

Nebulised Epoprostenol (FLOLAN®)

DERBY ONLY

Indication	To improve oxygenation and V/Q mismatch caused by ARDS. Use should be on Consultant Intensivist instruction, if other management strategies have failed.
Dose	10 – 50 nanograms/Kg/min; usual starting rate of 30 nanograms/Kg/min. Titrate dose to effect; dose adjustments should be made at 15-minute intervals. See the dosing table on the second page of this monograph. Doses should be based on ideal body weight: <ul style="list-style-type: none"> • Males: Height (cm) – 100 • Females: Height (cm) – 105
Preparation	<p>Epoprostenol vials (Flolan®) contain 500 micrograms</p> <ol style="list-style-type: none"> 1. Using a BD syringe, draw up 10mL of the provided solvent and use this to reconstitute the vial containing the Flolan® (epoprostenol) powder, then shake gently until the powder is dissolved. 2. Draw up the resulting solution into a 50mL BD syringe. 3. Make up to 25mL using the remaining solvent, then mix thoroughly. 4. Transfer this solution from the BD syringe into the blue 'Aerogen® Solo Continuous Nebulisation Syringe'. <p>This gives a concentration of 20,000 nanograms/mL</p>
Administration	See the 'Inhaled Prostacyclin (Epoprostenol) - ICU Clinical Guideline' on Koha for full instructions. Use only with the Aerogen® Solo nebuliser. The drug must be administered via the blue Aerogen® syringe to avoid inadvertent intravenous administration.
Shelf-life	<p>Applies to the FLOLAN® product only:</p> <p>72h at room temperature, once reconstituted</p> <p>Reconstituted solution can be stored in the vial for up to 8 days at 2-8°C</p> <p>The unopened vials and reconstituted solution must be protected from light.</p>
Common Compatibility Issues	N/A

Additional information	<p>Epoprostenol should be discontinued if no beneficial clinical effect is seen after 15 minutes.</p> <p>Epoprostenol should not be stopped suddenly, to avoid rebound pulmonary hypertension – titrate dose down by 10nanograms/Kg/min every 2 hours before stopping.</p> <p>Flow rates >12mL/h may lead to accumulation of drug in the nebuliser reservoir (6mL volume).</p> <p>Filters will become saturated with glycine – change every 8h at flow rates ≤10 mL/h and every 4h at rates >10 mL/h</p>																				
Sample Label	<table border="1" style="width: 100%; border-collapse: collapse;"> <thead> <tr style="background-color: yellow;"> <th colspan="4" style="text-align: center;">DRUGS ADDED TO THIS INFUSION</th> </tr> </thead> <tbody> <tr> <td colspan="3"> PATIENT <i>A.patient (A.number)</i> </td> <td> Ward <i>ICU</i> </td> </tr> <tr> <td> Drug <i>Epoprostenol in 25mL diluent</i> </td> <td> Amount <i>500 micrograms (20,000 nanograms/mL)</i> </td> <td> Add By </td> <td> Checked By </td> </tr> <tr> <td> Date Added Time Added </td> <td> Exp. Date Exp. Time </td> <td colspan="2"> Batch No. </td> </tr> <tr style="background-color: yellow;"> <td colspan="4" style="text-align: center;">DISCONTINUE IF CLOUDINESS OR PRECIPITATE DEVELOPS</td> </tr> </tbody> </table>	DRUGS ADDED TO THIS INFUSION				PATIENT <i>A.patient (A.number)</i>			Ward <i>ICU</i>	Drug <i>Epoprostenol in 25mL diluent</i>	Amount <i>500 micrograms (20,000 nanograms/mL)</i>	Add By	Checked By	Date Added Time Added	Exp. Date Exp. Time	Batch No.		DISCONTINUE IF CLOUDINESS OR PRECIPITATE DEVELOPS			
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Appendix 1: Nebulised Epoprostenol Dosing Table

Epoprostenol Concentration: 20,000 nanograms/mL (0.5mg vial in 25mL solvent)						
Height (cm)		Dose (nanograms/Kg/min)				
Male	Female	≈10	≈20	≈30	≈40	≈50
<150	<155	1.4	2.7	4.1	5.4	6.8
150-159	155-164	1.5	3.3	5.0	6.6	8.3
160-169	165-174	1.8	3.9	5.9	7.8	9.8
170-179	175-184	2.1	4.5	6.8	9.0	11.3
180-189	185-194	2.4	5.1	7.7	10.2	12.0*
>190	>195	2.7	5.7	8.6	11.4	12.0*
Values above give the required flow rate (mL/h). Values followed by * have been capped at this rate.						

Documentation Controls

Development of Guideline:	Pharmacist – Critical Care & Theatres
Consultation with:	Pharmacy Department, Critical Care Nursing & Medical teams
Approved By:	Adult Drug Monograph Process Written/Reviewed by Munthar Miah December 2023 Checked By: Tien Vu December 2023
Review Date:	December 2026
Key contact:	Pharmacist – Critical Care & Theatres

References

GlaxoSmithKline UK. Summary of Product Characteristics (SPC) - Flolan 0.5 mg Powder and Solvent for Solution for Infusion (with pH 12 solvent). medicines.org.uk. [Online] [Accessed: 12/12/23] <https://www.medicines.org.uk/emc/product/7396/smpc>.

Aerogen. Aerogen Solo Instruction Manual. Aerogen.com. [Online] [Accessed: 12/05/20] [https://u5i6p3z8.stackpathcdn.com/wp-content/uploads/2019/11/30-354-Rev-T-Aerogen-Solo-System-UK WEB.pdf](https://u5i6p3z8.stackpathcdn.com/wp-content/uploads/2019/11/30-354-Rev-T-Aerogen-Solo-System-UK_WEB.pdf).

Buckley, M. S., Agarwal, S. K., Garcia-Orr, R., Saggat, R., & MacLaren, R. (2020). Comparison of Fixed-Dose Inhaled Epoprostenol and Inhaled Nitric Oxide for Acute Respiratory Distress Syndrome in Critically Ill Adults. *Journal of intensive care medicine*. <https://doi.org/10.1177/0885066620906800>

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