

HIGHLIGHTS OF PRESCRIBING INFORMATION

These highlights do not include all the information needed to use HepaGam B® safely and effectively. See full prescribing information for HepaGam B®.

HepaGam B® [Hepatitis B Immune Globulin Intravenous (Human)],
Sterile Solution for Intravenous or Intramuscular Injection Solvent/Detergent Treated and Filtered. **>312 IU per milliliter** (Measured potency stamped on the vial label)

Initial U.S. Approval: 2006

RECENT MAJOR CHANGES

Dosage and Administration, Preparation (2.3)	1/2012
Warning and Precautions, Thrombotic events (5.7)	1/2012
Clinical Trials in Liver Transplant Patients (14.1)	1/2012

INDICATIONS AND USAGE

- For Intravenous or Intramuscular Administration Only
- Prevention of Hepatitis B recurrence following Liver Transplantation in HBsAg-positive liver transplant patients (1.1).
- Postexposure Prophylaxis (1.2) in the following settings:
 - Acute Exposure to Blood Containing HBsAg
 - Perinatal Exposure of Infants Born to HBsAg-positive Mothers
 - Sexual Exposure to HBsAg-positive Persons
 - Household Exposure to Persons with Acute HBV Infection

DOSAGE AND ADMINISTRATION

Prevention of Hepatitis B recurrence following liver transplantation (2.1)

HepaGam B is administered intravenously at doses of 20,000 IU (calculated from the measured potency stamped on the vial label) according to the following regimen to attain serum anti-HBs > 500 IU per liter:

Anhepatic Phase	Week 1 Post-Operative	Weeks 2-12 Post-Operative	Month 4 onwards
First dose	Daily from Day 1-7	Every two weeks from Day 14	Monthly

Regularly monitor serum anti-HBs to allow for treatment adjustments.

Postexposure Prophylaxis (2.2)

HepaGam B must be administered intramuscularly only as directed below:

Acute Exposure to Blood Containing HBsAg	0.06 milliliter per kilogram	Administer as soon as possible after exposure and within 24 hours if possible.
Perinatal Exposure of Infants Born to HBsAg-positive Mothers	0.5 milliliter	Administer after physiologic stabilization of the infant and preferably within 12 hours of birth.
Sexual Exposure to HBsAg-positive Persons	0.06 milliliter per kilogram	Administer HepaGam B within 14 days of the last sexual contact or if sexual contact with the infected person will continue.

Household Exposure to Persons with Acute HBV Infection	0.5 milliliter	Infants < 12 months: Administer HepaGam B + Hepatitis B vaccine if primary caregiver has acute HBV infection.
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DOSAGE FORMS AND STRENGTHS

1.0 milliliter sterile solution; single use vial (> 312 IU per milliliter). (3)
5.0 milliliters sterile solution; single use vial (> 312 IU per milliliter). (3)
The measured potency of each lot is stamped on the vial label.

CONTRAINDICATIONS

- History of anaphylactic or severe systemic reactions to human globulins (4)
- IgA deficient individuals may have the potential to develop IgA antibodies and have an anaphylactoid reaction. (4)
- IM injections may be contraindicated in patients with coagulation disorders. (4)

WARNINGS and PRECAUTIONS

- Hypersensitivity (5.1)
- Interference with Blood Glucose Testing (5.2)
- Monitoring: Serum Anti-HBs Antibody Levels (5.3)
- Infusion Reactions (5.4)
- Transmissible Infectious Agents (5.5)
- Coagulation Disorders (5.6)
- Thrombotic Events (5.7)

ADVERSE REACTIONS

The only adverse reactions observed in clinical trial subjects were hypotension and nausea (2% of clinical trial subjects). (6)

To report SUSPECTED ADVERSE REACTIONS, contact Cangene Corporation at 1-800-768-2304 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch

DRUG INTERACTIONS

- Efficacy of live attenuated virus vaccines may be impaired by immune globulin administration; revaccination may be necessary. (7.1)
- Antibodies in HepaGam B may interfere with some serological tests. (7.2)
- Maltose in HepaGam B may interfere with non-glucose specific blood glucose testing systems. (7.3)

