

LUMASON- sulfur hexafluoride

LUMASON- sulfur hexafluoride injection, powder, lyophilized, for suspension BRACCO DIAGNOSTICS INC

HIGHLIGHTS OF PRESCRIBING INFORMATION

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

LUMASON (sulfur hexafluoride lipid-type A microspheres) for injectable suspension, for intravenous use or intravesical use
Initial U.S. Approval: 2014

WARNING: SERIOUS CARDIOPULMONARY REACTIONS

See full prescribing information for complete boxed warning

Serious cardiopulmonary reactions, including fatalities, have occurred uncommonly during or following the injection of ultrasound contrast agents, including sulfur hexafluoride lipid microspheres (5.1). Most serious reactions occur within 30 minutes of administration (5.1).

- Assess all patients for the presence of any condition that precludes administration (4).
- Always have resuscitation equipment and trained personnel readily available (5.1).

RECENT MAJOR CHANGES

Contraindications (4)

4/2021

Warnings and Precautions, Hypersensitivity Reactions (5.2)

4/2021

INDICATIONS AND USAGE

Lumason is an ultrasound contrast agent indicated for use

- in echocardiography to opacify the left ventricular chamber and to improve the delineation of the left ventricular endocardial border in adult and pediatric patients with suboptimal echocardiograms (1)
- in ultrasonography of the liver for characterization of focal liver lesions in adult and pediatric patients (1)
- in ultrasonography of the urinary tract for the evaluation of suspected or known vesicoureteral reflux in pediatric patients (1)

DOSAGE AND ADMINISTRATION

Avoid intra-arterial injection (2.1, 5.3)

See Full Prescribing Information for reconstitution instructions (2.3)

For intravenous injection:

- Echocardiography in adults: After reconstitution, administer 2 mL as an intravenous injection (2.2, 2.4)
- Echocardiography in pediatric patients: After reconstitution, administer 0.03 mL per kg as an intravenous injection up to a maximum of 2 mL per injection (2.2, 2.4)
- Ultrasonography of the liver in adults: After reconstitution, administer 2.4 mL as an intravenous injection (2.2, 2.4)
- Ultrasonography of the liver in pediatric patients: After reconstitution, administer 0.03 mL per kg as an intravenous injection, up to a maximum of 2.4 mL per injection (2.2, 2.4)
- May repeat dose one time during a single examination (2.2, 2.4)
- Follow each injection with an intravenous flush of 0.9% Sodium Chloride Injection, USP (2.2, 2.4)

For intravesical administration in pediatric patients:

- Ultrasonography of the urinary tract: After reconstitution, administer 1 mL via sterile 6 to 8F urinary catheter. Bladder should be first emptied and then partially filled with 0.9% Sodium Chloride Injection, USP before injection of Lumason (2.2, 2.4)
- After Lumason administration, continue filling the bladder with 0.9% Sodium Chloride Injection, USP until the patient has the urge to micturate or at the first sign of back pressure to the infusion (2.4)

DOSAGE FORMS AND STRENGTHS

- For injectable suspension: 25 mg of lipid-type A lyophilized powder with headspace fill of 60.7 mg sulfur hexafluoride in a single-patient use vial for reconstitution (3)

CONTRAINDICATIONS

- Hypersensitivity to sulfur hexafluoride lipid microspheres or its components, such as polyethylene glycol (PEG) (4)

WARNINGS AND PRECAUTIONS

- Cardiopulmonary reactions, including fatalities. Always have resuscitation equipment and trained personnel readily available (5.1)
- Hypersensitivity reactions. Serious acute hypersensitivity reactions have occurred in patients with no prior exposure to sulfur hexafluoride lipid-containing microsphere products, including patients with prior hypersensitivity reaction(s) to PEG (5.2, 6)

ADVERSE REACTIONS

Most common adverse reactions (incidence $\geq 0.5\%$) are headache and nausea (6.1).

To report SUSPECTED ADVERSE REACTIONS, contact Bracco Diagnostics Inc at 1-800-257-5181 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch

See 17 for PATIENT COUNSELING INFORMATION.

Revised: 7/2023

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FULL PRESCRIBING INFORMATION

WARNING: SERIOUS CARDIOPULMONARY REACTIONS

Serious cardiopulmonary reactions, including fatalities, have occurred uncommonly during or following the injection of ultrasound contrast agents, including sulfur hexafluoride lipid microspheres [see *Warnings and Precautions (5.1)*]. Most serious reactions occur within 30 minutes of administration [see *Warnings and Precautions (5.1)*].

- Assess all patients for the presence of any condition that precludes administration [see *Contraindications (4)*].
- Always have resuscitation equipment and trained personnel readily available [see *Warnings and Precautions (5.1)*].

1 INDICATIONS AND USAGE

Echocardiography

Lumason is indicated for use in adult and pediatric patients with suboptimal echocardiograms to opacify the left ventricular chamber and to improve the delineation of the left ventricular endocardial border.

Ultrasonography of the Liver

Lumason is indicated for use with ultrasound of the liver in adult and pediatric patients to characterize focal liver lesions.

Ultrasonography of the Urinary Tract

Lumason is indicated for use in ultrasonography of the urinary tract in pediatric patients for the evaluation of suspected or known vesicoureteral reflux.

2 DOSAGE AND ADMINISTRATION

2.1 Important Administration Instructions

Do not administer Lumason by intra-arterial injection [see *Warnings and Precautions* (5.3)].

2.2 Recommended Dosage

Echocardiography

Adults

The recommended dose of Lumason after reconstitution is 2 mL administered as an intravenous bolus injection during echocardiography. During a single examination, a second injection of 2 mL may be administered to prolong contrast enhancement. Follow each Lumason injection with an intravenous flush using 5 mL of 0.9% Sodium Chloride Injection, USP.

Pediatric Patients

The recommended dose of Lumason after reconstitution in pediatric patients is 0.03 mL per kg administered as an intravenous injection during echocardiography. During a single examination, a second injection of 0.03 mL per kg may be administered, if needed. Do not exceed 2 mL per injection. Follow Lumason injection with an intravenous flush using 5 mL of 0.9% Sodium Chloride Injection, USP.

Ultrasonography of the Liver

Adults

The recommended dose of Lumason after reconstitution in adult patients is 2.4 mL administered as an intravenous injection during ultrasonography of the liver. During a single examination, a second injection of 2.4 mL may be administered, if needed. Follow Lumason injection with an intravenous flush using 5 mL of 0.9% Sodium Chloride Injection, USP.

Pediatric Patients

The recommended dose of Lumason after reconstitution in pediatric patients is 0.03 mL per kg administered as an intravenous injection during ultrasonography of the liver. During a single examination, a second injection of 0.03 mL per kg may be administered, if needed. Do not exceed 2.4 mL per injection. Follow Lumason injection with an intravenous flush of 0.9% Sodium Chloride Injection, USP.

Ultrasonography of the Urinary Tract

Pediatric Patients

The recommended dose of Lumason after reconstitution is 1 mL. The bladder may be refilled with 0.9% Sodium Chloride Injection, USP for a second cycle of voiding and imaging, without the need of a second Lumason administration.

