FREQUENTLY ASKED QUESTIONS (FAQs)
[Reference: TMR notification *vide* S.O. 5690(E) dated 17.10.2026]

S. No.	Question	Response
1	When will the IPDMS portal be live for submission of Annexure-I information?	Annexure-I is live on IPDMS Portal.
2	Are Form V and Annexure-I required to be submitted even when there is no change in MRP?	Manufacturers are required to submit information in Form-V and Annexure-I through IPDMS within seven days of publication of the Order, irrespective of whether there is change in MRP or not.
3	What is required to be mentioned in Newspaper advertisement?	The newspaper advertisement is required to be given for all products where there is reduction in prices. The advertisement must mention pre-revised MRP and revised MRP. The companies can choose to mention those products where there is no revision in MRP.
4	What constitutes Government sales for the purpose of this order?	Government sales shall mean the sales where: (i) MRP is not printed on the pack; Or (ii) MRP is printed with clear labelling as *"For supply to Government; not for sale"*.
5	Are sales returns to be adjusted while calculating net sales for PTS/PTD?	No adjustment on account of sales return is permissible.
6	Does TMR apply if a drug listed in Table B is used individually or in combination or for other therapeutic purposes?	Yes, as stated in note (o) of the Order, TMR for non-scheduled formulations shall be applicable for all strengths and dosage form (whether individual or in combination, irrespective of dosage strength, dosage form and /or route of administration) containing any of the drugs listed in Table B even if the formulation is being used for other therapeutic uses.
7	By when must the undertakings relating to the Patient Assistance Programme (PAP) be submitted to NPPA?	The undertakings mentioned in note (d)(ii) & (iii) of the Order relating to PAP has to be submitted initially by all manufacturers before 29.10.2026 and also every time whenever there is a revision in MRP. This information may be submitted on pricing-nppa@gov.in
8	Is recalling, re-labelling or re-stickering mandatory?	Recalling, re-labelling or re-stickering on the label of container or pack of stocks released in the market prior to implementation of revised MRP is not mandatory, if manufacturer is able to ensure price compliance at the retailer level.
9	When is the information required in note q(iii) & (iv) of the Order required to be submitted to NPPA?	At the request of industry associations, the information as specified in note q(iii) & (iv) may be submitted not later than 09.11.2026.

S. No.	Question	Response
		The information pertaining to note q(iv) may be provided for as on 31.10.2026. This information may be submitted on pricing-nppa@gov.in
10	When can a manufacturer avail the annual price increase under Paragraph 20(1) of DPCO, 2013 for drugs covered under the TMR Order?	As per the S.O. 5690(E) dated 17.10.2026, the revised prices become effective from 09.11.2026. Where there is a revision in MRP, the manufacturer shall be eligible to avail annual price increase under Paragraph 20(1) of DPCO, 2013 from the date of implementation of revised price which is not later than 09.11.2026. Where there is no revision in MRP, the manufacturer shall be eligible to avail annual price increase under Paragraph 20(1) of DPCO, 2013 after expiry of 12 months' period from last revision in MRP.