USE IN SPECIFIC POPULATIONS

- Pregnancy: No human or animal data. Use only if clearly needed. (8.1)
- Nursing mothers: Caution should be exercised (8.3)

See 17 for PATIENT COUNSELING INFORMATION

FULL PRESCRIBING INFORMATION: CONTENTS*

Revised: 01/2012

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FULL PRESCRIBING INFORMATION

1 INDICATIONS AND USAGE

HepaGam B [Hepatitis B immune globulin intravenous (Human)] is an intravenous immune globulin indicated for the following:

1.1 Prevention of Hepatitis B recurrence following liver transplant in HBsAg-positive liver transplant patients

1.2 Post-exposure prophylaxis including

- acute exposure to HBsAg-positive blood, plasma, or serum (parenteral exposure, direct mucus membrane contact, oral ingestion, etc),
- perinatal exposure of infants born to HBsAg-positive mothers,
- sexual exposure to HBsAg-positive persons, and
- household exposure to persons with acute HBV infection.

2 DOSAGE AND ADMINISTRATION

2.1 Prevention of Hepatitis B recurrence following liver transplantation

Administer the first dose of HepaGam B during the grafting of the transplanted liver (the anhepatic phase) with subsequent dosing as recommended in Table 1.

Calculate the dosing from the measured potency of the particular lot of HepaGam B as stamped on the vial label.

Administer by intravenous infusion (Table 2).

Table 1 - HepaGam B Dosing Regimen for HBV-Related Liver Transplant Patients

Anhepatic Phase	Week 1 Post-Operative	Weeks 2-12 Post-Operative	Month 4 onwards
First dose	Daily from Day 1-7	Every two weeks from Day 14	Monthly

* Each dose should contain 20,000 IU calculated from the measured potency as stamped on the vial label [see *Dosage Forms and Strengths* (3)].

Table 2 – HepaGam B Intravenous Infusion Rate

Route of Administration	Dosage	Infusion Rate
Intravenous	20,000 IU per dose	2 milliliters per minute. Decrease to 1 milliliter per minute or slower if the patient develops discomfort or infusion-related adverse reactions.

HepaGam B dose adjustments may be required in patients who fail to reach anti-HBs levels of 500 International Units per liter within the first week post-liver transplantation¹. Patients who have surgical bleeding or abdominal fluid drainage (> 500 milliliters) or patients who undergo plasmapheresis are particularly susceptible to extensive loss of circulated anti-HBs. In these cases, the dosing regimen should be increased to a half-dose (10,000 International Units calculated from the measured potency as stamped on the vial label) intravenously every 6 hours until the target anti-HBs is reached.

2.2 Postexposure Prophylaxis

Administer HepaGam B intramuscularly as recommended in Table 3.

Table 3 – HepaGam B Dosing Regimen for Postexposure Prophylaxis (Intramuscular)

Indication	Dosage	Instructions
Acute Exposure to Blood Containing HBsAg	0.06 milliliter per kilogram	Administer HepaGam B as soon as possible after exposure. The value after seven days following exposure is unclear ^{2,3} . For persons who refuse Hepatitis B vaccine or who are known non-responders to vaccine, give a second dose of HepaGam B one month after the first dose ² .
Perinatal exposure of Infants Born to HBsAg-positive mothers	0.5 milliliter	Administer after physiologic stabilization of the infant and preferably within twelve hours of birth. Administer concurrently with Hepatitis B vaccine.
Sexual Exposure to HBsAg-	0.06 milliliter per	Administer HepaGam B and Hepatitis B Vaccine series within 14 days of sexual

Indication	Dosage	Instructions
Positive Persons	kilogram	contact or if sexual contact with the infected person will continue.
Household Exposure to Person with Acute HBV Infection	0.5 milliliter	For infants less than twelve months of age administered concurrently with Hepatitis B Vaccine. Prophylaxis of other household contacts of persons with acute HBV infection is not indicated unless there is an identifiable blood exposure to the index patient, such as by sharing toothbrushes or razors. Treat such exposures like sexual exposures.