2.3 Reconstitution Instructions

- Refer to Section 2.3.1 for instructions for using the single patient use kit with diluent provided
- Refer to Section 2.3.2 for instructions for using the 20-vial pack without diluent provided

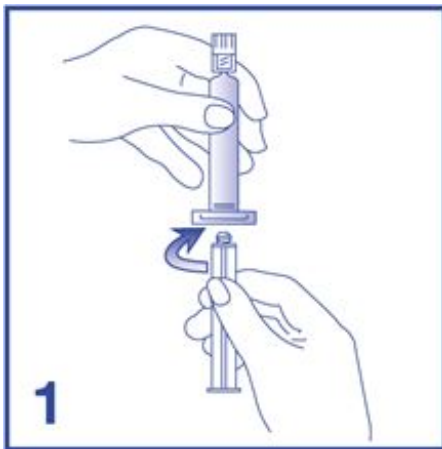
2.3.1 Lumason Kit (single patient use kit)

Contents of Lumason Kit

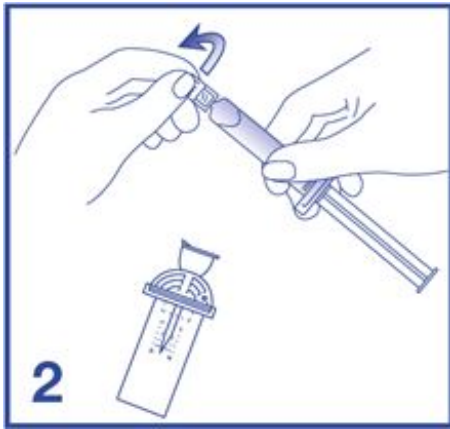


- Inspect the Lumason kit and its components for signs of damage. Do not use the kit if the protective caps on the Lumason vial and prefilled syringe with 5 mL 0.9% Sodium Chloride Injection, USP are not intact or if the kit shows other signs of damage.
- Under aseptic conditions, reconstitute the Lumason vial using the following illustrated steps:

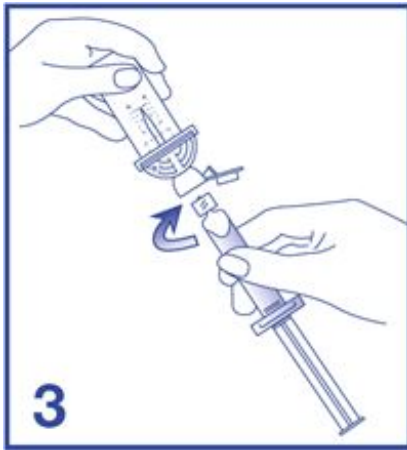
1. Connect the plunger rod to the prefilled 0.9% Sodium Chloride Injection, USP syringe barrel by screwing it clockwise into the syringe (see Figure 1).



2. Open the Mini-Spike blister and remove the syringe tip cap (see Figure 2).



3. Remove the Mini-Spike green cap and connect the syringe to the Mini-Spike by screwing it in clockwise (see Figure 3).

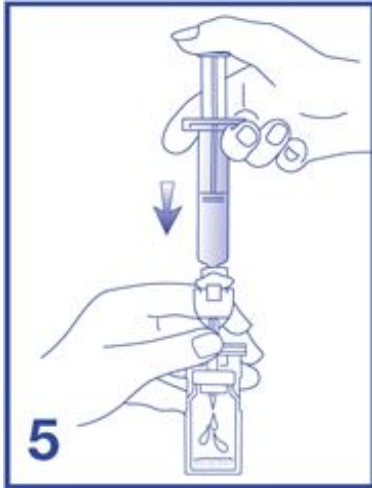


4. Remove the Mini-Spike spike protection and position the spike in the center of the rubber stopper of the vial. Press firmly inward until the spike is fully inserted in the stopper (see Figure 4).



5. Empty the content of the syringe into the vial by pushing on the plunger rod (see

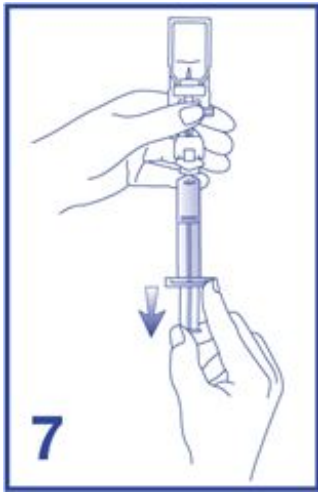
Figure 5).



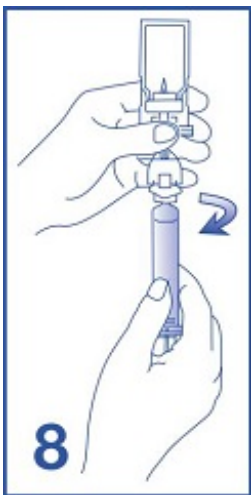
6. Shake vigorously for 20 seconds, mixing all the contents in the vial (see Figure 6). A homogeneous white milky liquid indicates formation of sulfur hexafluoride lipid microspheres.



7. For preparation of doses greater than or equal to 1 mL, invert the system and slowly withdraw the intended volume of suspension into the syringe (see Figure 7). For preparation of doses less than 1 mL, withdraw 2 mL of the reconstituted suspension into the 5 mL syringe and measure the volume of Lumason to inject by using the 0.2 mL graduations between the 1 mL and 2 mL marks.



8. Unscrew the syringe from the Mini-Spike (see Figure 8). Peel and remove the diluent label to display the reconstituted product label. For intravenous administration, immediately connect the syringe to a dose administration line (20 G) and administer as directed under the Administration Instructions below. For intravesical administration, immediately connect the syringe to a sterile urinary catheter (6 French to 8 French) and administer as directed under the Administration Instructions below.



- Following reconstitution, Lumason suspension contains 1.5 to 5.6×10^8 microspheres/mL with 45 mcg/mL of sulfur hexafluoride.
- Use immediately after reconstitution. If the suspension is not used immediately after reconstitution, resuspend the microspheres for a few seconds by hand agitation before the suspension is drawn into the syringe. Reconstituted suspension within a vial may be used for up to 3 hours from the time of its reconstitution. Maintain the vial containing the reconstituted suspension at room temperature 25°C (77°F); excursions permitted to 15° to 30°C (59° to 86°F).

2.3.2 Lumason Pack (20-vial pack)

Contents of Lumason Pack



20 x Glass vials
(Lumason)



20 x Mini-Spikes



20 x peel-off syringe labels

*Please note: This presentation does not include pre-filled syringes of 0.9% Sodium Chloride Injection, USP (Diluent).

Lumason vials are to be used with the supplied Mini-Spike only.

Use only additive-free 0.9% Sodium Chloride Injection, USP for the reconstitution of Lumason.

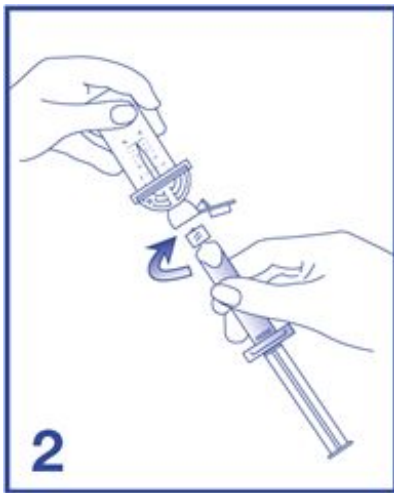
Reconstitution

- Inspect the Lumason components for signs of damage. Do not use the Lumason vial if the protective cap on the vial is not intact or other components in the pack show signs of damage.
 - Use aseptic conditions for the preparation and administration of Lumason.
1. Obtain a 5 mL syringe, with luer lock tip, and fill with 5 mL of additive-free 0.9% Sodium Chloride Injection, USP (diluent) (see Figure 1).
 - Two healthcare professionals (HCPs) should verify that the solution selected for reconstitution of Lumason is additive-free 0.9% Sodium Chloride Injection, USP.
 - a. Ensure that any air in the syringe is expelled.



[Note: A prefilled syringe containing additive-free 0.9% Sodium Chloride Injection, USP may be used. Ensure that any air in the syringe is expelled.]

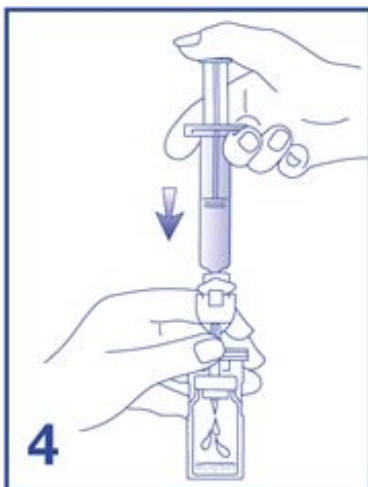
2. Remove the Mini-Spike green cap and connect the syringe to the Mini-Spike by screwing it in clockwise (see Figure 2).



