

## **FLEXBUMIN 25% [Albumin (Human)]**

- FLEXBUMIN 25% [Albumin (Human)], USP, 25% Solution is contraindicated in patients with cardiac failure, in patients with severe anemia and in patients with a history of allergic reactions to human albumin.
- FLEXBUMIN 25% is made from human plasma. Products made from human plasma may contain infectious agents, such as viruses, that can cause disease. Intravenous infusion rates should not exceed 1mL/min to patients with normal blood volume, due to the risk of developing circulatory overload and pulmonary edema. The quick rise in blood pressure, which may follow rapid administration, necessitates careful observation of the injured or post-operative patient to detect and treat severed blood vessels which failed to bleed at lower blood pressure.
- There exists a risk of potentially fatal hemolysis and acute renal failure from the inappropriate use of Sterile Water for Injection as a diluent.
- Adverse reactions to albumin are rare and may include nausea, fever, chills or urticaria, which may disappear when the infusion rate is slowed or stopped for a short period of time.

**CAUTION:** Do not use plastic containers in series connections. Such use could result in air embolism due to residual air being drawn from the primary container before the administration of the fluid from the secondary container is complete.

Please see Preparation for FLEXBUMIN 25% [Albumin (Human)] Administration in the [Prescribing Information](#).