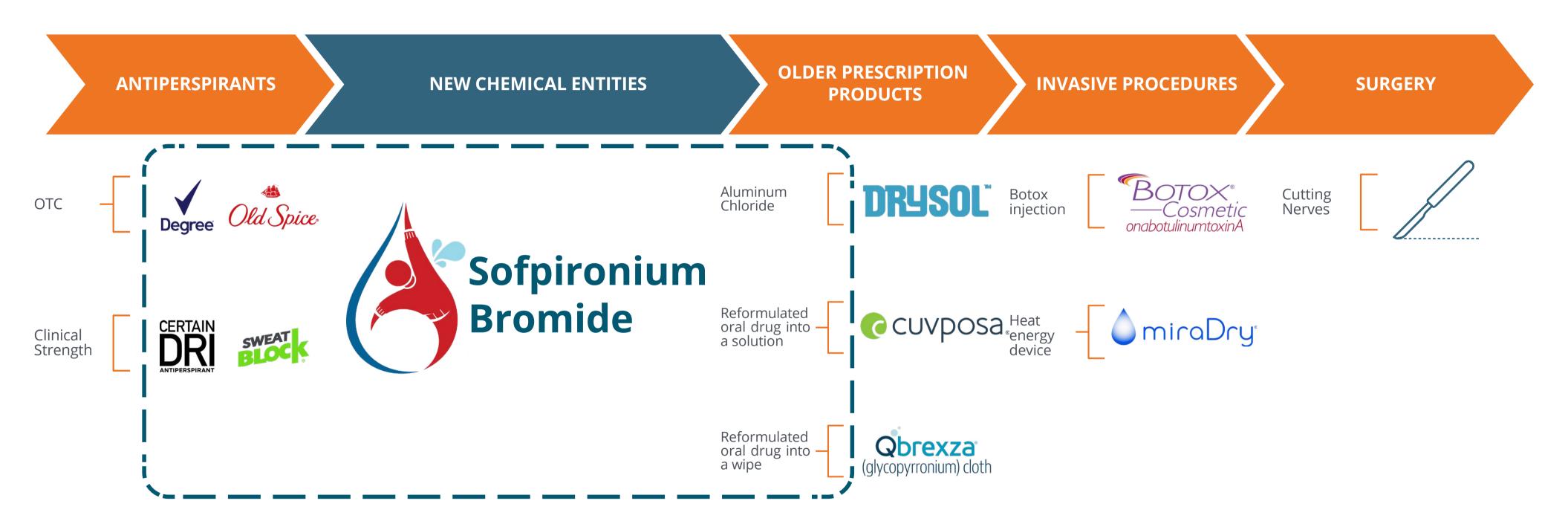
Source: 1. Data on File

Hyperhidrosis treatment continuum

No new chemical entities have been approved for hyperhidrosis



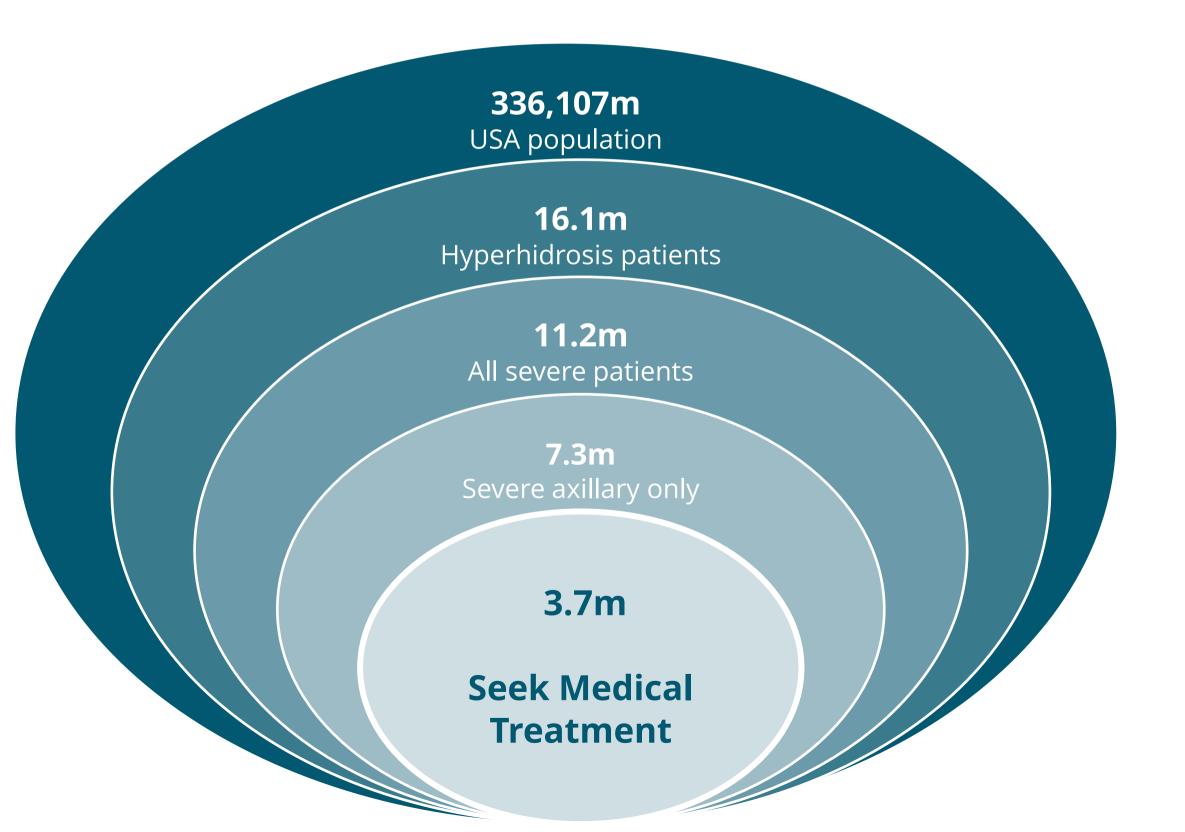
Significant opportunity for a new topical agent with class leading efficacy and safety



Due to its significant psychological impact, 54% of respondents suffering from hyperhidrosis say that they would <u>pay anything</u> for a treatment to stop their excessive sweating¹

Market opportunity for hyperhidrosis¹

Even a modest market share provides a significant financial opportunity



Share of patients already seeking treatment	Patients	Potential gross sales*
0.5%	18,500	\$144,300,000
1.0%	36,700	\$288,600,000
1.5%	55,500	\$432,900,000
2.0%	74,000	\$577,200,000
2.5%	92,500	\$721,500,000
3.0%	111,000	\$865,800,000

^{*} Current yearly cost of topical treatment is ~US\$7,800

Executing on key commercial and clinical milestones

Calendar year

Sofpironium Bromide

FDA filing for approval in 3Q 2022

Rosacea BTX 1702

Enrolment complete, data 3Q 2022 **Dermatitis BTX 1204A**

Enrolment complete, data 3Q 2022 Antimicrobial BTX 1801

Phase 2 study preparing to launch 2H 2022

Acne BTX 1503

Study start pending completion of BTX 1702 study

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