3. Remove the Mini-Spike spike protection and position the spike in the center of the rubber stopper of the vial. Press firmly inward until the spike is fully inserted in the stopper (see Figure 3).



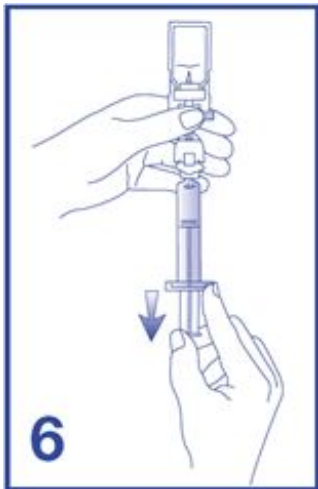
4. Empty the entire 5 mL content of the syringe into the vial by pushing on the plunger rod (see Figure 4).



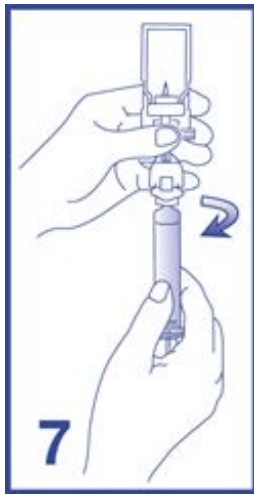
5. Shake vigorously for 20 seconds, mixing all the contents in the vial (see Figure 5). A homogeneous white milky liquid indicates formation of sulfur hexafluoride lipid microspheres.



6. To obtain required dose, invert the system and slowly withdraw the intended volume of suspension into the syringe (see Figure 6).



7. Unscrew the syringe from the Mini-Spike (see Figure 7).



8. Label the syringe using the peel-off sticker provided.

9. For intravenous administration, immediately connect the syringe to a dose administration line (20G) and administer as directed under the Administration Instructions below. For intravesical administration, immediately connect the syringe to a sterile urinary catheter (6 French to 8 French) and administer as directed under the Administration Instructions below.

- Following reconstitution, Lumason suspension contains 1.5 to 5.6×10^8 microspheres/mL with 45 mcg/mL of sulfur hexafluoride.
- Use immediately after reconstitution. If the suspension is not used immediately after reconstitution, resuspend the microspheres for a few seconds by hand agitation before the suspension is drawn into the syringe. Reconstituted suspension within a vial may be used for up to 3 hours from the time of its reconstitution. Maintain the vial containing the reconstituted suspension at room temperature 25°C (77°F); excursions permitted to 15° to 30°C (59° to 86°F).

2.4 Administration Instructions

Inspect visually for particulate matter and discoloration prior to administration, whenever solution and container permit. The reconstituted suspension is milky-white, and does not contain visible particulate matter. Do not use the single-patient-use vial for more than one patient.

Intravenous Administration

Administer Lumason as an intravenous bolus injection.

Intravesical Administration in Pediatric Patients

1. Insert a sterile 6 French to 8 French urinary catheter into the bladder under sterile conditions;
2. Empty the bladder of urine, and then fill the bladder with sterile 0.9% Sodium Chloride Injection, USP to approximately one third or half of its predicted total volume. The total bladder volume in children is calculated as $[(\text{age in years} + 2) \times 30]$ mL;
3. Administer Lumason as an intravesical bolus injection through the urinary catheter;
4. Continue filling the bladder with 0.9% Sodium Chloride Injection, USP until the patient has the urge to micturate or at the first sign of back pressure to the infusion.
5. Immediately following the first voiding, the bladder may be refilled with 0.9% Sodium Chloride Injection, USP for a second cycle of voiding and imaging, without the need of

a second Lumason administration.

2.5 Imaging Guidelines

Echocardiography

After baseline non-contrast echocardiography is complete, adjust the mechanical index for the ultrasound device to 0.8 or lower. Continue ultrasound imaging following Lumason injection.

Ultrasonography of the Liver

After identification of the target focal lesion on non-contrast ultrasound examination, hold transducer still while switching scanner to low mechanical index (≤ 0.4) contrast-specific imaging. Continue ultrasound imaging following Lumason injection.

Ultrasonography of the Urinary Tract

After baseline non-contrast ultrasound examination of the kidney and bladder, switch the scanner to low mechanical index (≤ 0.4) contrast specific imaging. Perform continuous alternate ultrasound imaging of the bladder, ureters, and kidneys during filling and voiding of the bladder.

3 DOSAGE FORMS AND STRENGTHS

For injectable suspension: Lumason is supplied in two presentations (single patient use kit or 20-vial pack):

3-part single patient use kit comprised of:

- one Lumason clear vial containing 25 mg of lipid-type A sterile white lyophilized powder with headspace filled with 60.7 mg of sulfur hexafluoride gas
- one prefilled syringe containing 5 mL of 0.9% Sodium Chloride Injection, USP (Diluent)
- one Mini-Spike

20-vial pack comprised of:

- twenty Lumason clear vials, each containing 25 mg of lipid-type A sterile white lyophilized powder with headspace filled with 60.7 mg of sulfur hexafluoride gas
- twenty Mini-Spikes
- twenty peel-off syringe labels

Following reconstitution, Lumason is a homogeneous, milky white suspension containing 1.5 to 5.6×10^8 microspheres/mL with 45 mcg/mL of sulfur hexafluoride.

4 CONTRAINDICATIONS

Lumason is contraindicated in patients with known or suspected:

- Hypersensitivity to sulfur hexafluoride lipid microsphere or its components, such as polyethylene glycol (PEG) [see *Warnings and Precautions (5.2) and Description (11)*].

5 WARNINGS AND PRECAUTIONS

5.1 Serious Cardiopulmonary Reactions

Serious cardiopulmonary reactions, including fatalities have occurred uncommonly during or shortly following administration of ultrasound contrast agents, including Lumason. These reactions typically occurred within 30 minutes of administration. The risk for these reactions may be increased among patients with unstable cardiopulmonary conditions (acute myocardial infarction, acute coronary artery syndromes, worsening or unstable congestive heart failure, or serious ventricular arrhythmias). Always have cardiopulmonary resuscitation personnel and equipment readily available prior to Lumason administration and monitor all patients for acute reactions.

The reported reactions that may follow the administration of ultrasound contrast agents include: fatal cardiac or respiratory arrest, shock, syncope, symptomatic arrhythmias (atrial fibrillation, tachycardia, bradycardia, supraventricular tachycardia, ventricular fibrillation, and ventricular tachycardia), hypertension, hypotension, dyspnea, hypoxia, chest pain, respiratory distress, stridor, wheezing, loss of consciousness, and convulsions.

5.2 Hypersensitivity Reactions

In postmarketing use, serious hypersensitivity reactions were observed during or shortly following sulfur hexafluoride lipid-containing microsphere administration including:

Anaphylaxis, with manifestations that may include death, shock, bronchospasm, dyspnea, throat tightness, angioedema, edema (pharyngeal, palatal, mouth, peripheral, localized), swelling (face, eye, lip, tongue, upper airway), facial hypoesthesia, rash, urticaria, pruritus, flushing, and erythema.

These reactions may occur in patients with no history of prior exposure to sulfur hexafluoride lipid-containing microspheres. Lumason contains PEG. There may be increased risk of serious reactions including death in patients with prior hypersensitivity reaction(s) to PEG [see *Adverse Reactions* (6.2)]. Clinically assess patients for prior hypersensitivity reactions to products containing PEG, such as certain colonoscopy bowel preparations and laxatives. Always have cardiopulmonary resuscitation personnel and equipment readily available prior to Lumason administration and monitor all patients for hypersensitivity reactions.

