

10th April, 2017

To, **National Stock Exchange of India Limited** Exchange Plaza Bandra Kurla Complex Bandra (E), Mumbai – 400051

Scrip Code: MARKSANS

To. **BSE Limited** P. J. Towers Dalal Street Mumbai - 400 001

Scrip Code: **524404**

Dear Sirs,

We wish to inform you that the United States Food and Drug Administration (US FDA) inspected our manufacturing facility located at L-82 & L-83, Verna Industrial Estate, Verna, Goa – 403722 from 3rd to 7th April, 2017.

At the end of the inspection, there were 4 (four) observations given under Form 483. The Company has already started addressing the same and is confident of satisfying the FDA within the stipulated time.

Further, on 7th April, 2017, US FDA has also granted final approval of the Abbreviated New Drug Application for Dutasteride Soft Gelatin Capsules 0.5 mg, which is therapeutically equivalent to the reference listed drug (RLD), Avodart® Capsules, 0.5 mg of GlaxoSmithKline. Dutasteride is indicated for the treatment of symptomatic benign prostatic hyperplasia (BPH) in men with an enlarged prostate. Marksans will launch the product immediately.

This is for your information and records.

Thanking you.

Yours faithfully,

For Marksans Pharma Limited

Harshavardhan Panigrahi

Company Secretary & Manager - Legal

CIN: L24110MH1992PLC066364

www.marksanspharma.com