

SPARC/Sec/SE/2019-20/014

1st July 2019

To

National Stock Exchange of India Ltd.

Exchange Plaza,
Plot No. C/1, G Block,
Bandra Kurla Complex,
Bandra (East),
Mumbai – 400 051.

BSE Limited

P J Towers,
Dalal street,
Mumbai - 400001

Ref: Scrip Code: NSE: SPARC; BSE: 532872

Sub: Press Release: SPARC Announces U.S. FDA Acceptance of NDA for Taclantis™
(Paclitaxel Injection Concentrate for Suspension) for Filing and Regulatory Review

Dear Sir/Madam,

Pursuant to regulation 30 of the SEBI (Listing Obligation & Disclosure Requirements) Regulations, 2015, we enclose herewith the Press Release on the above mentioned subject being released by the Company, which is self-explanatory.

We request you to kindly take the same on record.

Yours faithfully,

For **Sun Pharma Advanced Research Company Limited**

A handwritten signature in black ink, appearing to read "Debashis Dey".

Debashis Dey
Company Secretary

Encls: A/a.



FOR IMMEDIATE RELEASE

SPARC Announces U.S. FDA Acceptance of NDA for Taclantis™ (Paclitaxel Injection Concentrate for Suspension) for Filing and Regulatory Review

MUMBAI – July 01, 2019, Sun Pharma Advanced Research Company Ltd. (SPARC) (Reuters: SPRC.BO, Bloomberg: SPADV IN, NSE: SPARC, BSE: 532872) today announced that the U.S. Food and Drug Administration (USFDA) has accepted for review SPARC's New Drug Application (NDA) for Taclantis™ (Paclitaxel Injection Concentrate for Suspension). The NDA filing is based on successful demonstration of clinical bioequivalence of Taclantis™ with Abraxane® and associated clinical safety data. SPARC seeks the same indications as Abraxane® for Taclantis™ in the NDA. The USFDA confirmed that this NDA will be a standard review.

About Taclantis™ (Paclitaxel Injection Concentrate for Suspension):

Taclantis™ (Paclitaxel Injection Concentrate for Suspension) is a Cremophor® and Albumin-free formulation of Paclitaxel. It should be diluted with an appropriate volume of 5% w/v Dextrose injection in either a PVC or non-PVC type sterile infusion bag. Premedication to prevent hypersensitivity is generally not needed prior to administration of Taclantis™.

About SPARC (CIN: L73100GJ2006PLC047837):

Sun Pharma Advanced Research Company Ltd. (SPARC) is a global pharmaceutical company focused on continuously improving standards of care for patients globally through innovation in therapeutics and delivery. SPARC aims to consistently lower costs and improve operational efficiencies to advance availability and affordability of cures for patients across the world. More information about the company can be found at www.sparc.life

Disclaimer:

Statements in this document describing the Company's objectives, projections, estimates, expectations, plans or predictions or industry conditions or events may be "forward looking statements" within the meaning of applicable securities laws and regulations. Actual results, performance or achievements could differ materially from those expressed or implied.

Media Contacts:

Jaydeep Issrani
Tel +91 22 66455645, Extn: 5787
Tel Direct +91 22 6645 5787
Mobile +91-9820216916
E mail jaydeep.issrani@sparcmail.com

*Taclantis™ is conditionally approved by USFDA as trade name for US market.
®, TM - All brand names and trademarks are the property of respective owners.*

Sun Pharma Advanced Research Company Ltd.

17/B, Mahal Industrial Estate, Mahakali Caves Road, Andheri (East), Mumbai 400 093, Maharashtra, India.
Tel.: (91-22) 6645 5645 | Fax.: (91-22) 6645 5685 | CIN: L73100GJ2006PLC047837 | Website: www.sparc.life

Registered Office : SPARC, Akota Road, Akota, Vadodara - 390 020, Gujarat, India.