HepaGam B may be administered at the same time (but at a different site), or up to one month preceding Hepatitis B vaccination without impairing the active immune response to Hepatitis B vaccine^{2,3}.

2.3 Preparation

- Parenteral drug products should be inspected visually for particulate matter and discoloration prior to administration, whenever solution and container permit. Do not use if turbid.
- Do not shake vials during preparation to avoid foaming.
- The HepaGam B vial is for single use only. HepaGam B contains no preservatives.
- Promptly use any vial of HepaGam B that has been entered. Do not reuse or save for future use.

- For intravenous administration, administer HepaGam B through a separate intravenous line using an infusion pump.
- Use normal saline as the diluent if dilution of HepaGam B is preferred prior to intravenous administration. [see *Clinical Trials in Liver Transplant Patients (14.1)*]
- Do not use dextrose (5%) in water (D5W).
- Use a separate vial, sterile syringe, and needle for each individual patient, to prevent transmission of infectious agents from one person to another.

3 DOSAGE FORMS AND STRENGTHS

- HepaGam B is a sterile solution of purified gamma globulin (5% or 50 milligrams per milliliter) that contains greater than 312 International Units per milliliter of anti-HBs.
- The measured potency of each lot is stamped on the vial label.
- To ensure that the label claim of >312 International Unit per milliliter is maintained over the product shelf life, a higher potency of 550 International Unit per milliliter is targeted at the time of manufacture.
 - This higher target potency is a manufacturing requirement to account for variability in the potency assay and changes in potency over time.
 - The potency assay has a relative standard deviation (RSD) of approximately 10%.
 - The actual potency test result may vary from approximately 400 to 700 International Units per milliliter (3x RSD) based on statistical assessment of manufactured lots with a target potency of 550 International Unit per milliliter.
- Calculate the dosing for the prevention of hepatitis B recurrence following liver transplantation from the measured potency of the particular lot of HepaGam B as stamped on the vial label.

4 CONTRAINDICATIONS

- Individuals known to have anaphylactic or severe systemic reactions to the parenteral administration of human globulin preparations should not receive HepaGam B.
- Individuals who are deficient in IgA may have the potential to develop anti-IgA antibodies and have an anaphylactoid reaction.
 - HepaGam B contains less than 40 micrograms per milliliter of IgA.
- For postexposure prophylaxis indications, HepaGam B must be administered intramuscularly only. In patients who have severe thrombocytopenia or any coagulation disorder that would contraindicate intramuscular injections, HepaGam B should be given only if the expected benefits outweigh the potential risks.

5 WARNINGS AND PRECAUTIONS

5.1 Hypersensitivity

Severe hypersensitivity reactions may occur with HepaGam B. HepaGam B should be administered in a setting with appropriate equipment, medication and personnel trained in the management of hypersensitivity, anaphylaxis and shock. In case of hypersensitivity, discontinue HepaGam B infusion immediately and begin appropriate emergency treatment. Medications such as epinephrine and antihistamines should be available for immediate treatment of acute hypersensitivity reactions.

HepaGam B contains trace amounts of IgA (<40 micrograms per milliliter). Patients with known antibodies to IgA may have a greater risk of severe hypersensitivity and anaphylactic reactions. HepaGam B is contraindicated in IgA deficient patients with antibodies against IgA and history of hypersensitivity reaction. (see

CONTRAINDICATIONS [4])

5.2 Interference with Blood Glucose Testing

The maltose contained in HepaGam B can interfere with some types of blood glucose monitoring systems, i.e., those based on the glucose dehydrogenase pyrroloquinone (GDH-PQQ) method. This can result in falsely elevated glucose readings and, consequently, in the inappropriate administration of insulin, resulting in life-threatening hypoglycemia. Cases of true hypoglycemia may go untreated if the hypoglycemic state is masked by falsely elevated results.

