

SUN PHARMA
ADVANCED RESEARCH
COMPANY LTD.

FOR IMMEDIATE RELEASE

SPARC receives US FDA response on Levetiracetam NDA

May 4, 2013, Mumbai: Sun Pharma Advanced Research Company Ltd. (SPARC, NSE: SPARC, BSE: 532872) announced receipt of a Complete Response Letter from the U.S. Food and Drug Administration (FDA) on its New Drug Application (NDA) for Levetiracetam Extended-release Tablets, 1000mg and 1500mg, an anti-epileptic product. A Complete Response Letter is a communication from the FDA to companies that an NDA cannot be approved in its present form.

In the Complete Response Letter, the FDA specified that the clinical data submitted by SPARC establishes bioequivalence in the fasted state. However, the FDA has raised certain queries on the Pharmacokinetic data in the fed state. SPARC is evaluating the contents of the letter and plans further discussions with the FDA.

About SPARC

Sun Pharma Advanced Research Company Ltd is an international pharmaceutical company engaged in research and development of drugs and delivery systems. More Information about the company can be found at www.sunpharma.in.

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For the purpose of this letter,
the following information is
being provided for your
reference.

SUN PHARMA
ADVANCED RESEARCH
COMPANY LTD.

4 May 2013.

National Stock Exchange of India Ltd,
Exchange Plaza, 5th Floor,
Plot No C/1, G Block,
Bandra Kurla Complex,
Bandra (East),
Mumbai - 400 051.

BSE Limited.,
Market Operations Dept.
P. J. Towers,
Dalal Street,
Mumbai - 400 001.

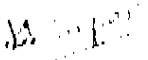
Dear Sirs,

Sub: Press Release

We are pleased to enclose herewith our Press Release relating to update on New Drug Application (NDA) for Levetiracetam Extended-release Tablets which we shall be releasing after faxing this letter to you. This is for your information and record.

Thanking you,

Yours faithfully,
For Sun Pharma Advanced Research Company Ltd.


Meetal Sampat
Company Secretary