November 25, 2022

The Manager- Listing
BSE Limited
Corporate Relationship Dept., 5th Floor, New Trading Ring
Rotunda Building, P J Towers, Dalal Street, Fort,
Mumbai - 400001

The Manager- Listing
National Stock Exchange of India Limited
Exchange plaza, 5th Floor, Plot No. C/1, G Block
Bandra –Kurla Complex, Bandra (E),
Mumbai - 400051

Dear Sir,

Sub: Intimation under Regulation 30 of SEBI (Listing Obligations and Disclosure Requirements) Regulation, 2015

This is to inform that AstraZeneca Pharma India Limited has received an Import and Market permission in Form CT-20 (Subsequent New Drug Approval) from the Central Drugs Standard Control Organisation, Directorate General of Health Services, Government of India for Dapagliflozin (Forxiga®) tablets of 10 mg - additional indication.

Through this approval, the Company is permitted to deal in Dapagliflozin (Forxiga®) tablets of 10 mg in additional/expanded indication "to reduce the risk of sustained eGFR and kidney disease, cardiovascular death and hospitalization for heart failure in adult with chronic kidney disease at risk of progression" **with condition that:-** it is indicated in adults for the treatment of patients of chronic kidney disease (CKD) up to eGFR of greater than or equal to 25ml/min/1.73m2. Below this, initiation of the treatment is not recommended however the patients may continue 10mg orally once daily to reduce the risk of eGFR decline, ESKD, CV death and hHF.

The receipt of this permission paves way for the launch of Dapagliflozin (Forxiga®) tablets of 10 mg in India for the specified additional/expanded indication, subject to the receipt of related statutory approvals

We request you to take the above intimation on record.

Thanking you
Yours faithfully

For **AstraZeneca Pharma India Limited**

MANASA RAMA
Digitally signed by MANASA RAMA
Date: 2022.11.25 20:44:35 +05'30'

Manasa. R
Company Secretary

CORPORATE & REGD. OFFICE
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