Zeal & Innovation in Medicine

Ref No.: ZLL/CS/BSE/NSE

Date: 01.08.2025

BSE Limited, Corporate Relationship Department P. J. Towers, Dalal Street, Mumbai- 400 001 Company Code- 541400	National Stock Exchange of India Limited Listing Compliance Department Exchange Plaza, Bandra-Kurla Complex, Bandra (E), Mumbai – 400 051 (Symbol - ZIMLAB)

Dear Sir/Madam,

Sub: EU GMP Inspection Outcome

Ref: Regulation 30 of the SEBI (Listing Obligations and Disclosure Requirements) Regulations 2015

This is to inform that the manufacturing facility of ZIM Laboratories Limited ("The Company"), located at B-21/22, MIDC Area, Kalmeshwar – 441 501, District Nagpur, Maharashtra, underwent an inspection conducted by the German and Portuguese regulatory authorities ("authorities"). The inspection was held from 30th June 2025 to 04th July 2025, as part of the European Good Manufacturing Practices (EU GMP) compliance process.

Following the inspection, the authorities have issued a report dated 31st July 2025 indicating that the facility is in non-compliance with EU-GMP requirements. The inspection identified 2 critical, 8 major, and 18 minor deficiencies.

The Company is required to submit a detailed and binding Corrective and Preventive Action (CAPA) plan within 4 weeks from the date of the report. A re-inspection will be conducted following the implementation of the CAPA to validate the resolution of the observed deficiencies.

The Company remains committed to the highest standards of quality and regulatory compliance across all its operations and will address the observations in a comprehensive and timely manner working closely with the regulatory authorities to achieve full compliance

We will continue to update stakeholders on the progress of this matter as appropriate.

Kindly take the intimation on record.

Thanking you,

Yours faithfully,
For ZIM LABORATORIES LIMITED

PIYUSH SIDDHESHWAR
NIKHADE

Digitally signed by PIYUSH SIDDHESHWAR NIKHADE
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(Piyush Nikhade)
Company Secretary and Compliance Officer
Membership No. A38972