

- 1 **TRISENOX™**
- 2 (arsenic trioxide) injection
- 3 For Intravenous Use Only
- 4 10 mg/10 mL (1 mg/mL) ampule

**Rx only**

## **WARNING**

**Experienced Physician and Institution:** TRISENOX™ (arsenic trioxide) injection should be administered under the supervision of a physician who is experienced in the management of patients with acute leukemia.

**APL Differentiation Syndrome:** Some patients with APL treated with TRISENOX™ have experienced symptoms similar to a syndrome called the retinoic-acid-Acute Promyelocytic Leukemia (RA-APL) or APL differentiation syndrome, characterized by fever, dyspnea, weight gain, pulmonary infiltrates and pleural or pericardial effusions, with or without leukocytosis. This syndrome can be fatal. The management of the syndrome has not been fully studied, but high-dose steroids have been used at the first suspicion of the APL differentiation syndrome and appear to mitigate signs and symptoms. At the first signs that could suggest the syndrome (unexplained fever, dyspnea and/or weight gain, abnormal chest auscultatory findings or radiographic abnormalities), high-dose steroids (dexamethasone 10 mg intravenously BID) should be immediately initiated, irrespective of the leukocyte count and continued for at least 3 days or longer until signs and symptoms have abated. The majority of patients do not require termination of TRISENOX™ therapy during treatment of the APL differentiation syndrome.

**ECG Abnormalities:** Arsenic trioxide can cause QT interval prolongation and complete atrioventricular block. QT prolongation can lead to a torsade de pointes-type ventricular arrhythmia, which can be fatal. The risk of torsade de pointes is related to the extent of QT prolongation, concomitant administration of QT prolonging drugs, a history of torsade de pointes, preexisting QT interval prolongation, congestive heart failure, administration of potassium-wasting diuretics, or other conditions that result in hypokalemia or hypomagnesemia. One patient (also receiving amphotericin B) had torsade de pointe during induction therapy for relapsed APL with arsenic trioxide.

**ECG and Electrolyte Monitoring Recommendations:** Prior to initiating therapy with TRISENOX™, a 12-lead ECG should be performed and serum electrolytes (potassium, calcium, and magnesium) and creatinine should be assessed; preexisting electrolyte abnormalities should be corrected and, if possible, drugs that are known to prolong the QT interval should be discontinued. For QTc greater than 500 msec, corrective measures should be completed and the QTc reassessed with serial ECGs prior to considering using TRISENOX™. During therapy with TRISENOX™, potassium concentrations should be kept above 4 mEq/dL and magnesium concentrations should be kept above 1.8 mg/dL. Patients who reach an absolute QT interval value > 500 msec should be reassessed and immediate action should be taken to correct concomitant risk factors, if any, while the risk/benefit of continuing versus suspending TRISENOX™ therapy should be considered. If syncope, rapid

or irregular heartbeat develops, the patient should be hospitalized for monitoring, serum electrolytes should be assessed, TRISENOX™ therapy should be temporarily discontinued until the QTc interval regresses to below 460 msec, electrolyte abnormalities are corrected, and the syncope and irregular heartbeat cease. There are no data on the effect of TRISENOX™ on the QTc interval during the infusion.

## 5 DESCRIPTION

6 TRISENOX™ is a sterile injectable solution of arsenic trioxide. The molecular formula of the  
7 drug substance in the solid state is As<sub>2</sub>O<sub>3</sub>, with a molecular weight of 197.8 g.

8 TRISENOX™ is available in 10 mL, single-use ampules containing 10 mg of arsenic trioxide.  
9 TRISENOX™ is formulated as a sterile, nonpyrogenic, clear solution of arsenic trioxide in  
10 water-for-injection using sodium hydroxide and dilute hydrochloric acid to adjust to pH 8.  
11 TRISENOX™ is preservative-free. Arsenic trioxide, the active ingredient, is present at a  
12 concentration of 1.0 mg/mL. Inactive ingredients and their respective approximate  
13 concentrations are sodium hydroxide (1.2 mg/mL) and hydrochloric acid, which is used to  
14 adjust the pH to 7.0 - 9.0.

