

16 October 2025

## Botanix Webinar – Monday 20 October – Registration Information

**Philadelphia PA and Phoenix AZ 16 October 2025:** Commercial dermatology company, Botanix Pharmaceuticals Limited (ABN 70 009 109 755) (ASX:BOT, “Botanix” or “the Company”), announces that it will host a webinar to provide a comprehensive update on the Company’s Quarterly Activity Report and 4C Quarterly Cash Flow Report. Executive Chairman Vince Ippolito, Chief Executive Officer Dr Howie McKibbin and US Chief Financial Officer Chris Lesovitz will include an update on the launch momentum for *Sofdra*® (sofpironium) topical gel, 12.45%.

The webinar will be held on Monday, 20 October 2025, at 11.00 am AEDT (Sydney/Melbourne)/8.00 am AWST (Perth). Interested participants **must register before the webinar using the link below**. Dial-in details will be sent in return.

### Webinar Details

**Date:** 20 October 2025

**Time:** 11.00 am AEDT (Sydney/Melbourne), 8.00 am AWST (Perth)

**To register:** [https://us06web.zoom.us/webinar/register/WN\\_-oGtp6viSHG5NwrKmitAsw](https://us06web.zoom.us/webinar/register/WN_-oGtp6viSHG5NwrKmitAsw)

**Dial in details:** Will be sent to you directly upon registration

Release authorised by the Botanix Board.

### About Botanix Pharmaceuticals

Botanix Pharmaceuticals Limited (ASX:BOT) is a dermatology company based in Philadelphia and Phoenix (US) which has received FDA approval for its lead product *Sofdra* for the treatment of primary axillary hyperhidrosis. *Sofdra* is the first and only new chemical entity approved by FDA to treat primary axillary hyperhidrosis and presents a novel safe and effective solution for patients who have lacked treatment options for this socially challenging medical condition.

To learn more please visit: <http://www.botanixpharma.com/>

**For more information, please contact:**

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**Cautionary Note on Forward-Looking Statements**

Any statements in this press release about future expectations, plans and prospects for the Company, the Company's strategy, future operations, 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 ability to successfully develop its product candidates and timely complete its planned clinical programs, the Company's ability to obtain marketing approvals for its 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.

## ***Sofdra* Important Safety Information & Indication**

### **Indication**

*Sofdra* (sofipirionium) topical gel, 12.45% is a prescription anticholinergic medicine used on the skin (topical) to treat excessive underarm sweating (primary axillary hyperhidrosis) in adults and children 9 years of age and older.

### **IMPORTANT SAFETY INFORMATION**

***Sofdra* is for use on the skin in the underarm area only. Wash your hands right away after you apply *Sofdra*. Do not touch your underarms after applying *Sofdra*. *Sofdra* is flammable. Avoid heat and flame while applying *Sofdra*.**

### **Who should not use *Sofdra*?**

Do not use *Sofdra* if you have certain medical conditions that can be made worse by taking an anticholinergic medicine such as glaucoma, severe ulcerative colitis (UC) or certain other serious bowel problems associated with severe UC, myasthenia gravis, and Sjogren's syndrome.

### **What should I tell my healthcare provider before using *Sofdra*?**

- **Tell your healthcare provider about all of your medical conditions**, including bladder or kidney problems, problems passing urine, if you are pregnant or breastfeeding, or plan to become pregnant or breastfeed. It is not known if *Sofdra* will harm your unborn baby or pass into your breast milk.
- **Tell your healthcare provider about all the medicines you take**, including prescription and over-the-counter medicines, especially any anticholinergic medicines.

### **What are possible side effects of *Sofdra*?**

#### **Serious side effects may include:**

- **Blurred vision.** Stop using *Sofdra*, call your healthcare provider right away, and do not drive or operate machinery or do hazardous work until your vision is clear.
- **New or worsened urinary retention.** Stop using *Sofdra* and call your healthcare provider right away if you experience difficulty urinating, urinating frequently, urination in a weak stream or drips, full bladder or difficulty emptying your bladder.

**The most common side effects of *Sofdra* include** dry mouth; blurred vision; pain, redness, swelling, itching, and irritation in the underarm area; dilation of the pupils of your eyes (mydriasis); and problems with urination. These are not all of the possible side effects of *Sofdra*. Call your doctor for medical advice about side effects.

You are encouraged to report negative side effects of prescription drugs to the FDA. Visit [www.fda.gov/medwatch](http://www.fda.gov/medwatch) or call 1-800-FDA-1088. You may also report side effects to Botanix at 1-866-763-6337.

**Keep *Sofdra* and all medicines out of the reach of children.**