

2<sup>nd</sup> July 2026

National Stock Exchange of India Limited  
Exchange Plaza,  
Bandra Kurla Complex,  
Bandra (East), Mumbai – 400 051

BSE Limited  
Phiroze Jeejeebhoy Towers,  
Dalal Street,  
Mumbai – 400001

**Scrip Code: AJANTPHARM EQ**

**Scrip Code: AJANTPHARM 532331**

**Sub.: Update on US FDA inspection at Paithan manufacturing facility**

Dear Sir/Madam,

We are pleased to inform you that the Company has received the Establishment Inspection Report (EIR) from the United States Food and Drug Administration (USFDA) for its manufacturing facility at Paithan, Maharashtra. The inspection has been classified as Voluntary Action Indicated (VAI).

The current Good Manufacturing Practices (cGMP) inspection of the Paithan facility was conducted by the USFDA from 13 April 2026 to 21 April 2026. The Company had earlier intimated the stock exchanges regarding the inspection on 22 April 2026.

This is for your information and records.

Thanking you,

Yours faithfully,

**Gaurang Shah**

*Sr. VP – Legal & Company Secretary*