

# U.S. FDA Extends Review for TLANDO™; Advisory Committee Meeting Date of January 10, 2018 Remains Unchanged

SALT LAKE CITY, Nov. 17, 2017 (GLOBE NEWSWIRE) -- [Lipocine Inc.](#) (NASDAQ:LPCN), a specialty pharmaceutical company, today announced the U.S. Food and Drug Administration ("FDA") has extended the review period for the New Drug Application ("NDA") for TLANDO, the Company's oral testosterone product candidate for the proposed indication of testosterone replacement therapy ("TRT") in adult males for conditions associated with a deficiency of endogenous testosterone, also known as hypogonadism. The FDA has assigned a new Prescription Drug User Fee Act ("PDUFA") goal date of May 8, 2018.

The FDA extended the action goal date to allow time to review additional data analyses recently submitted as an amendment to the NDA in response to the FDA's Information Requests. Upon preliminary review of the amendment by the FDA, the submission of the requested information has been determined to represent a Major Amendment to the NDA resulting in an extension of the PDUFA goal date by three months to provide time for a full review.

The previously scheduled Bone, Reproductive and Urologic Drugs Advisory Committee ("BRUDAC") meeting on January 10, 2018 to discuss the NDA for TLANDO remains unchanged.

"We believe TLANDO addresses a significant unmet medical need for men with hypogonadism and we will continue to work closely with the FDA throughout the review process," said Dr. Mahesh Patel, Chairman, President and Chief Executive Officer of Lipocine.

## 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. Lipocine's clinical development pipeline includes three development programs TLANDO, LPCN 1111 and LPCN 1107. TLANDO, a novel oral prodrug of testosterone containing testosterone undecanoate, is designed to help restore normal testosterone levels in hypogonadal men. TLANDO was well tolerated and met the primary efficacy end-points in Phase 3 testing with twice daily dosing and is currently under FDA review. LPCN 1111, a novel oral prodrug of testosterone, originated and is being developed by Lipocine as a next-generation oral testosterone product with potential for once-daily dosing and is currently in Phase 2 testing. LPCN 1107 is potentially the first oral hydroxyprogesterone caproate product candidate indicated for the prevention of recurrent preterm birth and has been granted orphan drug designation by the FDA. An End of Phase 2 meeting with the FDA has been completed. 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 Lipocine's product candidates and related clinical trials and the FDA review process relating to its product candidates, the expected timing of the FDA review process related to our resubmitted NDA, the path to approvability by the FDA of Lipocine's development programs, the potential uses and benefits of our product candidates, and our product development efforts. Investors are cautioned that all such forward-looking statements involve risks and uncertainties, including, without limitation, the risks that the FDA will not approve any of our products, the risk that Bone, Reproductive, Urologic and Drug Advisory Committee ("BRUDAC") may make a negative recommendation to the Commissioner of the FDA with respect to TLANDO, 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 that the FDA will determine there are deficiencies in our resubmitted NDA, the receipt of regulatory approvals, 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.

## CONTACT:

Morgan Brown  
Executive Vice President & Chief Financial Officer  
Phone: (801) 994-7383  
[mb@lipocine.com](mailto:mb@lipocine.com)

Investors:  
Hans Vitzhum  
Phone: (646) 597-6979  
[hans@lifesciadvisors.com](mailto:hans@lifesciadvisors.com)

---

<https://ir.lipocine.com/U-S-FDA-Extends-Review-for-TLANDO-TM-Advisory-Committee-Meeting-Date-of-January-10-2018-Remains-Unchanged>