



Mallinckrodt™
Pharmaceuticals

Technical Bulletin

Ref: **TB-17002-ENG**

Product: INOmax DS_{IR}®, INOmax DS_{IR}® Plus,
INOmax DS_{IR}® Plus MRI and INOflo DS®

Affected parts: INOmax DS_{IR}, INOmax DS_{IR} Plus,
INOmax DS_{IR} Plus MRI and INOflo DS

Subject: **Purging the Regulator Supply Line and/or the Delivery System**

Issue date: February 2022

Priority: Low

Classification: Information for Distribution

This technical bulletin is written to advise clinicians that testing has been completed which demonstrates that the INOmax DS_{IR}, INOmax DS_{IR} Plus, INOmax DS_{IR} Plus MRI and INOflo DS may stand idle for 24 hours after a successful Pre-Use procedure and depressurization of the regulator(s) has been completed.

If the device is not placed into use within these 24 hours repeat the Pre-Use procedure.



WARNING: If the INOmax DS_{IR}, INOmax DS_{IR} Plus, INOmax DS_{IR} Plus MRI or INOflo DS is depressurized and is not used on a patient within 24 hours, repeat the pre-use procedure.

For technical assistance regarding any of the above devices, please contact Technical Support at 1-877-566-9466 (North America) or your specific country manager.

Mallinckrodt Manufacturing LLC
6603 Femrite Drive,
Madison, WI 53718-6801 USA

Mallinckrodt, the "M" brand mark and the Mallinckrodt Pharmaceuticals logo are trademark of a Mallinckrodt company. Other brands are trademarks of a Mallinckrodt company or their respective owner. ©2022 Mallinckrodt



Europe
European Authorized Representative
MDSS GmbH
Schiffgraben 41
30175 Hannover, Germany