5.3 Systemic Embolization

When administering Lumason to patients with cardiac shunt, microspheres can bypass filtering by the lung and enter the arterial circulation. Assess patients with shunts for embolic phenomena following Lumason administration. Lumason is only for intravenous and/or intravesical administration; do not administer Lumason by intra-arterial injection [see *Dosage and Administration* (2.1)].

5.4 Ventricular Arrhythmia Related to High Mechanical Index

High ultrasound mechanical index values may cause microsphere cavitation or rupture and lead to ventricular arrhythmias. Additionally, end-systolic triggering with high mechanical indices has been reported to cause ventricular arrhythmias.

Lumason is not recommended for use at mechanical indices greater than 0.8.

6 ADVERSE REACTIONS

The following serious adverse reactions are discussed elsewhere in the labeling:

- Cardiopulmonary reactions [see *Warnings and Precautions* (5.1)]
- Hypersensitivity reactions [see *Warnings and Precautions* (5.2)]

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.

Adults

In completed clinical trials, a total of 6984 adult subjects (128 healthy volunteers and 6856 patients) received Lumason at cumulative doses ranging from 0.2 to 161 mL (mean 9.8 mL). Lumason was administered mainly as single or multiple injections; however, some subjects received infusion dosing. The majority (75%) of subjects received Lumason at cumulative doses of 10 mL or less. There were 64% men and 36% women, with an average age of 59 years (range 17 to 99 years). A total of 79% subjects were White; 4% were Black; 16% were Asian; <1% were Hispanic; and <1% were in other racial groups or race was not reported.

In the clinical trials, serious adverse reactions were observed in 2 subjects; one who experienced a hypersensitivity-type rash and presyncope and another who experienced anaphylactic shock shortly following Lumason administration.

The most commonly reported adverse reactions among patients (occurring among at least 0.2% of patients) are listed below (Table 1). Most adverse reactions were mild to moderate in intensity and resolved spontaneously.

Table 1. Adverse Reactions in Adult Patients* n = 6856	
Number (%) of Patients with Adverse Reactions	340 (5%)
Headache	65 (1%)
Nausea	37 (0.5%)
Dysgeusia	29 (0.4%)
Injection site pain	23 (0.3%)
Feeling Hot	18 (0.3%)
Chest discomfort	17 (0.2%)
Chest pain	12 (0.2%)
Dizziness	11 (0.2%)
Injection Site Warmth	11 (0.2%)

*occurring in at least 0.2% of patients

Pediatric Patients:

In completed clinical trials for echocardiography, a total of 12 pediatric patients received Lumason at a dose of 0.03 mL/kg. No adverse reactions were identified in pediatric patients [see *Clinical Studies* (14.1)].

6.2 Postmarketing Experience

In the international postmarketing clinical experience and clinical trials, serious adverse reactions have uncommonly been reported following administration of Lumason. 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. The serious adverse reactions include fatalities, especially in a pattern of symptoms suggestive of anaphylactoid/hypersensitivity reactions. Other serious reactions included arrhythmias and hypertensive episodes. These reactions typically occurred within 30 minutes of Lumason administration. These serious reactions may be increased among patients with pre-existing PEG hypersensitivity and/or unstable cardiopulmonary conditions (acute myocardial infarction, acute coronary artery syndromes, worsening or unstable congestive heart failure, or serious ventricular arrhythmias) [see *Warnings and Precautions* (5.1, 5.2)].

Hypersensitivity

Anaphylaxis, with manifestations that may include death, shock, bronchospasm, dyspnea, throat tightness, angioedema, edema (pharyngeal, palatal, mouth, peripheral, localized), swelling (face, eye, lip, tongue, upper airway), facial hypoesthesia, rash, urticaria, pruritus, flushing, and erythema.

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary

There are no data with Lumason use in pregnant women to inform any drug-associated risks. No adverse developmental outcomes were observed in animal reproduction studies with administration of sulfur hexafluoride lipid-type A microspheres in pregnant rats and rabbits during organogenesis at doses up to at least 10 and 20 times, respectively, the maximum human dose of 4.8 mL based on body surface area (see *Data*).

In the U.S. general population, the estimated background risk of major birth defects and miscarriage in clinically recognized pregnancies is 2-4% and 15-20%, respectively.

Data

Animal Data

Lumason was administered intravenously to rats at doses of 0.2, 1, and 5 mL/kg (approximately 0.4, 2, and 10 times the recommended maximum human dose of 4.8 mL, respectively, based on body surface area); Lumason doses were administered daily for about 30 consecutive days, from two weeks before pairing until the end of organogenesis. Lumason was administered intravenously to rabbits at doses of 0.2, 1, and 5 mL/kg (approximately 0.8, 4, and 20 times the recommended maximum human dose, respectively, based on body surface area); Lumason doses were administered daily from gestation day 6 to day 19 inclusive. No significant findings on the fetus were observed.

8.2 Lactation

Risk Summary

There are no data on the presence of Lumason in human milk, the effects on the breastfed infant, or the effects on milk production. The developmental and health benefits of breastfeeding should be considered along with the mother's clinical need for Lumason and any potential adverse effects on the breastfed infant from Lumason or from the underlying maternal condition.

8.4 Pediatric Use

Echocardiography

Safety and effectiveness have been established for use in pediatric patients with suboptimal echocardiograms to opacify the left ventricular chamber and to improve delineation of the left endocardial border. Safety and effectiveness in pediatric patients are based on adequate and well-controlled studies in adults and are supported by a clinical study in 12 pediatric patients (mean age: 13.8 years) with extrapolation of efficacy to younger pediatric patients. No new adverse reactions were reported in the pediatric study [see *Adverse Reactions (6.1)* and *Clinical Studies (14.1)*]. Safety of intravenous use of Lumason was based on evaluation of published literature involving the use of Lumason in over 1400 pediatric patients (0 to 17 years).

Ultrasonography of the Liver

Safety and effectiveness in pediatric patients has been established for use in ultrasonography of the liver for characterization of focal liver lesions from adequate and well controlled trials in adult patients and a clinical study of 44 pediatric patients [see *Clinical Studies (14)*]. Safety of intravenous use of Lumason was based on evaluation of published literature involving use of Lumason in over 1400 pediatric patients. Non-fatal anaphylaxis was reported in one pediatric patient.

Ultrasonography of the Urinary Tract

Safety and effectiveness in pediatric patients has been established for use in ultrasonography of the urinary tract for the evaluation of suspected or known vesicoureteral reflux from two published studies comprising a total of 411 pediatric patients [see *Clinical Studies (14)*]. Safety of intravesical use of Lumason was based on evaluation of published literature involving use of Lumason in over 6000 pediatric patients. No adverse reactions were reported.

8.5 Geriatric Use

Of the total number of 6856 adult patients in clinical studies of Lumason, 39% were 65 and over, while 11% were 75 and older. No overall differences in safety or effectiveness were observed between these subjects and younger subjects, and other reported clinical experience has not identified differences in responses between the elderly or younger patients, but greater sensitivity of some older individuals cannot be ruled out.

