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TheraCryf plc
("TheraCryf", the "Company" or the "Group")

Lead Ox-1 Programme Advances to Final Toxicology Studies Before Human Phase 1 Clinical Study

*Ox-1 blocker well tolerated at the highest regulatory dose permissible
Programme remains on track for clinic readiness by Q4 2026*

TheraCryf plc (AIM: TCF), the biotech company developing new medicines for addiction and other neuropsychiatric disorders, today announces completion of preclinical dose range finding studies for its lead orexin-1 (Ox-1) receptor antagonist programme being developed for addiction, a market worth US 42 billion¹ and an area attracting significant clinical and commercial interest from large pharmaceutical companies.

Dose range finding is an essential step in moving a drug into the clinic that identifies the doses for the pivotal 28-day toxicology study. TheraCryf's Ox-1 blocker was well tolerated at doses up to 1g per 1kg of body weight, the highest that is permissible by regulators in such a study. Findings from this study are consistent with data reported for other Ox-1 and dual Ox-1/2 antagonists that have achieved full marketing approval.

Based on these results, doses have been selected for the pivotal 28-day toxicology studies, the final major pre-clinical studies required for submission for regulatory approval for the first in human clinical study. These studies will commence imminently, with final reporting on schedule in Q3 2026.

Dr Huw Jones, Chief Executive Officer of TheraCryf, commented:

"We continue to deliver a potentially class leading asset on target for clinic readiness in the fourth quarter of this year, exactly in line with the plan outlined at the start of this project. The substance use disorder market is already worth over US 42 billion growing to over US 71 billion¹ over the next seven years.

These conditions cause more deaths than road accidents and substance abusers lose over 20 years of life compared to non-abusers². The licensing market to large pharma for such treatments is considerable, generating substantial returns to originating companies at the right stage of development. With a class leading profile, we anticipate generating significant commercial interest in TheraCryf's Ox-1 asset."

1. Substance Use Disorder Treatment Market Size and Share Forecast Outlook 2025 to 2035. Future Market Insights Inc, November 2025

<https://www.futuremarketinsights.com/reports/substance-use-disorder-treatment-market>

2. Gov.uk and Chan *et al* Lancet 2023

<https://www.gov.uk/government/statistics/substance-misuse-treatment-for-adults-statistics-2023-to-2024/adult-substance-misuse-treatment-statistics-2023-to-2024-report>

[https://www.thelancet.com/journals/lanpub/article/PIIS2468-2667\(25\)00174-4/fulltext#:~:text=Variations%20in%20estimates%20of%20alcohol,impact%20policies%20must%20be%20intensified](https://www.thelancet.com/journals/lanpub/article/PIIS2468-2667(25)00174-4/fulltext#:~:text=Variations%20in%20estimates%20of%20alcohol,impact%20policies%20must%20be%20intensified)

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About TheraCryf

TheraCryf plc is a biotechnology company developing new medicines for addiction and other neuropsychiatric disorders, areas of significant unmet medical need within central nervous system (CNS) disorders.

The Group's lead programme is a novel, best-in-class orexin-1 receptor antagonist being developed as a potential treatment for addiction, including binge eating, alcohol and other substance use disorders.

The programme has already been heavily de-risked for both safety/tolerability and efficacy in previous testing and is fully funded through final pre-clinical trials to clinical readiness, with regulatory submissions for first in man studies targeted for 2026.

TheraCryf also has a dopamine transporter (DAT) modulator programme addressing fatigue of brain origin, including fatigue associated with multiple sclerosis, chemotherapy and narcolepsy. The Group also has a legacy, grant-funded, oncology programme in glioblastoma with SFX-01.

The Group operates a capital-light, virtual development model advancing programmes to early clinical or proof-of-concept stage before partnering with larger pharmaceutical or biotechnology companies.

TheraCryf's headquarters and registered office are at Alderley Park, Cheshire.

For further information, visit: <https://theracryf.com>

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