5.3 Monitoring: Serum Anti-HBs Antibody Levels

Liver transplant patients should be monitored regularly for serum anti-HBs antibody levels using a quantitative assay to ensure that adequate protective levels are maintained.

5.4 Infusion Reactions

Certain adverse drug reactions may be related to the rate of infusion. The recommended infusion rate given under dosage and administration (2.1) must be closely followed. Patients must be closely monitored and carefully observed for any symptoms throughout the infusion period and immediately following an infusion.

5.5 Transmissible Infectious Agents

Because HepaGam B is made from human plasma, it may carry a risk of transmitting infectious agents, e.g. viruses, and, theoretically, the Creutzfeldt-Jakob disease (CJD) agent. No cases of transmission of viral diseases or CJD have been associated with the use of HepaGam B. All infections suspected by a physician possibly to have been transmitted by this product should be reported by the physician or other healthcare provider to Cangene Corporation at 1-800-768-2304.

5.6 Coagulation Disorders

For postexposure prophylaxis indications, HepaGam B must be administered intramuscularly only. In patients who have severe

thrombocytopenia or any coagulation disorder that would contraindicate intramuscular injections, HepaGam B should be given only if the expected benefits outweigh the potential risks.

5.7 Thrombotic Events

Thrombotic events may occur during or following treatment with IGIV products^{4,5}. Patients at risk include those with a history of atherosclerosis, multiple cardiovascular risk factors, advanced age, impaired cardiac output, coagulation disorders, prolonged periods of immobilization, and/or known/suspected hyperviscosity.

Consider baseline assessment of blood viscosity in patients at risk for hyperviscosity including those with cryoglobulins, fasting chylomicronemia/markedly high triacylglycerols (triglycerides), or monoclonal gammopathies. For patients who are at risk of developing thrombotic events, administer HepaGam B at the minimum rate of infusion practicable.

6 ADVERSE REACTIONS

The only adverse reactions observed in clinical trial subjects were hypotension and nausea (2% of clinical trial subjects).

6.1 Clinical Trials Experience

Because clinical trials are conducted under widely varying conditions, adverse reaction rates observed in the clinical trials of a drug cannot be directly compared to rates in the clinical trials of another drug and may not reflect the rates observed in practice.

Hepatitis B-Related Liver Transplantation

In a clinical trial with 27 liver transplant patients, one adverse drug reaction was reported following the 578 (<1%) HepaGam B infusions. This study utilized the recommended dosing regimen outlined in Table 1 [see Dosage and Administration (2.1)]. The attributed adverse drug reaction of hypotension was reported in one patient. The reaction was associated with a single HepaGam B infusion during the first day post-transplant. The

reaction resolved on the same day and did not recur with subsequent HepaGam B infusions.

Healthy Volunteer Studies

Seventy healthy male and female volunteers received a single dose of HepaGam B intramuscularly in clinical trials⁶. Only one adverse drug reaction, an episode of nausea, was reported.

6.2 Postmarketing Experience

The following adverse reactions have been identified during postapproval use of HepaGam B. Because these reactions are reported voluntarily from a population of uncertain size, it is not always possible to reliably estimate their frequency or establish a causal relationship to drug exposure.

Postexposure Prophylaxis:

Dizziness has been reported in the postmarketing surveillance of HepaGam B for the postexposure prophylaxis indication.

Hepatitis B-Related Liver Transplantation:

The system organ classification of reported adverse reactions is provided below:

Cardiac disorders:	Sinus tachycardia
Gastrointestinal disorders:	Abdominal pain upper Nausea
General disorders and administration site conditions:	Chills Feeling cold Influenza like illness Pyrexia
Immune system disorders:	Anaphylactoid reaction Hypersensitivity
Investigations:	Lipase increased Transaminases increased
Musculoskeletal and	Back pain

connective tissue disorders:

Nervous system disorders: Dizziness
 Headache

**Respiratory, thoracic and
mediastinal disorders:** Dyspnoea

**Skin and subcutaneous
tissue disorders:** Cold sweat

Healthcare professionals should report adverse reactions following the administration of HepaGam B to Cangene Corporation at 1-800-768-2304 or FDA's MedWatch reporting system at 1-800-FDA-1088 or www.fda.gov/medwatch.

