

Albumin (Normal Human Serum)

Brand names

50 mg/mL: Albuked 5, Albuminar-5, AlbuRx 5, Albumin (Human) 5% Solution, Albutein 5%, Buminate 5%, Flexbumin 5%, Plasbumin-5

200 mg/mL: Albuminar-20, Human Albumin Grifols 20%, Plasbumin-20

250 mg/mL: Albuked 25, Albuminar-25, Albumin (Human) 25% Solution, Albutein 25%, AlbuRx 25, Buminate 25%, Flexbumin 25%, Human Albumin Grifols 25%, Kedbumin, Plasbumin-25

Medication error potential None noted

Contraindications and warnings

Contraindications: Patients who (1) are severely anemic, (2) have cardiac failure, or (3) have known hypersensitivity to albumin products.⁽³²⁾

When administering concentrated albumin (20% to 25%) to patients with marked dehydration, monitor closely as these patients may require additional fluids.^(19,33)

Concentrated albumin (20% to 25%) solutions are relatively low in electrolytes compared to 4% to 5% solutions. Monitor patient electrolyte status regularly.⁽¹⁹⁾

Some albumin products (Buminate 5%, Buminate 25%) contain components made with natural rubber latex.^(34,35)

Anaphylaxis has been reported in an *adult*.⁽²⁾

Infusion-related cautions

Too rapid infusion may result in acute hypertension or vascular overload, causing pulmonary edema or cardiac failure.^(18,32) Infusion over a longer time (8–12 hours in patients with nephrotic syndrome) decreases the risk for the development of CHF.⁽³⁾

Monitor blood pressure in trauma patients and postoperative patients being resuscitated with albumin 5% to detect rebleeding secondary to clot disruption.⁽³⁴⁾

Premature neonates are at risk for intraventricular hemorrhage from rapid intravascular volume expansion.⁽⁴⁾

Allergic reactions may result in chills, fever, nausea, vomiting, or urticaria.⁽²⁵⁾

Dosage

Hypovolemia: 0.5–1.25 g/kg (or generally 10–20 mL/kg).^(25,35,36) 20 mL/kg of 4.5% (0.9 g/kg)⁽⁵⁾ and 5% albumin (1 g/kg)⁽⁶⁾ have been given over 20 minutes in premature neonates.

A post hoc analysis of an *adult* randomized controlled trial found that NS was preferable to albumin 4% for acute resuscitation of patients with severe traumatic brain injury (improved mortality and neurologic outcomes).⁽⁷⁾

Hypoalbuminemia: Albumin deficits have been replaced by intermittent infusions of up to 1 g/kg of albumin 25%.⁽⁸⁾ Manufacturers recommend that doses should not exceed 2 g/kg/day^(34,35,37) or 25 g/day (in children) in the absence of ongoing albumin losses.⁽³³⁾ Continuous infusion of albumin results in a more sustained increase in serum albumin concentration.⁽⁹⁾ The albumin deficit can be estimated with the following equation:

$$\text{g albumin} = \text{weight (kg)} \times 3 \text{ dL/kg} \times (3.5 - \text{observed serum albumin in g/dL})^{(10,11)}$$

Nephrotic syndrome (controversial): 0.25–1 g/kg of albumin 25% infused over ≥ 1 –12 hours.^(3,12,13) Furosemide (0.5–2 mg/kg)^(3,12-14) infusions may accompany or follow the albumin infusion and have been shown to result in better weight loss, diuresis, and sodium excretion compared to either therapy alone.^(14,15) Alternatively, 1 g/kg of albumin 20% over 4 hours with furosemide continuous infusion of 0.3 mg/kg/hr has also been used.⁽¹⁵⁾

Neonatal hyperbilirubinemia: In one randomized study, neonates receiving 1 g/kg of albumin 20% 1 hour before exchange transfusion had a greater reduction in total serum bilirubin and required a shorter duration of phototherapy compared to those receiving exchange transfusions without albumin.⁽¹⁶⁾ Albumin 25% may be used.⁽²¹⁾



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Dosage adjustment in organ dysfunction	None noted. ⁽²⁵⁾ Consider using low aluminum-containing products in patients with chronic kidney failure. ⁽³¹⁾
Maximum dosage	For hypoalbuminemia, should not exceed 2 g/kg/day ^(34,35,37) or 25 g/day (in children) in the absence of ongoing albumin losses. ⁽³³⁾ Should not exceed 250 g in 48 hours in <i>adults</i> . ⁽¹⁾
Additives	Contains 130–160 mEq sodium/L of albumin. ^(17,24) Aluminum is present as a contaminant. Formulations containing ≤ 200 mcg/L aluminum include AlbuRx 5, AlbuRx 25, Albutein 5%, Albutein 25%, Human Albumin Grifols 20%, Human Albumin Grifols 25%, Kedbumin, and Plasbumin-25. ^(18-24,26) Contains no preservatives or antimicrobials. ^(17,24,26)
Suitable diluents	Manufacturers recommend dilution with D5W or NS. ^(21,26,38,39) Albumin is stable in D10W. ⁽¹⁷⁾ In patients who require sodium restriction, a 5% albumin solution that contains less sodium can be prepared by diluting each 1 mL of albumin 25% in 4 mL of D5W or D10W. However, infusion of large amounts of albumin diluted with D5W can result in hyponatremia; therefore, when using large volumes of albumin, dilution in NS is preferred. ^(26,33) Kidney failure and fatal hemolysis may occur if SW is used as a diluent. ^(17,27,28)
Maximum concentration	Undiluted (25%, 250 mg/mL)
Preparation and delivery	Compatibility: <i>Parenteral products should be visually inspected for particulate matter and discoloration before use. Refer to appropriate reference for more information on compatibility with other drugs and solutions, compatibility following Y-site delivery, and suggested storage and extended stability.</i> ⁽¹⁷⁾ Delivery: Do not use if solution is turbid. ⁽¹⁷⁾ The infusion should begin within 4 hours of opening the bottle. ^(17,34) Institution-specific protocols for blood product infusions should be used to guide infusion duration from an opened bottle. Solutions containing >25 g/L of albumin are more likely to occlude 0.22-micron inline filters; however, PN solutions containing as little as 10.8 g/L caused filter occlusion. ^(29,30) Manufacturers recommend that albumin not be mixed with protein hydrolysates because it may cause proteins to precipitate. ^(34,35,39)
IV push	When used as a plasma volume expander in the treatment of hypovolemic shock, the rate of administration should be adapted to the patient response. ^(18,36) (See the Infusion-Related Cautions section.)
Intermittent infusion	In the absence of overt shock, infuse albumin 5% at 1–2 mL/min (60–120 mL/hr). ⁽²⁰⁾ In severe hypovolemia, faster rates may be required. ⁽²⁶⁾ Manufacturers recommend infusion rates of 200–600 mL/hr for albumin 5% (faster if necessary in patients with decreased plasma volume except in patients with cardiovascular disease), ^(32,34) and rates not to exceed 120 mL/hr for albumin 25%. ⁽³³⁾ (See the Infusion-Related Cautions section.)
Continuous infusion	Can be infused continuously. (See the Preparation and Delivery section.)
Other routes of administration	Not indicated

