

# Lipocine to Present Clinical Data on LPCN 1148 at The Liver Meeting® 2023

"Late Breaking" abstract scheduled for oral presentation on November 13

SALT LAKE CITY, Nov. 1, 2023 /PRNewswire/ -- Lipocine Inc. (NASDAQ: LPCN), a biopharmaceutical company focused on treating Central Nervous System (CNS) disorders, today announced that an abstract on LPCN 1148 has been selected for a late breaking oral presentation at [The American Association for the Study of Liver Diseases \(AASLD\) - The Liver Meeting® 2023](#), to take place in Boston MA, November 10 to 14, 2023.

## Presentation Details

Title	<b>Oral LPCN 1148 improves sarcopenia and hepatic encephalopathy in patients with cirrhosis</b>
Presenter:	Dr. Arun J. Sanyal, MD, Director, Stravitz-Sanyal Institute for Liver Disease and Metabolic Health, Virginia Commonwealth University Advisor to Lipocine
Session Title:	Late Breaking Abstract #1
Session Date and Time:	Monday, November 13, 2023, 2:00 PM - 3:30 PM ET
Presentation Time:	2:45 PM ET
Abstract Number:	5003

## **About LPCN 1148**

Lipocine is currently evaluating LPCN 1148 comprising testosterone laurate ("TL") for the management of decompensated cirrhosis. The Company believes LPCN 1148 targets unmet needs for patients with advanced cirrhosis including improvement in the quality of life of patients while on the liver transplant waitlist, prevention or reduction in the occurrence of new decompensation events such as hepatic encephalopathy ("HE"), and improvement in post liver transplant outcomes including survival and costs.

## **About Lipocine**

Lipocine is a biopharmaceutical company leveraging its proprietary technology platform to augment therapeutics through effective oral delivery to develop differentiated products for CNS disorders. Lipocine has drug candidates in development as well as drug candidates for which we are exploring partnering. Our drug candidates represent enablement of differentiated, patient friendly oral delivery options for favorable benefit to risk profile which target large addressable markets with significant unmet medical needs.

Lipocine's clinical development candidates include: LPCN 1154, oral brexanolone, for the potential treatment of postpartum depression, LPCN 2101 for the potential treatment of epilepsy and LPCN 1148, a novel androgen receptor agonist prodrug for oral administration targeted for the management of symptoms associated with liver cirrhosis. Lipocine is exploring partnering opportunities for LPCN 1107, our candidate for prevention of preterm birth, LPCN1154, for rapid relief of postpartum depression, LPCN 1148, for the management of decompensated cirrhosis, LPCN 1144, our candidate for treatment of non-cirrhotic NASH, and LPCN 1111, a once-a-day therapy candidate for testosterone replacement therapy (TRT). TLANDO, a novel oral prodrug of testosterone containing testosterone undecanoate developed by Lipocine, is approved by the FDA for conditions associated with a deficiency of endogenous testosterone, also known as hypogonadism, in adult males. For more information, please visit [www.lipocine.com](http://www.lipocine.com).

## **Forward-Looking Statements**

This release contains "forward-looking statements" that are made pursuant to the safe harbor provisions of the Private Securities Litigation Reform Act of 1995 and include statements that are not historical facts regarding our product development efforts, the application of our proprietary platform in developing new treatments for CNS disorders, our product candidates and related clinical trials, our development of and filing of a NDA with the FDA for LPCN 1148, and the potential uses and benefits of our product candidates. Investors are cautioned that all such forward-looking statements involve risks and uncertainties, including, without limitation, the risks that we may not be successful in developing product candidates to treat CNS disorders, we may not have sufficient

capital to complete the development processes for our product candidates, we may not be able to enter into partnerships or other strategic relationships to monetize our non-core assets, the FDA will not approve any of our products, risks related to our products, expected product benefits not being realized, clinical and regulatory expectations and plans not being realized, new regulatory developments and requirements, risks related to the FDA approval process including the receipt of regulatory approvals and our ability to utilize a streamlined approval pathway for LPCN 1154, the results and timing of clinical trials, patient acceptance of Lipocine's products, the manufacturing and commercialization of Lipocine's products, and other risks detailed in Lipocine's filings with the SEC, including, without limitation, its Form 10-K and other reports on Forms 8-K and 10-Q, all of which can be obtained on the SEC website at [www.sec.gov](http://www.sec.gov). Lipocine assumes no obligation to update or revise publicly any forward-looking statements contained in this release, except as required by law.

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