7 DRUG INTERACTIONS

7.1 Live Attenuated Virus Vaccines

Immune globulin administration may impair the efficacy of live attenuated virus vaccines such as measles, rubella, mumps and varicella^{2,3,7}. Vaccination with live virus vaccines should be deferred until approximately three months after administration of HepaGam B, Hepatitis B Immune Globulin Intravenous (Human). Persons who received HepaGam B less than 14 days after live virus vaccination should be revaccinated 3 months after the administration of the immune globulin, unless serologic test results indicate that antibodies were produced^{2,3}.

There are no available data on drug interactions of HepaGam B with other medications.

7.2 Drug-Laboratory Interactions: Serological Testing

Antibodies present in HepaGam B may interfere with some serological tests. After administration of immune globulins like HepaGam B, a transitory increase of passively transferred antibodies in the patient's blood may result in misleading positive results in serological testing (e.g. Coombs' test).

7.3 Drug-Laboratory Interactions: Blood Glucose Testing

HepaGam B contains maltose which can interfere with certain types of blood glucose monitoring systems. [See *Warnings and Precautions* (5.3).] Only testing systems that are glucose-specific should be used in patients receiving HepaGam B. This interference can result in falsely elevated glucose readings that can lead to untreated hypoglycemia or to inappropriate insulin administration, resulting in life-threatening hypoglycemia.

The product information of the blood glucose testing system, including that of the test strips, should be carefully reviewed to determine if the system is appropriate for use with maltose-containing parenteral products. If any uncertainty exists, contact the manufacturer of the testing system to determine if the system is appropriate for use with maltose-containing parenteral products.

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Pregnancy Category C. Animal reproduction studies have not been conducted with HepaGam B. It is also not known whether HepaGam B can cause fetal harm when administered to a pregnant woman or can affect reproductive capacity. HepaGam B should be given to a pregnant woman only if clearly indicated.

8.3 Nursing Mothers

It is not known whether HepaGam B is excreted in human milk. Because many drugs are excreted in human milk, caution should be exercised when HepaGam B is administered to a nursing mother.

8.4 Pediatric Use

Safety and effectiveness have not been established in pediatric patients. However, for postexposure prophylaxis, the safety and effectiveness of similar hepatitis B immune globulins have been demonstrated in infants and children⁸.

8.5 Geriatric Use

Clinical studies of HepaGam B did not include sufficient numbers of subjects aged 65 and over to determine whether they respond differently from younger subjects. Other reported clinical experience has not identified differences in responses between the elderly and younger patients. In general, dose selection for an elderly patient should be cautious, usually starting at the low end of the dosing range, reflecting the greater frequency of decreased hepatic, renal, or cardiac function, and of concomitant disease or other drug therapy.

10 OVERDOSAGE

Consequences of an overdose are not known. For intramuscular administration of HepaGam B, the only manifestations of overdose would be pain and tenderness at the injection site.

11 DESCRIPTION

HepaGam B, Hepatitis B Immune Globulin Intravenous (Human), is a solvent/detergent-treated sterile solution of purified gamma globulin containing anti-HBs. It is prepared from plasma donated by healthy, screened donors with high titers of anti-HBs that is purified by an anion-exchange column chromatography manufacturing method^{9,10}. HepaGam B is formulated as a 5% (50 milligrams per milliliter) protein solution with 10% maltose and 0.03% polysorbate 80 at pH 5.6. It is available in 1 milliliter and 5 milliliters single dose vials. The product appears as a clear to opalescent liquid. HepaGam B does not contain mercury. It contains no preservatives. This product is intended for single use. HepaGam B may be administered intravenously or intramuscularly dependent upon indication [see *Dosage and Administration* (2.)]. The source plasma used in the manufacture of this product was tested by FDA licensed Nucleic Acid testing (NAT) for HIV-1, HBV and HCV and found to be negative. Plasma also has been tested by in-process NAT for hepatitis A virus (HAV) and parvovirus B19 (B19) via minipool testing and the limit for B19 in the manufacturing pool is set not to exceed 10⁴ IU of B19 DNA per milliliter.