11 DESCRIPTION

Lumason (sulfur hexafluoride lipid-type A microspheres) for injectable suspension, for intravenous or intravesical use is used to prepare the ultrasound contrast agent. Lumason is supplied in two presentations (single patient use kit or 20-vial pack):

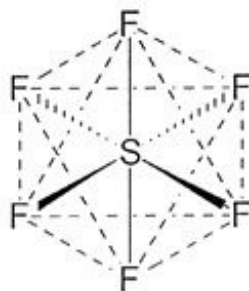
- The single patient use kit contains the following three items:
 1. one clear glass 10 mL vial containing 25 mg of white lyophilized powder lipid-type A, 60.7 mg of sulfur hexafluoride gas and capped with a blue flip-cap
 2. one prefilled syringe containing 5 mL 0.9% Sodium Chloride Injection, USP (Diluent)
Each prefilled syringe with 5 mL of diluent 0.9% Sodium Chloride Injection, USP is sterile, nonpyrogenic, and additive-free containing 9 mg sodium chloride per mL.
 3. one Mini-Spike
- The 20-vial pack is comprised of:
 1. twenty Lumason clear vials, each containing 25 mg of lipid-type A sterile white lyophilized powder with headspace filled with 60.7 mg of sulfur hexafluoride gas
 2. twenty Mini-Spikes
 3. twenty peel-off syringe labels

Each vial is formulated as a 25 mg sterile, pyrogen-free lyophilized powder containing 24.56 mg of polyethylene glycol 4000, 0.19 mg of distearoylphosphatidyl-choline (DSPC), 0.19 mg of dipalmitoylphosphatidylglycerol sodium (DPPG-Na) and 0.04 mg of palmitic acid. The headspace of each vial contains 6.07 mg/mL ($\pm 2\%$) sulfur hexafluoride, SF₆, or 60.7 mg per vial.

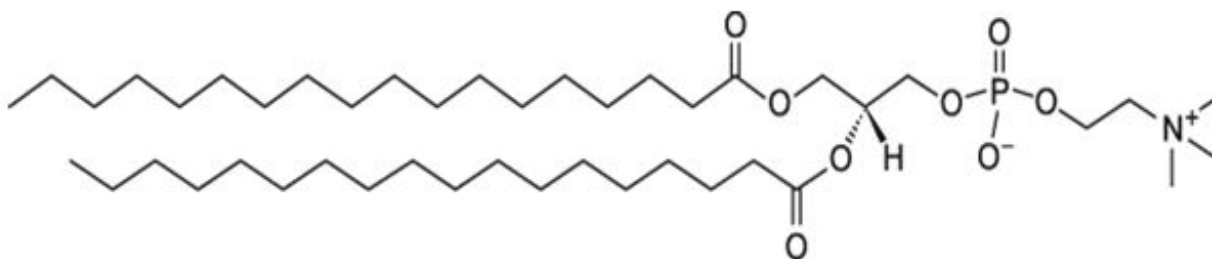
Upon reconstitution with 5mL diluent, Lumason is a milky white, homogeneous suspension containing sulfur hexafluoride lipid-type A microspheres. The suspension is isotonic and has a pH of 4.5 to 7.5.

The sulfur hexafluoride lipid microspheres are composed of SF₆ gas in the core surrounded by an outer shell monolayer of phospholipids consisting DSPC and DPPG-Na with palmitic acid as a stabilizer.

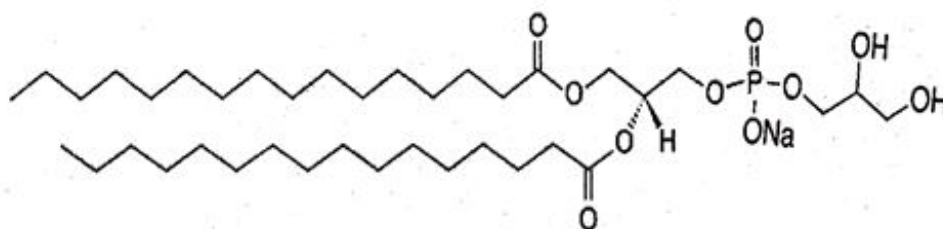
Sulfur hexafluoride has a molecular weight of 145.9 and the following chemical structure:



1,2-Distearoyl-*sn*-glycero-3-phosphocholine (DSPC), with empirical formula C₄₄H₈₈NO₈P, has a molecular weight of 790.6 and the following chemical structure:



1,2-Dipalmitoyl-*sn*-glycero-3-phospho-*rac*-glycerol sodium (DPPG-Na), with empirical formula $C_{38}H_{74}NaO_{10}P$, has a molecular weight of 745 and the following chemical structure:



Each milliliter of reconstituted Lumason suspension contains 1.5 to 5.6×10^8 microspheres, 68 mcg SF_6 (12 mcL), 0.038 mg DSPC, 0.038 mg DPPG-Na, 4.91 mg polyethylene glycol 4000 and 0.008 mg palmitic acid. The sulphur hexafluoride associated with the microspheres suspension is 45 mcg/mL. Fifteen to twenty three percent of the total lipids in the suspension are associated with the microspheres.

The sulfur hexafluoride lipid microsphere characteristics are listed in Table 2:

Table 2. Microsphere Characteristics	
Mean diameter range	1.5 – 2.5 μm
Percent of microspheres $\leq 10 \mu m$	$\geq 99\%$
Upper size limit	$100.0\% \leq 20 \mu m$

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

Within the blood, the acoustic impedance of Lumason microspheres is lower than that of the surrounding non-aqueous tissue. Therefore, an ultrasound beam is reflected from the interface between the microspheres and the surrounding tissue. The reflected ultrasound signal provides a visual image that shows a contrast between the blood and the surrounding tissues.

For ultrasonography of the urinary tract in pediatric patients, the intravesically administered Lumason microspheres increase signal intensity of fluids within the

urethra, bladder, ureters, and renal pelvis.

12.2 Pharmacodynamics

Lumason provides useful echocardiographic signal intensity for two minutes after intravenous injection. Lumason microspheres are destroyed and contrast enhancement decreases as the mechanical index increases (values of 0.8 or less are recommended).

For ultrasonography of the liver, Lumason provides dynamic patterns of differential signal intensity enhancement between focal liver lesions and liver parenchyma during the arterial, portal venous, and late phase of signal intensity enhancement of the microvasculature.

In ultrasonography of the urinary tract, Lumason facilitates the detection of reflux of fluid from the bladder into the ureters.

Pulmonary Hemodynamic Effects

The effect of Lumason on pulmonary hemodynamics was studied in a prospective, open-label study of 36 patients scheduled for right heart catheterization, including 18 with mean pulmonary arterial pressure (MPAP) > 25 mmHg and 18 with MPAP ≤ 25 mmHg. No clinically important pulmonary hemodynamic changes were observed. This study did not assess the effect of Lumason on visualization of cardiac or pulmonary structures.

12.3 Pharmacokinetics

The pharmacokinetic of the SF₆ gas component of Lumason was evaluated in 12 healthy adult subjects. After intravenous bolus injections of 0.03 mL/kg and 0.3 mL/kg of Lumason, corresponding to approximately 1 and 10 times the recommended doses, concentrations of SF₆ in blood peaked within 1 to 2 minutes for both doses. The terminal half-life of SF₆ in blood was approximately 10 minutes for the 0.3 mL/kg dose. The area-under-the-curve of SF₆ was dose-proportional over the dose range studied.

Distribution

In a study of healthy subjects, the mean values for the apparent steady-state volume of distribution of SF₆ following intravenous administration, were 341 mL and 710 mL for Lumason doses of 0.03 mL/kg and 0.3 mL/kg, respectively. Preferential distribution to the lung is likely responsible for these values.

Elimination

Following intravenous administration, the SF₆ component of Lumason is eliminated via the lungs. In a clinical study that examined SF₆ elimination twenty minutes following Lumason injection, the mean cumulative recovery of SF₆ in expired air was 82 ± 20% (SD) at the 0.03 mL/kg dose and 88 ± 26% (SD) at the 0.3 mL/kg dose.

SF₆ undergoes first pass elimination within the pulmonary circulation; approximately 40% to 50% of the SF₆ content was eliminated in the expired air during the first minute following Lumason injection.

Metabolism

SF₆ undergoes little or no biotransformation; following intravenous administration, 88% of an administered dose is recovered unchanged in expired air.

Pharmacokinetics in Specific Populations

Pulmonary Impairment:

In a study of patients with pulmonary impairment, blood concentrations of SF₆ peaked at 1 to 4 minutes following intravenous Lumason administration. The cumulative recovery of SF₆ in expired air was 102 ± 18% (mean ± standard deviation), and the terminal half-life of SF₆ in blood was similar to that measured in healthy subjects.