## 15 CLINICAL PHARMACOLOGY

### 16 Mechanism of Action

17 The mechanism of action of TRISENOX™ is not completely understood. Arsenic trioxide  
18 causes morphological changes and DNA fragmentation characteristic of apoptosis in NB4  
19 human promyelocytic leukemia cells *in vitro*. Arsenic trioxide also causes damage or  
20 degradation of the fusion protein PML-RAR alpha.

### 21 Pharmacokinetics

22 The pharmacokinetics of trivalent arsenic, the active species of TRISENOX™, has not been  
23 characterized.

### 24 Metabolism

25 The metabolism of arsenic trioxide involves reduction of pentavalent arsenic to trivalent arsenic  
26 by arsenate reductase and methylation of trivalent arsenic to monomethylarsonic acid and  
27 monomethylarsonic acid to dimethylarsinic acid by methyltransferases. The main site of  
28 methylation reactions appears to be the liver. Arsenic is stored mainly in liver, kidney, heart,  
29 lung, hair and nails.

### 30 Excretion

31 Disposition of arsenic following intravenous administration has not been studied. Trivalent  
32 arsenic is mostly methylated in humans and excreted in urine.

### 33 Special Populations

34 The effects of renal or hepatic impairment or gender, age and race on the pharmacokinetics of  
35 TRISENOX™ have not been studied (see PRECAUTIONS).

### 36 **Drug Interactions**

37 No formal assessments of pharmacokinetic drug-drug interactions between TRISENOX™ and  
38 other drugs have been conducted. The methyltransferases responsible for metabolizing arsenic  
39 trioxide are not members of the cytochrome P450 family of isoenzymes. (see  
40 PRECAUTIONS).

### 41 **Clinical Studies**

42 TRISENOX™ has been investigated in 40 relapsed or refractory APL patients, previously  
43 treated with an anthracycline and a retinoid regimen, in an open-label, single-arm, non-  
44 comparative study. Patients received 0.15 mg/kg/day intravenously over 1 to 2 hours until the  
45 bone marrow was cleared of leukemic cells or up to a maximum of 60 days. The CR (absence  
46 of visible leukemic cells in bone marrow and peripheral recovery of platelets and white blood  
47 cells with a confirmatory bone marrow  $\geq$  30 days later) rate in this population of previously  
48 treated patients was 28 of 40 (70%). Among the 22 patients who had relapsed less than one  
49 year after treatment with ATRA, there were 18 complete responders (82%). Of the 18 patients  
50 receiving TRISENOX™  $\geq$  one year from ATRA treatment, there were 10 complete  
51 responders (55%). The median time to bone marrow remission was 44 days and to onset of  
52 CR was 53 days. Three of 5 children 5 years or older achieved CR. No children less than 5  
53 years old were treated.

54 Three to six weeks following bone marrow remission, thirty-one patients received consolidation  
55 therapy with TRISENOX™, at the same dose, for 25 additional days over a period up to 5  
56 weeks. In follow-up treatment, eighteen patients received further arsenic trioxide as a  
57 maintenance course. Fifteen patients had bone marrow transplants. At last follow-up, 27 of 40  
58 patients were alive with a median follow-up time of 484 days (range 280 to 755) and 23 of 40  
59 patients remained in complete response with a median follow-up time of 483 days (range 280 to  
60 755).

61 Cytogenetic conversion to no detection of the APL chromosome rearrangement was observed  
62 in 24 of 28 (86%) patients who met the response criteria defined above, in 5 of 5 (100%)  
63 patients who met some but not all of the response criteria, and 3 of 7 (43%) of patients who did  
64 not respond. Reverse Transcriptase – Polymerase Chain Reaction conversions to no detection  
65 of the APL gene rearrangement were demonstrated in 22 of 28 (79%) of patients who met the  
66 response criteria, in 3 of 5 (60%) of patients who met some but not all of the response criteria,  
67 and in 2 of 7 (29%) of patients who did not respond.

68 Responses were seen across all age groups tested, ranging from 6 to 72 years. The ability to  
69 achieve a CR was similar for both genders. There were insufficient patients of black, Hispanic  
70 or Asian derivation to estimate relative response rates in these groups, but responses were seen  
71 in members of each group.

72 Another single center study in 12 patients with relapsed or refractory APL, where patients  
73 received TRISENOX™ doses generally similar to the recommended dose, had similar results  
74 with 9 of 12 (75%) patients attaining a CR.