The manufacturing process contains two steps implemented specifically for virus clearance. The solvent and detergent step (using tri-n-butyl phosphate and Triton X-100) is effective in the inactivation of enveloped viruses, such as hepatitis B, hepatitis C and HIV¹¹. Virus filtration, using a Planova 20N virus filter, is effective for the removal of viruses based on their size, including some non-enveloped viruses¹². These two viral clearance steps are designed to increase product safety by reducing the risk of transmission of enveloped and non-enveloped viruses. In addition to these two specific steps, the process step of anion-exchange chromatography was identified as contributing to the overall viral clearance capacity for small non-enveloped viruses.

The inactivation and reduction of known enveloped and non-enveloped model viruses were validated in laboratory studies as summarized in Table 4. The viruses employed for spiking studies were selected to represent those viruses that are potential contaminants in the product, and to represent a wide range of physiochemical properties in order to challenge the manufacturing process's ability for viral clearance in general.

Table 4 - Virus Reduction Values Obtained Through Validation Studies⁸

Enveloped	Enveloped			Non-Enveloped			
Genome	RNA		DNA	RNA		DNA	
Virus	HIV-1	BVDV	PRV	HAV	EMC	MMV	PPV
Family	retro	flavi	herpes	picorna		parvo	
Size (nm)	80-100	50-70	120-200	25-30	30	20-25	18-24
Anion Exchange Chromatography (partitioning)	Not evaluated			2.3	n.e.	3.4	n.e.
20N Filtration (size exclusion)	≥ 4.7	≥ 3.5	≥ 5.6 ^a	n.e.	4.8	n.e.	4.1
Solvent/Detergent (inactivation)	≥ 4.7	≥ 7.3	≥ 5.5	Not evaluated			
Total Reduction (log ₁₀)	≥ 9.4	≥ 10.8	≥ 11.1	2.3	4.8	3.4	4.1

Abbreviations:

HIV-1: human immunodeficiency virus-1; relevant virus for human immunodeficiency virus-1 and model for HIV-2

BVDV: bovine viral diarrhea virus; model virus for hepatitis C virus (HCV) and West Nile virus (WNV)

PRV: pseudorabies virus; model for large enveloped DNA viruses, including herpes

HAV: human hepatitis A virus; relevant virus for HAV and model for small non-enveloped viruses in general

EMC: encephalomyocarditis virus; model for HAV and for small non-enveloped viruses in general

MMV: murine minute virus; model for human parvovirus B19 and for small non-enveloped viruses in general

PPV: porcine parvovirus; model for human parvovirus B19 and for small non-enveloped viruses in general

n.e.: not evaluated

^a The PRV was retained by the 0.1 µm pre-filter during the virus validation. Since manufacturing employs a 0.1 µm pre-filter before the 20N filter, the claim of ≥ 5.6 reduction is considered applicable.

The product potency is expressed in international units (IU) by comparison to the World Health Organization (WHO) standard Hepatitis B Immune Globulin. Each vial contains greater than 312 IU per milliliter. The measured potency of each lot is stamped on the vial label [*see Dosage Forms and Strengths (3)*].

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

HepaGam B provides passive immunization for individuals exposed to the hepatitis B virus, by binding to the surface antigen and reducing the rate of hepatitis B infection¹³⁻¹⁶.