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

No long-term animal studies were performed to evaluate the carcinogenic potential of Lumason. No evidence of genotoxicity was found in the following studies conducted with Lumason: 1) a bacterial mutagenesis (Ames) assay, 2) an in vitro human lymphocyte chromosome aberration assay, and 3) an in vivo mouse micronucleus assay.

No impairment of fertility was observed in rats receiving Lumason at doses up to 8 times the human dose based on body surface area.

14 CLINICAL STUDIES

14.1 Echocardiography

Adults

A total of 191 patients with suspected cardiac disease and suboptimal non-contrast echocardiography received Lumason in three multi-center controlled clinical trials (76 patients in Study A, 62 patients in Study B, and 53 patients in Study C). Among these patients, there were 127 men and 64 women. The mean age was 59 years (range 22 to 96 years). The racial and ethnic representations were 79% White, 16% Black, 4% Hispanic, < 1% Asian, and < 1% other racial or ethnic groups. The mean weight was 204 lbs (range 92 to 405 lbs). Approximately 20% of the patients had a chronic pulmonary disorder and 30% had a history of heart failure. Of the 106 patients for whom a New York Heart Association (NYHA) classification of heart failure was assigned, 49% were Class I, 33% were Class II, and 18% were Class III. Patients with NYHA Class IV heart failure were not included in these studies.

In Studies A and B, each patient received four intravenous bolus injections of Lumason (0.5, 1, 2, and 4 mL), in randomized order. In Study C, each patient received two doses of Lumason (1 mL and 2 mL) in randomized order. All three studies assessed endocardial border delineation and left ventricular opacification. For each patient in each study, echocardiography with Lumason was compared to non-contrast (baseline) echocardiography. A recording of 2D echocardiography was obtained from 30 seconds prior to each injection to at least 15 minutes after dosing or until the disappearance of the contrast effect, whichever was longer. Contrast and non-contrast echocardiographic images for each patient were evaluated by two independent reviewers, who were blinded to clinical information and the Lumason dose. Evaluation of left ventricular endocardial border consisted of segment based assessment involving six endocardial segments and using two apical views (2- and 4-chamber views).

Endocardial Border Delineation and Duration of Useful Contrast Effect

In all three studies, administration of Lumason improved left ventricular endocardial border delineation. The majority of the patients who received a 2.0 mL dose of Lumason had improvement in endocardial border delineation manifested as visualization of at least two additional endocardial border segments. Table 3 demonstrates the improvement in endocardial border delineation following Lumason administration as a reduction in percentage of patients with inadequate border delineation in at least one pair of adjacent segments (combined 2-chamber and 4-chamber view). The results are shown by reader.

Table 3. Reduction in Percentage of Patients with Inadequate Border Delineation

Reader	Study A N = 76		Study B N = 62		Study C N = 53	
	Non-contrast	Lumason	Non-contrast	Lumason	Non-contrast	Lumason
A	60 (79%)	22 (33%)	31 (50%)	12 (19%)	12 (23%)	10 (19%)
B	62 (82%)	29 (37%)	54 (87%)	6 (10%)	45 (85%)	20 (38%)

Following the first appearance of contrast within the left ventricle the mean duration of useful contrast effect ranged from 1.7 to 3.1 minutes.

Left Ventricular Opacification

In all three studies, complete left ventricular opacification was observed in 52% to 80% of the patients following administration of a 2.0 mL dose of Lumason. The studies did not sufficiently assess the effect of Lumason upon measures of left ventricular ejection fraction and wall motion.

Pediatric patients:

Twelve pediatric patients 9 to 17 years of age with suspected cardiac disease and suboptimal non-contrast echocardiography received Lumason in one prospective multicenter clinical trial. Patients received Lumason at a dose of 0.03 mL/kg (mean 1.83 mL). There were 7 female, 10 white, and 2 black patients.

For both the non-contrast and contrast-enhanced images, standard apical 4-, 2-, and 3-chamber views with harmonic imaging were acquired. Contrast and non-contrast images for each patient were evaluated by three independent reviewers, who were blinded to clinical information.

Endocardial Border Delineation

Evaluation of left ventricular endocardial border delineation consisted of segment-based assessment of the left ventricle divided into 17 endocardial segments. The delineation of each segment's endocardial border was rated as inadequate, sufficient, or good. An exam was considered suboptimal if any of the patient's apical views had 2 or more adjacent segments with inadequate delineation scores.

The majority of screened patients had adequate delineation of the left ventricular endocardial border without administration of contrast. The number of patients with inadequate left ventricular endocardial border delineation without contrast and after

Lumason are shown for the 12 patients, by reader, in Table 4.

Table 4. Number of Pediatric Patients with Inadequate Border Delineation with and without Lumason			
	Reader A	Reader B	Reader C
Non-contrast	12/12	11/11 ^b	12/12
Lumason	1/11 ^a	0/9 ^{bb}	0/11 ^c

^a Reader A had missing segment data with contrast echocardiography for one patient; ^b Reader B had missing segment data with non-contrast echocardiography for one patient; ^{bb} Reader B had missing segment data with contrast echocardiography for three patients; ^c Reader C had missing segment data with contrast echocardiography for one patient

Left Ventricular Opacification

Complete left ventricular opacification was observed in all the patients by all 3 readers following administration of Lumason.

14.2 Ultrasonography of the Liver

Adults

A total of 499 patients with at least 1 focal liver lesion requiring characterization were evaluated in two studies (259 patients in Study A, 240 patients in Study B). Among these patients, there were 259 men and 240 women. The mean age was 56 years (range 19 to 93 years). The racial and ethnic representations were 74% White, 11% Black, 9% Hispanic, 5% Asian, and 1% other racial or ethnic groups. The mean weight was 80 kg (range 44 to 173 kg).

In both studies, prior to Lumason administration, gray scale and Doppler (color or power imaging) ultrasound examinations of the target lesion were performed using commercially available ultrasound equipment and using standard techniques. Each patient received an intravenous injection of 2.4 mL of Lumason (up to 2 injections were allowed, 91% patients received 1 injection). Following Lumason administration, ultrasound examination of the target lesion was carried out using contrast-specific imaging modes operating at $MI \leq 0.4$. The probe was positioned to provide optimal visualization over the target lesion and was kept in the same position for at least 180 seconds.

Truth standard included: histology/surgery, contrast CT, contrast MRI, and/or 6 month follow-up.

For each study, the interpretation of images was conducted by three independent readers who were blinded to clinical data. Lesions were characterized as malignant or benign. Separate blinded readers assessed the truth standard images.

Results of both studies demonstrated an improvement in characterization of focal liver lesions using Lumason ultrasound compared to non-contrast ultrasound images. Table 5 summarizes the efficacy results by reader.

Table 5. Diagnostic Performance of Lumason Ultrasound for Characterization

of Focal Liver Lesions						
Study A:						
	Sensitivity (patients with malignant lesions) N=119			Specificity (patients with benign lesions) N=140		
	Lumason %	Non-contrast %	Difference (95% CI)	Lumason %	Non-contrast %	Difference (95% CI)
Reader 1	87*	49	38 (30, 54)	71	63	8 (-4, 21)
Reader 2	76*	35	41 (29, 52)	83*	54	29 (21, 44)
Reader 3	92*	16	76 (67, 84)	73*	22	51 (40, 61)
Study B:						
	Sensitivity (patients with malignant lesions) N=124			Specificity (patients with benign lesions) N=116		
	Lumason %	Non-contrast %	Difference (95% CI)	Lumason %	Non-contrast %	Difference (95% CI)
Reader 4	65	53	12 (-1, 23)	72*	24	48 (35, 58)
Reader 5	61*	41	20 (7, 32)	67*	7	60 (50, 70)
Reader 6	47	66	-19 (-31, -7)	88*	59	29 (18, 40)

* Statistically significant improvement from non-contrast (p<0.05 based on McNemar's test)

Pediatric patients

In one published study, 44 patients with an indeterminate focal liver lesion (23 males, 21 females, age range: 4-18 years, median 11.5 years) were evaluated after intravenous bolus administration of 1.2 to 2.4 mL of Lumason. The findings of Lumason ultrasound images were compared to CT, MRI or histology. Specificity was 98% (43/44 patients).

