## Lipocine to Present at 36th Annual Canaccord Genuity Growth Conference

SALT LAKE CITY, Aug. 04, 2016 (GLOBE NEWSWIRE) -- <u>Lipocine Inc.</u> (NASDAQ:LPCN), a specialty pharmaceutical company, today announced that Mahesh Patel, President and CEO, and Morgan Brown, Executive Vice President and CFO, will present a company overview at the 36<sup>th</sup> Annual Canaccord Genuity Growth Conference on Thursday, August 11, 2016 at 9:30 a.m. ET.

A live audio webcast of the presentation will be available via the "Investor Relations" page of the Lipocine website, <a href="www.lipocine.com">www.lipocine.com</a>. Please log on through Lipocine's website approximately 10 minutes prior to the scheduled start time. A replay of the webcast will also be archived on Lipocine's website for 90 days following the presentation.

## **About Lipocine**

Lipocine Inc. is a specialty pharmaceutical company developing innovative pharmaceutical products for use in men's and women's health using its proprietary drug delivery technologies. LPCN 1021, an oral testosterone replacement therapy product candidate, demonstrated positive efficacy and safety results in Phase 3 testing but received a Complete Response Letter from the FDA on June 28, 2016. LPCN 1111, a next-generation oral testosterone replacement therapy product with once-daily dosing, is currently in Phase 2 testing. LPCN 1107, which has the potential to become the first oral hydroxyprogesterone caproate product indicated for the prevention of recurrent preterm birth, has an End of Phase 2 meeting scheduled with the FDA during the third quarter of 2016 and has been granted orphan drug designation by the FDA. For more information, please visit <a href="https://www.lipocine.com">www.lipocine.com</a>.

## **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 relating to Lipocine's FDA review process relating to LPCN 1021 and the possible outcome of such process, clinical trials, the regulatory process for our other product candidates, the potential uses and benefits of our product candidates, the TRT market, product development and commercialization efforts and the projected timing and outcome of regulatory filings and actions. Investors are cautioned that all such forward-looking statements involve risks and uncertainties, including, without limitation, the risks related to our products, expected product benefits, clinical and regulatory expectations and plans, regulatory developments and requirements, risks related to the receipt of a CRL from the FDA for LPCN 1021, the receipt of regulatory approvals, the results of clinical trials, patient acceptance of Lipocine's products, the manufacturing and commercialization of Lipocine's products, the risks related to market conditions for Lipocine's common stock 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 <a href="https://www.sec.gov">www.sec.gov</a>. 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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