## 75 **INDICATIONS**

76 TRISENOX™ is indicated for induction of remission and consolidation in patients with acute  
77 promyelocytic leukemia (APL) who are refractory to, or have relapsed from, retinoid and  
78 anthracycline chemotherapy, and whose APL is characterized by the presence of the t(15;17)  
79 translocation or PML/RAR-alpha gene expression.

80 The response rate of other acute myelogenous leukemia subtypes to TRISENOX™ has not  
81 been examined.

## 82 **CONTRAINDICATIONS**

83 TRISENOX™ is contraindicated in patients who are hypersensitive to arsenic.

## 84 **WARNINGS (see boxed WARNING)**

85 TRISENOX™ should be administered under the supervision of a physician who is experienced  
86 in the management of patients with acute leukemia.

87 **APL Differentiation Syndrome (see boxed WARNING):** Nine of 40 patients with APL  
88 treated with TRISENOX™, at a dose of 0.15 mg/kg, experienced the APL differentiation  
89 syndrome (see box WARNING and ADVERSE REACTIONS).

90 **Hyperleukocytosis:** Treatment with TRISENOX™ has been associated with the  
91 development of hyperleukocytosis ( $\geq 10 \times 10^3/\mu\text{L}$ ) in 20 of 40 patients. A relationship did not  
92 exist between baseline WBC counts and development of hyperleukocytosis nor baseline WBC  
93 counts and peak WBC counts. Hyperleukocytosis was not treated with additional  
94 chemotherapy. WBC counts during consolidation were not as high as during induction  
95 treatment.

96

97 **QT Prolongation (see boxed WARNING):** QT/QTc prolongation should be expected  
98 during treatment with arsenic trioxide and torsade de pointes as well as complete heart block  
99 has been reported. Over 460 ECG tracings from 40 patients with refractory or relapsed APL  
100 treated with TRISENOX™ were evaluated for QTc prolongation. Sixteen of 40 patients  
101 (40%) had at least one ECG tracing with a QTc interval greater than 500 msec. Prolongation of  
102 the QTc was observed between 1 and 5 weeks after TRISENOX™ infusion, and then returned  
103 towards baseline by the end of 8 weeks after TRISENOX™ infusion. In these ECG  
104 evaluations, women did not experience more pronounced QT prolongation than men, and there  
105 was no correlation with age.

106

107 *Complete AV block:* Complete AV block has been reported with arsenic trioxide in the  
108 published literature including a case of a patient with APL.  
109

110 **Carcinogenesis:** Carcinogenicity studies have not been conducted with TRISENOX™ by  
111 intravenous administration. The active ingredient of TRISENOX™, arsenic trioxide, is a human  
112 carcinogen.  
113

114 **Pregnancy:** TRISENOX™ may cause fetal harm when administered to a pregnant woman.  
115 Studies in pregnant mice, rats, hamsters, and primates have shown that inorganic arsenicals  
116 cross the placental barrier when given orally or by injection. The reproductive toxicity of  
117 arsenic trioxide has been studied in a limited manner. An increase in resorptions, neural-tube  
118 defects, anophthalmia and microphthalmia were observed in rats administered 10 mg/kg of  
119 arsenic trioxide on gestation day 9 (approximately 10 times the recommended human daily dose  
120 on a mg/m<sup>2</sup> basis). Similar findings occurred in mice administered a 10 mg/kg dose of a related  
121 trivalent arsenic, sodium arsenite, (approximately 5 times the projected human dose on a mg/m<sup>2</sup>  
122 basis) on gestation days 6, 7, 8 or 9. Intravenous injection of 2 mg/kg sodium arsenite  
123 (approximately equivalent to the projected human daily dose on a mg/m<sup>2</sup> basis) on gestation day  
124 7 (the lowest dose tested) resulted in neural-tube defects in hamsters.

125 There are no studies in pregnant women using TRISENOX™. If this drug is used during  
126 pregnancy or if the patient becomes pregnant while taking this drug, the patient should be  
127 apprised of the potential harm to the fetus. One patient who became pregnant while receiving  
128 arsenic trioxide had a miscarriage. Women of childbearing potential should be advised to avoid  
129 becoming pregnant.  
130

## 131 **PRECAUTIONS**

132

133 **Laboratory Tests:** The patient's electrolyte, hematologic and coagulation profiles should be  
134 monitored at least twice weekly, and more frequently for clinically unstable patients during the  
135 induction phase and at least weekly during the consolidation phase. ECGs should be obtained  
136 weekly, and more frequently for clinically unstable patients, during induction and consolidation.