14.3 Ultrasonography of the Urinary Tract

Pediatric Patients

The efficacy of Lumason for the evaluation of pediatric patients with suspected or known vesicoureteral reflux was established in two published open-label single center studies (A and B). Patients received 1 mL of Lumason intravesically and underwent voiding urosonography (VUS). Patients were also evaluated with voiding cystourethrography (VCUG) as the reference standard. The presence or absence of urinary reflux with Lumason ultrasound was compared to the radiographic reference standard.

Study A evaluated 183 patients (94 male, 89 female; age 2 days - 44 months) with a

total of 366 kidney-ureter units. The images were interpreted by one on-site reader, blinded to the reference standard. Out of 103 reference standard-positive images, Lumason VUS was positive in 89 units and falsely negative in 14 units. In 263 units with negative reference standard, the Lumason ultrasonography was negative in 226 and falsely positive in 37.

Study B evaluated 228 patients (123 male, 105 female; age 6 days -13 years) with a total of 463 kidney-ureter units (some patients had more than 2 units). The images were interpreted independently by two on-site readers, blinded to the reference standard. Out of 71 reference standard positive images, Lumason ultrasonography was positive in 57 and falsely negative in 14. In 392 units with negative reference standard, Lumason ultrasonography was negative in 302 and falsely positive in 90.

16 HOW SUPPLIED/STORAGE AND HANDLING

16.1 How Supplied

Lumason (sulfur hexafluoride lipid-type A microspheres) for injectable suspension is supplied as a single patient use kit and as a 20-vial pack as follows:

- **5 single patient use kits** (NDC 0270-7099-16) with each kit containing:
 1. One Lumason vial of 25 mg lipid-type A white lyophilized powder with headspace fill of 60.7 mg of sulfur hexafluoride
 2. One prefilled syringe containing 5mL of 0.9% Sodium Chloride Injection, USP (Diluent)
 3. One Mini-SpikeEach kit is packaged in a clear plastic container.
- **20-vial pack** (NDC 0270-7097-07) containing:
 1. Twenty (20) Lumason vials of 25 mg lipid-type A white lyophilized powder with headspace fill of 60.7 mg of sulfur hexafluoride
 2. Twenty (20) Mini-Spikes
 3. Twenty (20) peel-off syringe labels

16.2 Storage and Handling

Store the kit at 25°C (77°F); excursions permitted to 15-30°C (59-86°F) [see USP Controlled Room Temperature].

17 PATIENT COUNSELING INFORMATION

Advise patients to inform their healthcare provider if they develop any symptoms of hypersensitivity after LUMASON administration including rash, wheezing, or shortness of breath.

Manufactured for:

Bracco Diagnostics Inc.

Princeton, NJ 08540

by

BRACCO Suisse SA

Plan-les-Ouates Geneve, Switzerland (Lumason lyophilized powder vial-25 mg lipid- type

A/60.7 mg sulfur hexafluoride gas)

Vetter Pharma-Fertigung GmbH & Co. KG
88212 Ravensburg, Germany (0.9% Sodium Chloride Injection, USP)
or

Bracco Imaging S.p.A.
Via Ribes, 5, 10010 Colletterto Giacosa (TO), Italy
(0.9% Sodium Chloride Injection, USP)


B. Braun Melsungen AG
34212 Melsungen, Germany (Mini-Spike)

This product is covered by US Patent No. 5,686,060

CL68D-11

Revised July 2024


Lumason - Box of 5 Kits
NDC: 0270-7099-16

**Lumason[®]**
(sulfur hexafluoride lipid-type A microspheres) for injectable suspension, 60.7 mg sulfur hexafluoride/25 mg lipid-type A per vial

NDC 0270-7099-16

Contents: 5 single use Lumason Kits with each kit containing:
1 vial of Lumason lyophilized powder for injectable suspension, 60.7 mg sulfur hexafluoride/25 mg lipid-type A
1 prefilled syringe containing 5 mL 0.9% Sodium Chloride Injection, USP for use during reconstitution (Diluent)
1 Mini-Spike

For intravenous or intravesical use only after reconstitution. See package insert for dosage, reconstitution and solution stability. Kit components intended for single use.
Each vial contains sterile lyophilized powder of lipid-type A blend consisting of 0.19 mg distereoylphosphatidylcholine, 0.19 mg dipalmitoylphosphatidylglycerol sodium, 24.56 mg polyethylene glycol 4000 and 0.04 mg palmitic acid. The headspace of the vial contains 60.7 mg sulfur hexafluoride.
Each mL of Diluent in prefilled syringe contains 9 mg of sodium chloride.
After vial reconstitution with 5 mL of diluent, each mL of Lumason contains 1.5-5.6 x 10⁶ microspheres, 68 mcg sulfur hexafluoride [45 mcg inside microspheres], 0.038 mg distereoylphosphatidylcholine, 0.038 mg dipalmitoylphosphatidylglycerol sodium, 4.1 mg polyethylene glycol 4000 and 0.008 mg palmitic acid. The pH of the reconstituted product is 4.5 to 7.5.
Store Lumason at 25°C (77°F); excursions permitted to 15° to 30°C (59° to 86°F) [see USP Controlled Room Temperature].
Manufactured for Bracco Diagnostics Inc., Princeton, NJ 08540
by Bracco Suisse SA, 31 Route de la Galaise, 1228 Plan-les-Ouates, Switzerland


(01)30302707099167(17)000000(10)000000L

Lot: Exp:

CE68C-05

Lumason - Kit Box Label
NDC: 0270-7099-16



Lumason®

NDC 0270-7099-16

(sulfur hexafluoride lipid-type A microspheres) for injectable suspension, 60.7 mg sulfur hexafluoride/25 mg lipid-type A per vial

Contents: 1 vial of Lumason lyophilized powder for injectable suspension, 60.7 mg sulfur hexafluoride/25 mg lipid-type A
1 prefilled syringe containing 5 mL 0.9% Sodium Chloride Injection, USP for use during reconstitution (Diluent)
1 Mini-Spike

For intravenous or intravesical use only after reconstitution. See package insert for dosage, reconstitution and solution stability. Kit components intended for single use. Each vial contains lyophilized powder of lipid-type A blend consisting of 0.19 mg distereoylphosphatidylcholine, 0.19 mg dipalmitoylphosphatidylglycerol sodium, 24.56 mg polyethylene glycol 4000 and 0.04 mg palmitic acid. The headspace of the vial contains 60.7 mg sulfur hexafluoride.

Each mL of Diluent in prefilled syringe contains 9 mg of sodium chloride.

Store Lumason at 25°C (77°F); excursions permitted to 15° to 30°C (59° to 86°F) [see USP Controlled Room Temperature].

Rx only

Manufactured for: Bracco Diagnostics Inc.,
Princeton, NJ 08540
by: Bracco Suisse SA, 31 Route de la Galaise,
1228 Plan-les-Ouates, Switzerland



(01)20302707099160

Exp:

Lot:

CF68B-07

Lumason Vial Label

Lumason®

NDC 0270-7099-16

(sulfur hexafluoride lipid-type A microspheres) lyophilized powder for injectable suspension, 60.7 mg sulfur hexafluoride/ 25 mg lipid-type A



Rx only

For intravenous or intravesical use after reconstitution with 0.9% Sodium Chloride Injection, USP. (Diluent)

Date & time of reconstitution _____

Single Patient Use Vial - Discard unused portion after 3 hours.

