



Date: 5th July 2023

To,
The Manager,
Listing Compliance Department,
National Stock Exchange of India Ltd,
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Symbol: SAKAR

SUB: WHO GMP approval received for Oncology Injection (small volume parenteral & lyophilized) unit at Sakar's vertically integrated oncology product manufacturing site

Reference: Announcement under Regulation 30 (LODR)

Dear All,

With great pleasure we would like to share that Sakar's research-driven API integrated oncology formulation manufacturing unit located at Bavla, Gujarat, India has grabbed WHO GMP approval from Food & Drugs Control Administration, Gujarat for its injection dosage form (small volume parenteral, liquid and lyophilised forms). The regulatory authority has earlier certified both the Oral Solid Dosage (OSD) and Active Pharmaceutical Ingredient (API) manufacturing units in cytotoxic category of Sakar with Good Manufacturing Practices (GMP).

With this certification, the entire oncology product-manufacturing site of Sakar is now WHO GMP approved and preparing to:

- Trigger audits by overseas regulatory bodies to export injection cytotoxic products in addition to oral solids and API in same category
- Complete dossiers with necessary Certificate of product permission (CoPP) for small volume parenteral, liquid and lyophilised injection range and apply for registering products overseas (Marketing Authorizations) for exports
- Offer to manufacture and supply the product mix in injection and oral solid category for renowned players in the segment at global level

With media filling, process validation, dossier compilation in progress with cytotoxic injections, Sakar is now heading towards export of anti-cancer products basket to multiple countries.

This is for your information, records and meeting the disclosure requirements under the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015, as amended from time to time.

Thanking you,

Yours faithfully,
for SAKAR HEALTHCARE LIMITED

BHARAT SONI
COMPANY SECRETARY &
COMPLIANCE OFFICER