137 **Drug Interactions:** No formal assessments of pharmacokinetic drug-drug interactions between  
138 TRISENOX™ and other agents have been conducted. Caution is advised when  
139 TRISENOX™ is coadministered with other medications that can prolong the QT interval (e.g.  
140 certain antiarrhythmics or thioridazine) or lead to electrolyte abnormalities (such as diuretics or  
141 amphotericin B).

142 **Carcinogenesis, Mutagenesis, Impairment of Fertility:** See WARNINGS section for  
143 information on carcinogenesis. Arsenic trioxide and trivalent arsenite salts have not been  
144 demonstrated to be mutagenic to bacteria, yeast or mammalian cells. Arsenite salts are  
145 clastogenic *in vitro* (human fibroblasts, human lymphocytes, Chinese hamster ovary cells,  
146 Chinese hamster V79 lung cells). Trivalent arsenic produced an increase in the incidence of

147 chromosome aberrations and micronuclei in bone marrow cells of mice. The effect of arsenic  
148 on fertility has not been adequately studied.

149 **Pregnancy:** Pregnancy Category D. See WARNINGS section.

150 **Nursing Mothers:** Arsenic is excreted in human milk. Because of the potential for serious  
151 adverse reactions in nursing infants from TRISENOX™, a decision should be made whether to  
152 discontinue nursing or to discontinue the drug, taking into account the importance of the drug to  
153 the mother.

154 **Pediatric Use:** There are limited clinical data on the pediatric use of TRISENOX™. Of 5  
155 patients below the age of 18 years (age range: 5 to 16 years) treated with TRISENOX™, at  
156 the recommended dose of 0.15 mg/kg/day, 3 achieved a complete response.

157 Safety and effectiveness in pediatric patients below the age of 5 years have not been studied.

158 **Patients with Renal or Hepatic Impairment:** Safety and effectiveness of TRISENOX™ in  
159 patients with renal and hepatic impairment have not been studied. Particular caution is needed  
160 in patients with renal failure receiving TRISENOX™, as renal excretion is the main route of  
161 elimination of arsenic.

## 162 **ADVERSE REACTIONS**

163 Safety information was available for 52 patients with relapsed or refractory APL who  
164 participated in clinical trials of TRISENOX™. Forty patients in the Phase 2 study received the  
165 recommended dose of 0.15 mg/kg of which 29 completed both induction and consolidation  
166 treatment cycles. An additional 12 patients with relapsed or refractory APL received doses  
167 generally similar to the recommended dose. Most patients experienced some drug-related  
168 toxicity, most commonly leukocytosis, gastrointestinal (nausea, vomiting, diarrhea, and  
169 abdominal pain), fatigue, edema, hyperglycemia, dyspnea, cough, rash or itching, headaches,  
170 and dizziness. These adverse effects have not been observed to be permanent or irreversible  
171 nor do they usually require interruption of therapy.

172 Serious adverse events (SAEs), grade 3 or 4 according to version 2 of the NCI Common  
173 Toxicity Criteria, were common. Those SAEs attributed to TRISENOX™ in the Phase 2 study  
174 of 40 patients with refractory or relapsed APL included APL differentiation syndrome (n=3),  
175 hyperleukocytosis (n=3), QTc interval ≥ 500 msec (n=16, 1 with torsade de pointes), atrial  
176 dysrhythmias (n=2), and hyperglycemia (n=2).

177 The following table describes the adverse events that were observed in patients treated for APL  
178 with TRISENOX™ at the recommended dose at a rate of 5% or more. Similar adverse event  
179 profiles were seen in the other patient populations who received TRISENOX™ .