12.2 Pharmacokinetics

The pharmacokinetic profile of HepaGam B has been evaluated in two 84-day clinical trials in which 70 healthy subjects received an intramuscular injection of 0.06 milliliter per

kilogram of HepaGam B. The mean peak concentrations (C_{max}) in both studies were comparable and occurred within 4-5 days of administration. Both studies demonstrated mean elimination half-lives ($t_{1/2}$) following IM administration of 22 to 25 days. The mean clearance rate was 0.21 to 0.24 liter per day and the volume of distribution was approximately 7.5 liter. Thus, HepaGam B demonstrates pharmacokinetic parameters similar to those reported by Scheiermann and Kuwert¹⁷.

The maximum concentration of anti-HBs achieved by HepaGam B was consistent with that of two other licensed comparator Hepatitis B Immune Globulin (Human) products⁶. Comparability of pharmacokinetics between HepaGam B and a commercially available hepatitis B immune globulin product administered IM indicates that comparable efficacy of HepaGam B should be inferred.

14 CLINICAL STUDIES

14.1 Clinical Trials in Liver Transplant Patients

A clinical trial examined the effectiveness of HepaGam B in the prevention of hepatitis B recurrence following liver transplantation. The study was a multi-center, open-labeled, superiority study involving HBsAg-positive/HBeAg-negative liver transplant patients. The study included two arms, an active treatment group of patients enrolled to receive the described dosing regimen of HepaGam B starting during transplant and continuing over the course of a year, and a retrospective untreated control group of historical patients with data gathered by chart review.

There were 27 liver transplant patients who received HepaGam B and 14 retrospective untreated control patients. The patients in both groups were HBsAg-positive/HBeAg-negative liver transplant patients who met similar entry criteria, had similar medical history and had similar status at transplant based on MELD and/or ChildPugh-Turcotte scores.

In the active treatment group, HepaGam B intravenous doses of 35 milliliters were initiated

during transplant according to the regimen identified in Table 1 [see *Dosing and Administration* (2.1)]. As a result of the targeted potency of 550 IU per milliliter at the time of manufacture [see *Dosage Forms and Strengths* (3.)], the 35 milliliter doses of HepaGam B used in this study actually contained between 17,000 and 23,000 IU anti-HBs. These 35 milliliter doses consistently yielded anti-HBs trough levels > 500 IU per liter (99% of all anti-HBs levels were > 500 IU per liter; see Figure 1). Patients received HepaGam B doses diluted with 50 mL of saline.

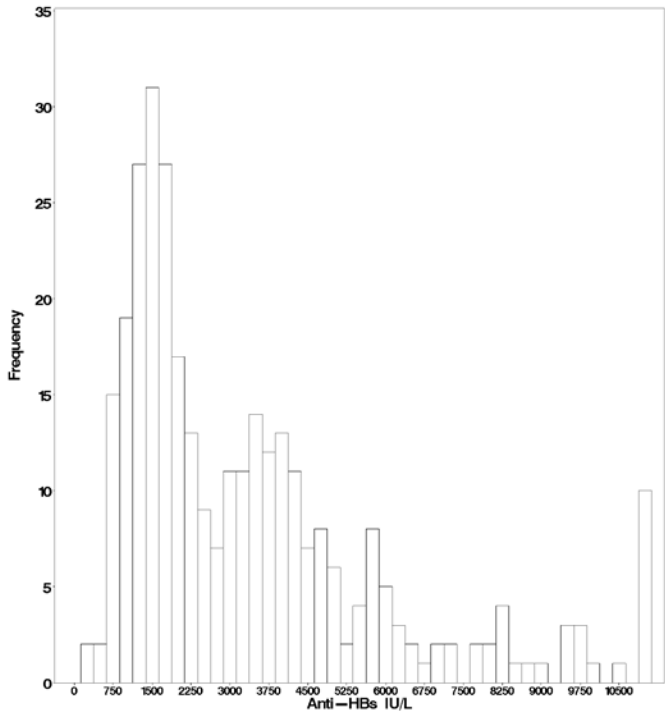


Figure 1: Frequency Histogram of Trough anti-HBs Levels more than 30 days after Transplant

Values below the target trough were only observed in the 2 patients with HBV recurrence who had anti-HBs levels <150 IU per liter at the time of seroconversion.