Store Lumason at 25°C (77°F); excursions permitted to 15° to 30°C (59° to 86°F) [see USP Controlled Room Temperature].

Distributed by Bracco Diagnostics Inc. - Princeton, NJ 08540



(01)20302707099160

C168A-06

Exp:

Lot:

Lumason 20 Pack Vial Label NDC 0270-7097-07

Lumason®

NDC 0270-7097-07

(sulfur hexafluoride lipid-type A microspheres) lyophilized powder for injectable suspension, 60.7 mg sulfur hexafluoride/ 25 mg lipid-type A



Rx only

For intravenous or intravesical use after reconstitution with 5 mL additive-free 0.9% Sodium Chloride Injection, USP.

Date & time of reconstitution _____

Single Patient Use Vial - Discard unused portion after 3 hours.

Store Lumason at 25°C (77°F); excursions permitted to 15° to 30°C (59° to 86°F) [see USP Controlled Room Temperature].

Distributed by Bracco Diagnostics Inc. - Princeton, NJ 08540



(01)20302707097074

C1171-03

Exp:

Lot:

Lumason 20 Pack Carton Label



Lumason[®]
(sulfur hexafluoride lipid-type A microspheres) for injectable
suspension, 60.7 mg sulfur hexafluoride/25 mg lipid-type A
per vial

Contents: Each Lumason 20-vial pack contains:

20 single patient use vials of Lumason lyophilized powder for injectable suspension,
 60.7 mg sulfur hexafluoride/25 mg lipid-type A
 20 Mini-Spikes
 20 labels

For intravenous or intravesical use only after reconstitution.

Recommended Dosage: See Prescribing Information.

Each vial contains sterile lyophilized powder of lipid-type A blend consisting of 0.19 mg distereoylphosphatidylcholine, 0.19 mg dipalmitoylphosphatidylglycerol sodium, 24.56 mg polyethylene glycol 4000 and 0.04 mg palmitic acid. The headspace of the vial contains 60.7 mg sulfur hexafluoride.

Reconstitution: Add 5 mL of additive free 0.9% Sodium Chloride Injection, USP to each single patient use vial.

Each mL of Lumason contains 1.5-5.6 x 10⁸ microspheres, 68 mcg sulfur hexafluoride [45 mcg inside microspheres], 0.038 mg distereoylphosphatidylcholine, 0.038 mg dipalmitoylphosphatidylglycerol sodium, 4.1 mg polyethylene glycol 4000 and 0.008 mg palmitic acid. The pH of the reconstituted product is 4.5 to 7.5.

Store Lumason at 25°C (77°F); excursions permitted to 15° to 30°C (59° to 86°F) [see USP Controlled Room Temperature].

Rx only

Manufactured for Bracco Diagnostics Inc.,
 Princeton, NJ 08540
 by Bracco Suisse SA, 31 Route de la Galaise,
 1228 Plan-les-Ouates, Switzerland

Lot:

Exp:



(01)30302707097071

CF1173-05

LUMASON

sulfur hexafluoride kit

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:0270-7099
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Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0270-7099-16	5 in 1 BOX	01/15/2015	
1		1 in 1 KIT; Type 1: Convenience Kit of Co-Package		

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 VIAL	25 mg
Part 2	1 SYRINGE	5 mL

Part 1 of 2

LUMASON

sulfur hexafluoride injection, powder, lyophilized, for suspension

Product Information

Route of Administration	INTRAVESICAL, INTRAVENOUS
-------------------------	---------------------------

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SULFUR HEXAFLUORIDE (UNII: WS7LR3I1D6) (SULFUR HEXAFLUORIDE - UNII:WS7LR3I1D6)	SULFUR HEXAFLUORIDE	60.7 mg in 1 mg
DISTEAROYLPHOSPHATIDYLCHOLINE, DL- (UNII: EAG959U971) (DISTEAROYLPHOSPHATIDYLCHOLINE, DL- - UNII:EAG959U971)	DISTEAROYLPHOSPHATIDYLCHOLINE, DL-	.19 mg in 1 mg
SODIUM 1,2-DIPALMITOYL-SN-GLYCERO-3-PHOSPHO-(1'-RAC-GLYCEROL) (UNII: 841B886EJ7) (SODIUM 1,2-DIPALMITOYL-SN-GLYCERO-3-PHOSPHO-(1'-RAC-GLYCEROL) - UNII:841B886EJ7)	SODIUM 1,2-DIPALMITOYL-SN-GLYCERO-3-PHOSPHO-(1'-RAC-GLYCEROL)	.19 mg in 1 mg

Inactive Ingredients

Ingredient Name	Strength
POLYETHYLENE GLYCOL 4000 (UNII: 4R4HFI6D95)	24.56 mg in 1 mg
PALMITIC ACID (UNII: 2V16EO95H1)	.04 mg in 1 mg

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		25 mg in 1 VIAL; Type 1: Convenience Kit of Co-Package		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA203684	01/15/2015	

Part 2 of 2

DILUENT

sodium chloride 0.9% injection,usp injection

Product Information

Route of Administration	INTRAVESICAL, INTRAVENOUS
-------------------------	---------------------------

Inactive Ingredients

Ingredient Name	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X)	.9 mg in 1 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		5 mL in 1 SYRINGE; Type 1: Convenience Kit of Co-Package		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA203684	01/15/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA203684	01/15/2015	

LUMASON

sulfur hexafluoride injection, powder, lyophilized, for suspension

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:0270-7097
Route of Administration	INTRAVESICAL, INTRAVENOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SULFUR HEXAFLUORIDE (UNII: WS7LR3I1D6) (SULFUR HEXAFLUORIDE - UNII:WS7LR3I1D6)	SULFUR HEXAFLUORIDE	60.7 mg in 1 mg
DISTEAROYLPHOSPHATIDYLCHOLINE, DL- (UNII: EAG959U971) (DISTEAROYLPHOSPHATIDYLCHOLINE, DL- - UNII:EAG959U971)	DISTEAROYLPHOSPHATIDYLCHOLINE, DL-	.19 mg in 1 mg

SODIUM 1,2-DIPALMITOYL-SN-GLYCERO-3-PHOSPHO-(1'-RAC-GLYCEROL) (UNII: 841B886EJ7) (SODIUM 1,2-DIPALMITOYL-SN-GLYCERO-3-PHOSPHO-(1'-RAC-GLYCEROL) - UNII:841B886EJ7)		SODIUM 1,2-DIPALMITOYL-SN-GLYCERO-3-PHOSPHO-(1'-RAC-GLYCEROL)	.19 mg in 1 mg	
Inactive Ingredients				
Ingredient Name			Strength	
POLYETHYLENE GLYCOL 4000 (UNII: 4R4HFI6D95)			24.56 mg in 1 mg	
PALMITIC ACID (UNII: 2V16EO95H1)			.04 mg in 1 mg	
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0270-7097-07	20 in 1 BOX	04/01/2021	
1		25 mg in 1 VIAL; Type 1: Convenience Kit of Co-Package		
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA		NDA203684	04/01/2021	

Labeler - BRACCO DIAGNOSTICS INC (849234661)

Registrant - BRACCO DIAGNOSTICS INC (849234661)

Establishment

Name	Address	ID/FEI	Business Operations
Bracco Suisse SA		485635705	MANUFACTURE(0270-7097, 0270-7099) , ANALYSIS(0270-7099, 0270-7097)

Establishment

Name	Address	ID/FEI	Business Operations
Vetter Pharma-Fertigung GmbH & Co. KG		316126754	MANUFACTURE(0270-7097, 0270-7099)

Establishment

Name	Address	ID/FEI	Business Operations
Bracco Imaging SPA		434384008	MANUFACTURE(0270-7097, 0270-7099) , ANALYSIS(0270-7099, 0270-7097)

Establishment

Name	Address	ID/FEI	Business Operations
Lipoid GmbH		312893951	API MANUFACTURE(0270-7097, 0270-7099)