**Adverse Events (any grade) Occurring in ≥5% of 40 Patients with APL who Received  
TRISENOX™ at a dose of 0.15 mg/kg/day**

System organ class / Adverse Event	All Adverse Events, Any Grade		Grade 3 & 4 Events	
	n	%	n	%
<b>General disorders and administration site conditions</b>				
Fatigue	25	63	2	5
Pyrexia (Fever)	25	63	2	5
Edema – non-specific	16	40		
Rigors	15	38		
Chest pain	10	25	2	5
Injection site pain	8	20		
Pain – non specific	6	15	1	3
Injection site erythema	5	13		
Injection site edema	4	10		
Weakness	4	10	2	5
Hemorrhage	3	8		
Weight gain	5	13		
Weight loss	3	8		
Drug hypersensitivity	2	5	1	3
<b>Gastrointestinal disorders</b>				
Nausea	30	75		
Anorexia	9	23		
Appetite decreased	6	15		
Diarrhea	21	53		
Vomiting	23	58		
Abdominal pain (lower & upper)	23	58	4	10
Sore throat	14	40		
Constipation	11	28	1	3
Loose stools	4	10		
Dyspepsia	4	10		
Oral blistering	3	8		
Fecal incontinence	3	8		
Gastrointestinal hemorrhage	3	8		
Dry mouth	3	8		
Abdominal tenderness	3	8		
Diarrhea hemorrhagic	3	8		
Abdominal distension	3	8		
<b>Metabolism and nutrition disorders</b>				
Hypokalemia	20	50	5	13
Hypomagnesemia	18	45	5	13
Hyperglycemia	18	45	5	13

**Adverse Events (any grade) Occurring in ≥5% of 40 Patients with APL who Received  
TRISENOX™ at a dose of 0.15 mg/kg/day**

System organ class / Adverse Event	All Adverse Events, Any Grade		Grade 3 & 4 Events	
	n	%	n	%
ALT increased	8	20	2	5
Hyperkalemia	7	18	2	5
AST increased	5	13	1	3
Hypocalcemia	4	10		
Hypoglycemia	3	8		
Acidosis	2	5		
<b>Nervous system disorders</b>				
Headache	24	60	1	3
Insomnia	17	43	1	3
Parasthesia	13	33	2	5
Dizziness (excluding vertigo)	9	23		
Tremor	5	13		
Convulsion	3	8	2	5
Somnolence	3	8		
Coma	2	5	2	5
<b>Respiratory</b>				
Cough	26	65		
Dyspnea	21	53	4	10
Epistaxis	10	25		
Hypoxia	9	23	4	10
Pleural effusion	8	20	1	3
Post nasal drip	5	13		
Wheezing	5	13		
Decreased breath sounds	4	10		
Crackles	4	10		
Rales	4	10		
Hemoptysis	3	8		
Tachypnea	3	8		
Rhonchi	3	8		
<b>Skin &amp; subcutaneous tissue disorders</b>				
Dermatitis	17	43		
Pruritus	13	33	1	2
Ecchymosis	8	20		
Dry Skin	6	13		
Erythema- non-specific	5	10		
Increased sweating	5	10		
Facial edema	3	8		
Night sweats	3	8		
Petechiae	3	8		



**Adverse Events (any grade) Occurring in ≥5% of 40 Patients with APL who Received  
TRISENOX™ at a dose of 0.15 mg/kg/day**

System organ class / Adverse Event	All Adverse Events, Any Grade		Grade 3 & 4 Events	
	n	%	n	%
Hyperpigmentation	3	8		
Non specific skin lesions	3	8		
Urticaria	3	8		
Local exfoliation	2	5		
Eyelid edema	2	5		
<b>Cardiac disorders</b>				
Tachycardia	22	55		
ECG QT corrected interval prolonged > 500msec	16	38		
Palpitations	4	10		
ECG abnormal other than QT interval prolongation	3	7		
<b>Infections and infestations</b>				
Sinusitis	8	20		
Herpes simplex	5	13		
Upper respiratory tract infection	5	13	1	3
Bacterial infection- non-specific	3	8	1	3
Herpes zoster	3	8		
Nasopharyngitis	2	5		
Oral candidiasis	2	5		
Sepsis	2	5	2	5
<b>Musculoskeletal, connective tissue and bone disorders</b>				
Arthralgia	13	33	3	8
Myalgia	10	25	2	5
Bone pain	9	23	4	10
Back pain	7	18	1	3
Neck Pain	5	13		
Pain in limb	5	13	2	5
<b>Hematologic Disorders</b>				
Leukocytosis	20	50	1	3
Anemia	8	14	2	5
Thrombocytopenia	7	19	5	12
Febrile neutropenia	5	13	3	8
Neutropenia	4	10	4	10
Disseminated intravascular coagulation	3	8	3	8
Lymphadenopathy	3	8		
<b>Vascular disorders</b>				
Hypotension	10	25	2	5

