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**Developed for Scotland by the
National Plasma Product Expert Advisory Group**

Clinical Guidelines for Human Albumin Use

National Plasma Products Expert Advisory Group

Guidelines for the Usage of Human Albumin Solution (HAS)

1. Ascites and large volume paracentesis:

Where management of ascites is **refractory** to sodium restriction (90mmol/day) HAS is indicated following paracentesis (British Society of Gastroenterology Guidelines, 2006; EASL clinical practice guidelines, 2010). Paracentesis without albumin replacement leads to a fall in pulmonary capillary wedge pressure, maximal at 6 hours, and can result in circulatory and renal dysfunction.

Where there is **normal** pre-morbid renal function:

- Administer **1 unit (100ml) HAS 20%** (Human Albumin Solution, 20% i.e. 20g albumin per 100ml) (STAT) following every **3 litres** of ascites drained.

Where renal function is **impaired** consider either:

- Administration of **100ml HAS 20%** per **2 litres** of ascites.

or adherence to the protocol for hepatorenal syndrome (see below).

2. Spontaneous bacterial peritonitis (SBP):

Administration of HAS in the setting of SBP reduces the incidence of renal failure and reduces mortality (British Society of Gastroenterology Guidelines, 2006; EASL clinical practice guidelines, 2010):

- **Day 1: 1.5g HAS / kg** given over a 6 hour period:
- **Day 3: 1g HAS / kg** given over 3 hours

Recent evidence suggests that this can be restricted to “high” risk patients (Poca et al 2012).

3. Hepatorenal syndrome (HRS)

Administration of HAS and vasoconstrictors are effective therapy in 60% of patients with HRS and is associated with improved survival (EASL clinical practice guidelines, 2010):

- **Terlipressin: 0.5 – 2mg iv every 4 hours, plus**
- **Day 1: 1g / kg HAS**
- **Day 2 - 16: 20 – 40 g HAS / day**

Rx continued until serum creatinine falls below 130mol/l. NB. Where creatinine is **rising** despite Rx, **60g HAS /day** may be clinically indicated.

Therapeutic apheresis (Therapeutic Plasma Exchange TPE)

TPE has a clearly defined role in a large variety of conditions that are presumed to be immunologically mediated. The indications for TPE are protean and listed in a recent guideline produced by the American Society for Apheresis (Schwartz et al 2013). The standard replacement fluid for TPE is 5% HAS with or without Gelofusine. FFP / Octaplas may be indicated dependent on the indication e.g. TTP. The volume treated per procedure is usually: 1–1.5 total plasma volumes (TPV). Procedures may continue daily for up to several weeks in some cases.

Other indications

There is **little evidence** to support the use of HAS in other circumstances (Cochrane Injuries Group 2011; Jacob *et al* 2008; Kuper *et al* 2007; Perel and Roberts 2011) and these require discussion with pharmacy or haematology medical staff before release can be sanctioned, unless specifically requested by a Consultant Physician with frequent experience of HAS administration (eg. Gastroenterology, Neonatology Consultant).

Volume expansion – Synthetic alternatives or Saline 0.9% may be used for temporary intravascular volume expansion. No studies have convincingly shown that 5% Albumin confers any survival advantage.

Use of albumin in patients with decompensated liver disease is the subject of ongoing research (ATTIRE trial 2016).

Requesting Human Albumin Solution:

Human Albumin Solution (HAS) is available from the **Hospital Pharmacy**. **The indication for its use should be documented in the case notes and there should be written documentation of its prescription.**

The products in stock are:

Human Albumin Solution 5% 500ml (approx. 25g albumin) & 100ml (approx. 5g albumin)
Human Albumin Solution 20% 100ml (approx. 20g albumin)

Administration:

20% Albumin is hyperoncotic. The 100ml volume will expand to approximately 400mls within 25 minutes of transfusion. Rapid administration can lead to rapid volume expansion and cardiac failure. There is no UK published data regarding 20% Albumin infusion rates; in clinical practice it is usual to infuse 100mls over 30 minutes (personal communication AL/RS), but infusion rate varies and depends on clinical circumstances.

References:

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EASL clinical practice guidelines on the management of ascites, spontaneous bacterial peritonitis and hepato-renal syndrome in cirrhosis. *Journal of Hepatology* 2010; 53: 397-417.

Jacob M et al: Small-volume resuscitation with hyperoncotic albumin: a systematic review of randomized clinical trials. *Critical Care* 2008 Vol 12 No 2 R34.

Kuper M et al: The short-term effect of hyperoncotic albumin, given alone or with furosemide, on oxygenation in sepsis-induced Acute Respiratory Distress Syndrome. *Anaesthesia* 2007; 62: 259 – 263.

Patel A, Laffan MA, Waheed U, Brett SJ. Randomised trials of human albumin for adults with sepsis: systematic review and meta-analysis with trial sequential analysis of all-cause mortality. *BMJ* 2014;349:g4561.

Perel P, Roberts I. Colloids versus crystalloids for fluid resuscitation in critically ill patients. *Cochrane Database Syst Rev* 2011; 30: CD000567.

Poca M et al: Role of albumin treatment in patients with spontaneous bacterial peritonitis. *Clin Gastroenterol Hepatol* 2012; 10(3): 309 – 315.

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SUMMARY OF INDICATIONS FOR HUMAN ALBUMIN SOLUTION

	INDICATION	ALBUMIN PRODUCT	MAY BE AUTHORISED BY
1.	Large volume paracentesis (drainage of ascites) in patient with chronic liver disease.	HAS 20% 100ml 1 bottle per every 3L ascites drained (increased to 1 bottle every per every 2L if renal function impaired).	Junior medical staff on advice of Consultant Gastroenterologist or ITU Consultant.
2.	Large volume paracentesis (drainage of ascites) in patient with ascites not due to chronic liver disease.	May be indicated dependent on clinical circumstances.	Refer request to duty Pharmacy / Consultant Haematologist.
3.	Spontaneous bacterial peritonitis (SBP).	HAS 20% 100ml 1.5g/kg on day one (usually 4-8 bottles), then 1g/kg on day 3 (usually 2-5 bottles) or HAS 5% 500ml 1g.5/kg on day one (usually 3-6 bottles), then 1g/kg on day 3 (usually 2-4 bottles).	Junior medical staff on advice of Consultant Gastroenterologist or ITU Consultant. 5% or 20% products at discretion of Gastroenterologist or ITU Consultant.
4.	Hepatorenal syndrome.	HAS 20% 100ml 1g/kg on day (usually 2-5 bottles) or HAS 5% 500ml 1g/kg on day 1 (usually 2-4 bottles) then HAS 20% 100ml or HAS 5% 500ml 1-2 bottles daily for 2-16 days (occasionally 3 bottles/day).	Junior medical staff on advice of Consultant Gastroenterologist or ITU Consultant. 5% or 20% products at discretion of Gastroenterologist or ITU Consultant.
5.	Therapeutic Apheresis	HAS 5% 500ml 1-1.5 plasma volumes daily	At request of Consultant Haematologist, Apheresis Unit staff
6.	Low serum albumin in haemodynamically unstable patient in critical care (HDU or ITU).	Occasionally HAS 20% may be indicated.	At request of Consultant Anaesthetist/Intensivist in ITU/HDU.
7.	Low serum albumin in the absence of ascites, SBP or hepatorenal syndrome.	Not usually indicated.	Refer request to duty Pharmacist.
8.	All other requests.	Other indications are rare.	Refer request to duty Pharmacist.