For the efficacy endpoint of the proportion of patients with HBV recurrence (HBsAg positive and/or HBeAg positive after 4 weeks post-OLT), a significant treatment effect was observed. As summarized in Table 5, HBV recurrence was seen in 2/24 or 8.3% of HepaGam B patients compared to 12/14 or 86% of retrospective untreated control patients (see Table 5). Two of the HepaGam B patients who died within 28 days post-transplant

were excluded from all efficacy analyses, but included for safety analyses. The deaths were not HBV or study drug related.

Table 5 - Results of Study HB-005 for the Prevention of Hepatitis B Recurrence Following Liver Transplantation

	HepaGam B	Retrospective Untreated Control	P-value (Fisher's Exact Test)
HBV Recurrence Proportion, % (95% confidence interval)	8.3 (0.1 -27.0)	85.7 (57.2 -98.2)	< 0.001

The conclusion that HepaGam B monotherapy post-OLT is effective at preventing HBV recurrence post-OLT is further supported by the secondary endpoints of time to recurrence, survival, anti-HBs levels, biochemical markers of liver inflammation, and liver biopsy. Time to recurrence for the HepaGam B treatment group was 358 days for two HBV recurrent patients. In comparison, the retrospective untreated control patients had a median time to recurrence of 88 days with a 95% confidence interval of 47 to 125 days. Survival calculations showed that 96% (23/24) of patients in the active treatment group survived for at least 1 year post-OLT compared to 43% (6/14) retrospective control patients. The endpoints for HBV recurrence were supported by an observed drop in anti-HBs levels, elevated liver function tests, and abnormal liver biopsy result at the time of recurrence.

HepaGam B is recommended in patients who have no or low levels of viral replication at the time of liver transplantation. The clinical trial evaluating HepaGam B in liver transplant patients selected patients with no or low replication status only. HepaGam B therapy has not been evaluated in combination with antiviral therapy post-transplantation.

15 REFERENCES

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16 HOW SUPPLIED/STORAGE AND HANDLING

NDC 53270-0052-1; a carton containing a 1.0 milliliter single dose vial (>312 IU per milliliter; measured potency of each lot is stamped on the vial label) and a package insert.

NDC 53270-0051-1; a carton containing a 5.0 milliliter single dose vial (>312 IU per milliliter; measured potency of each lot is stamped on the vial label) and a package insert.

Store at 36 to 46 °F (2 to 8 °C). Do not freeze. Do not use after expiration date. Use within 6 hours after the vial has been entered.

17 PATIENT COUNSELING INFORMATION

- Inform patients of the following:
 - HepaGam B is prepared from human plasma, and therefore, may contain infectious agents such as viruses that can cause disease.
 - The risk that such products will transmit an infectious agent has been reduced by screening plasma donors for prior exposure to certain viruses, by testing for the presence of certain current virus infections, and by inactivating and/or removing certain viruses during manufacturing.
 - Despite these measures, such products can still potentially transmit disease.
 - There is also the possibility that unknown infectious agents may be present in such products.
- Tell patients that persons known to have severe, potentially life-threatening reactions to human globulin products should not receive HepaGam B or any other immune globulin products unless the risk has been justified.
- Tell patients that persons who are deficient in IgA may have the potential for developing anti-IgA antibodies and have severe potentially life threatening allergic reactions.

- In case of allergic or anaphylactic reaction, the infusion should be stopped immediately.
- In case of shock, the current medical standards for treatment of shock should be observed.
- Advise Liver Transplant patients about the potential interference with non-glucose specific monitoring systems.
 - The maltose contained in HepaGam B can interfere with some types of blood glucose monitoring systems.
 - Only testing systems that are glucose-specific should be used in patients receiving HepaGam B.
 - This inference can result in falsely elevated glucose readings that can lead to untreated hypoglycemia or to inappropriate insulin administration, resulting in life-threatening hypoglycemia.

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