**Adverse Events (any grade) Occurring in  $\geq$ 5% of 40 Patients with APL who Received  
TRISENOX™ at a dose of 0.15 mg/kg/day**

System organ class / Adverse Event	All Adverse Events, Any Grade		Grade 3 & 4 Events	
	n	%	n	%
Flushing	4	10		
Hypertension	4	10		
Pallor	4	10		
<b>Psychiatric Disorders</b>				
Anxiety	12	30		
Depression	8	20		
Agitation	2	5		
Confusion	2	5		
<b>Ocular Disorders</b>				
Eye irritation	4	10		
Blurred vision	4	10		
Dry eye	3	8		
Painful red eye	2	5		
<b>Renal and Urinary Disorders</b>				
Renal failure	3	8	1	3
Renal impairment	3	8		
Oliguria	2	5		
Incontinence	2	5		
<b>Reproductive System Disorders</b>				
Vaginal hemorrhage	5	13		
Intermenstrual bleeding	3	8		
<b>Ear Disorders</b>				
Earache	3	8		
Tinnitus	2	5		

181

182 **OVERDOSAGE**

183 If symptoms suggestive of serious acute arsenic toxicity (e.g., convulsions, muscle weakness and  
184 confusion) appear, TRISENOX™ should be immediately discontinued and chelation therapy  
185 should be considered. A conventional protocol for acute arsenic intoxication includes  
186 dimercaprol administered at a dose of 3 mg/kg intramuscularly every 4 hours until immediate  
187 life-threatening toxicity has subsided. Thereafter, penicillamine at a dose of 250 mg orally, up to  
188 a maximum frequency of four times per day ( $\leq$  1 gm per day), may be given.

189

190 **DOSAGE AND ADMINISTRATION**

191 TRISENOX™ should be diluted with 100 to 250mL 5% dextrose injection, USP or 0.9%  
192 Sodium Chloride injection, USP, using proper aseptic technique, immediately after withdrawal  
193 from the ampule. The TRISENOX™ ampule is single-use and does not contain any  
194 preservatives. Unused portions of each ampule should be discarded properly. Do not save any  
195 unused portions for later administration. Do not mix TRISENOX™ with other medications.

196 TRISENOX™ should be administered intravenously over 1-2 hours. The infusion duration  
197 may be extended up to 4 hours if acute vasomotor reactions are observed. A central venous  
198 catheter is not required.

### 199 **Stability**

200 After dilution, TRISENOX™ is chemically and physically stable when stored for 24 hours at  
201 room temperature and 48 hours when refrigerated.

### 202 **Dosing Regimen**

203 TRISENOX™ is recommended to be given according to the following schedule:

204 **Induction Treatment Schedule:** TRISENOX™ should be administered intravenously at a  
205 dose of 0.15 mg/kg daily until bone marrow remission. Total induction dose should not exceed  
206 60 doses.

207 **Consolidation Treatment Schedule:** Consolidation treatment should begin 3 to 6 weeks after  
208 completion of induction therapy. TRISENOX™ should be administered intravenously at a  
209 dose of 0.15 mg/kg daily for 25 doses over a period up to 5 weeks.

### 210 **HANDLING AND DISPOSAL**

211 Procedures for proper handling and disposal of anticancer drugs should be considered. Several  
212 guidelines on this subject have been published.<sup>1-7</sup> There is no general agreement that all of the  
213 procedures recommended in the guidelines are necessary or appropriate.

### 214 **HOW SUPPLIED**

215 TRISENOX™ (arsenic trioxide) injection is supplied as a sterile, clear, colorless solution in 10  
216 mL glass, single use ampules.

217 **NDC 60553-111-10** 10 mg/10 mL (1 mg/mL) ampule in packages of ten ampules.

218 Store at 25°C (77°F); excursions permitted to 15 - 30°C (59 - 86°F). Do not freeze.

219 Do not use beyond expiration date printed on the label.

220 **REFERENCES**

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239 **Rx only**

240 For additional information, contact Cell Therapeutics, Inc.  
241 Professional Services at 1-800-715-0944  
242 Customer Service at 1-888-305-2289.

243 **Manufactured for:**

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