

VELETRI® (epoprostenol) **Sensitivity of VELETRI to Light in an Extension Line**

CLINICAL DATA

In a study that was performed to measure the stability of VELETRI for injection at different temperatures following dilution to various concentrations of epoprostenol, experimental samples were not shielded from light.¹

Additionally, during the studies EPITOME-1² and EPITOME-2³ comparing the administration of epoprostenol sodium for injection (e.g., Flolan®) and VELETRI for injection in patients with PAH, light-sensitive lines were not used.²⁻⁴

Direct exposure to sunlight may lead to warming of the VELETRI for injection solution and consequently, a negative impact on product potency. For this reason, VELETRI for injection solutions should be shielded from direct sunlight.⁵

LITERATURE SEARCH

A literature search of MEDLINE®, EMBASE®, BIOSIS Previews®, DERWENT® (and/or other resources, including internal/external databases) was conducted on 27 March 2024.

REFERENCES

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3. Sitbon O, Delcroix M, Bergot E, et al. EPITOME-2: An open-label study assessing the transition to a new formulation of intravenous epoprostenol in patients with pulmonary arterial hypertension. *Am Heart J.* 2014;167(2):210-217.
4. Data on File. Epoprostenol. AC-066A401/402 Clinical Study Report (EPITOME-1) and AC-066A301 Clinical Study Report (EPITOME-2). Janssen Scientific Affairs, LLC; 2021.
5. Epoprostenol. Company Core Data Sheet. Janssen Scientific Affairs, LLC. EDMS-ERI-205912254. 2021.