

January 05, 2026

To,
The Manager,
Dept. of Corporate Services,
BSE Limited,
Phiroze Jeejeebhoy Towers,
Dalal Street, Mumbai- 400001

Dear Sir/Madam,

Subject : Receipt of certificate of GLP compliance from National Good Laboratory Practice (GLP) Compliance Monitoring Authority (NGCMA), Department of Science and Technology, Government of India – Regulation 30 of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015

Scrip Code : 511509

Pursuant to Regulation 30 read with Schedule III of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015, we wish to inform that Vivo Bio Tech Limited has received the Certificate of Good Laboratory Practice (GLP) Compliance from the National Good Laboratory Practice (GLP) Compliance Monitoring Authority (NGCMA), Department of Science & Technology, Government of India.

The GLP certification specifically covers a wide range of non-clinical safety and regulatory studies, with includes the following critical and high-value study areas in addition to the other existing area(s) of expertise;

- Toxicity Studies in Dogs, including repeated dose and specialized toxicological evaluations;
- Carcinogenicity Studies, supporting long-term safety assessment and global regulatory submissions;
- Eye Irritation / Serious Eye Damage Studies, including *in vitro* models, aligned with international regulatory recommendation to replace animal models.

The receipt of GLP accreditation for dog studies, carcinogenicity studies and *in vitro* eye studies significantly enhances the Company's capabilities to conduct globally acceptable non-clinical safety

studies across multiple product lines and strengthens its credibility and positioning as a comprehensive GLP-compliant research test facility supporting international regulatory filings as a part of product registration mandate.

Kindly take the above information on your record.

Thanking You,

For Vivo Bio Tech Limited

**A V Kiran
Company Secretary**