



12 September 2024

AGM Date and Closing Date for Director Nominations



ANNUAL GENERAL MEETING (AGM) TO BE HELD ON THURSDAY, 14 NOVEMBER 2024

Zelira Therapeutics Ltd (ASX:ZLD, OTCQB:ZLDAF), a global leader in the research, development and commercialisation of clinically validated cannabinoid medicines, wishes to advise that, in accordance with ASX Listing Rule 3.13.1, it intends to hold its Annual General Meeting (AGM) on Thursday, 14 November 2024.

An item of business on the AGM agenda will be the re-election of Directors. The closing date for the receipt of nominations from persons wishing to be considered for election as a director is Thursday, 26 September 2024.

Any nominations must be received at the Company's registered office no later than 5.00pm (Perth time) on Thursday, 26 September 2024.

This announcement has been approved and authorised for release by the board of Zelira Therapeutics Limited.

For further information
please contact

Company

Dr Oludare Odumosu

Managing Director & CEO

☎ +1 909 855 0675

✉ oodumosu@zeliratx.com

Australia

Level 3, 101 St Georges Terrace

Perth WA 6000, AUSTRALIA

☎ +61 8 6558 0886

Fax: +61 8 6316 3337

✉ enquiries@zeliratx.com

www.zeliratx.com

ACN 103 782 378

Investors

Heidi Lord

Executive Director, Automic Group

☎ +61 404 216 403

✉ heidi.lord@automicgroup.com.au

USA

5110 Campus Drive, Suite 150

Plymouth Meeting, PA 19462

United States Of America

☎ +1 484 630 0650

Zelira Therapeutics Ltd (ASX:ZLD,

OTCQB:ZLDAF) Zelira is a leading global biopharmaceutical company in the research, development and commercialisation of clinically validated cannabinoid-based medicines. Zelira owns a portfolio of proprietary revenue generating products and a pipeline of candidates undergoing clinical development positioned to enter global markets. The Company is focused on developing and clinically validating branded cannabinoid-based medicines in its prescription [Rx] business for the treatment of a variety of medical conditions including insomnia, autism and chronic noncancer pain as well as offering over the counter [OTC] products.

Zelira has established a special purpose vehicle (SPV) to conduct FDA Phase 1, Phase 2 and Phase 3 clinical trials for Zelira's proprietary and patent protected HOPE® 1. Zelira has contributed to the SPV its HOPE® 1 product, IP and real-world data for 55% equity ownership of the SPV. Cash investors will contribute a total of circa US\$35 million to fund the SPV and US FDA trials for HOPE® 1 in exchange for a cumulative equity interest of 45% of the SPV. Zelira will manage the SPV as part of its business platform. The SPV has appointed iGENū CRO Pty Ltd (iGENū) as its Contract Research Organisation (CRO) to lead the clinical validation and regulatory registration of the study product with the US FDA through the submission of an Investigative New Drug (IND) application.

In May 2023, Zelira completed an IRB approved strategically designed multi-arm, head-to-head study targeting diabetic nerve pain. The clinical trial included a comprehensive comparison against the widely recognised and highly successful multi-

billion dollar revenue generating drug Lyrica® (Pregabalin). With the findings underscoring the exceptional efficacy of our treatments in managing pain, with ZLT-L-007 demonstrating the most substantial reduction in pain severity, particularly at the 60-day and 90-day follow-up periods. Zelira has developed Enhanced Distillate Capture and Dissolution Matrix (EDCDM) technology under the brand name Zyraydi™, that solves the problem of non-uniformity and separation of cannabinoid from powder bed, opening new ways to develop pharmaceutical grade solid oral dosage forms such as capsules and tablets. Zelira will be assessing opportunities for commercialisation of this technology.

Zelira's Rx business generates revenue from its proprietary medication, HOPE. The Company has two proprietary formulations under the HOPE® brand that are generating revenue in Australia, Washington, D.C., Pennsylvania and Louisiana. Zelira will also be expanding commercialisation of ZENIVOL® – the world's first clinically validated cannabinoid drug for treatment of chronic insomnia into Germany via its German commercialisation partner Adjupharm GmbH following recent approval from German regulatory authority BfArM. Zelira's OTC products in the oral and dermatology health care sectors are also generating revenue. Zelira, in partnership with SprinJeneCBD, launched a full line of oral care products, currently generating revenue in the US. Zelira also launched in 2021 the RAF FIVE™ brand, which consists of five OTC acne treatment products using a proprietary formulation incorporating cannabidiol (CBD).

For further information, please visit: zeliratx.com

