HIGHLIGHTS OF PRESCRIBING INFORMATION

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

ELEVIDYS (delandistrogene moxeparvovec-rokl) suspension, for intravenous infusion
Initial U.S. Approval: 2023

WARNING: ACUTE SERIOUS LIVER INJURY AND ACUTE LIVER FAILURE

See full prescribing information for complete boxed warning.

- **Acute serious liver injury, including life-threatening and fatal acute liver failure have occurred with ELEVIDYS. (5.1)**
- **Patients with preexisting liver impairment may be at higher risk. (5.1)**
- **Prior to infusion, assess liver function by clinical examination and laboratory testing. Administer systemic corticosteroids before and after ELEVIDYS infusion. Continue to monitor liver function weekly for the first 3 months after infusion and continue until results are unremarkable. (2.1, 2.2, 2.4)**
- **Instruct patients to maintain proximity to an appropriate healthcare facility, as determined by the healthcare provider, for at least 2 months following ELEVIDYS infusion. (2.1)**
- **Obtain prompt consultation with a specialist (e.g., gastroenterologist or hepatologist) if acute serious liver injury or impending acute liver failure is suspected. (2.2, 5.1)**

----------RECENT MAJOR CHANGES----------

Boxed Warning	11/2025
Indication and Usage (1)	11/2025
Dosage and Administration (2)	11/2025
Contraindications (4)	11/2025
Warnings and Precautions (5)	11/2025

----------INDICATIONS AND USAGE----------

ELEVIDYS is an adeno-associated virus vector-based gene therapy indicated for the treatment of patients 4 years of age and older with Duchenne muscular dystrophy (DMD) who are ambulatory and have a confirmed mutation in the *DMD* gene. (1,12.2,14)

Limitations of Use:

ELEVIDYS is not recommended in patients with:

- Preexisting liver impairment (defined as gamma-glutamyl transferase [GGT] > 2 x upper limit of normal or total bilirubin > the upper limit of normal not due to Gilbert's syndrome) or active hepatic viral infection due to the high risk of acute serious liver injury and acute liver failure.
- Recent vaccination (within 4 weeks of treatment) due to immunogenicity and potential safety concerns.
- Active or recent (within 4 weeks) infections due to safety concerns.

----------DOSAGE AND ADMINISTRATION----------

ELEVIDYS is for single-dose intravenous infusion only.

- Select patients for treatment with ELEVIDYS with anti-AAVrh74 total binding antibody titers <1:400. (2.1)
- Postpone in patients with active or recent (within 4 weeks) infections. (2.1)
- Assess liver function, platelet counts and troponin-I before ELEVIDYS infusion. (2.1)
- Recommended dosage: 10 to 70 kg: 1.33×10^{14} vector genomes (vg) per kg of body weight; 70 kg or greater: 9.31×10^{15} vg. (2.2)
- One day prior to infusion, initiate a corticosteroid regimen for a minimum of 60 days. Recommend modifying corticosteroid dose for patients with liver function abnormalities. (2.2)
- Administer as an intravenous infusion over 1-2 hours. Infuse at a rate of less than 10 mL/kg/hour. (2.4)

----------DOSAGE FORMS AND STRENGTHS----------

- ELEVIDYS is a suspension for intravenous infusion with a nominal concentration of 1.33×10^{13} vg/mL. (3)
- ELEVIDYS is provided in a customized kit containing ten to seventy 10 mL single-dose vials, with each kit constituting a dosage unit based on the patient's body weight. (3)

----------CONTRAINDICATIONS----------

- ELEVIDYS is contraindicated in patients with any deletion in exon 8 and/or exon 9, including a deletion of any portion or the entirety of these exons, in the *DMD* gene. (4)

----------WARNINGS AND PRECAUTIONS----------

- Serious Infections: Serious infections with fatal outcomes may occur due to concomitant administration of corticosteroids, additional immunosuppressants, and ELEVIDYS. Monitor patients for signs and symptoms of infection; treat appropriately. (5.2)
- Myocarditis: Acute, serious, life-threatening myocarditis and troponin-I elevations have been observed. Monitor troponin-I before ELEVIDYS infusion, and weekly for the first month after ELEVIDYS infusion. (5.3)
- Infusion-related Reactions: Infusion-related reactions, including hypersensitivity reactions and anaphylaxis, have occurred. Monitor during administration and for at least 3 hours after end of infusion. If symptoms occur, slow or stop the infusion and give appropriate treatment. Once symptoms resolve, restart infusion at a slower infusion rate. Discontinue infusion for anaphylaxis. (2.4, 5.4)
- Immune-mediated Myositis: Severe to life-threatening immune-mediated myositis has been reported with ELEVIDYS in patients with deletions including portions of exons 1 to 17 and /or exons 59 to 71 of the *DMD* gene. Consider additional immunomodulatory treatment if symptoms of myositis occur (e.g., unexplained increased muscle pain, tenderness, or weakness). (5.5)
- Pre-existing Immunity against AAVrh74: Perform baseline testing for presence of anti-AAVrh74 total binding antibodies prior to ELEVIDYS administration. (5.6)

----------ADVERSE REACTIONS----------

Most common adverse reactions (incidence ≥5%) were vomiting and nausea, liver injury, pyrexia, thrombocytopenia, and troponin-I increased. (6.1)

To report SUSPECTED ADVERSE REACTIONS, contact Sarepta Therapeutics, Inc., at 1-888-SAREPTA (1-888-727-3782) or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

See 17 for PATIENT COUNSELING INFORMATION and Medication Guide.

Revised: 11/2025

FULL PRESCRIBING INFORMATION: CONTENTS*

* Sections or subsections omitted from the full prescribing information are not listed.

FULL PRESCRIBING INFORMATION

WARNING: ACUTE SERIOUS LIVER INJURY AND ACUTE LIVER FAILURE

- **Acute serious liver injury, including life-threatening and fatal acute liver failure have occurred with ELEVIDYS *[see Warnings and Precautions (5.1)].***
- **Patients with preexisting liver impairment may be at higher risk *[see Warnings and Precautions (5.1)].***
- **Prior to infusion, assess liver function by clinical examination and laboratory testing. Administer systemic corticosteroids before and after ELEVIDYS infusion. Continue to monitor liver function weekly for the first 3 months after infusion and continue until results are unremarkable *[see Dosage and Administration (2.1, 2.2, 2.4)].***
- **Instruct patients to maintain proximity to an appropriate healthcare facility, as determined by the healthcare provider, for at least 2 months following ELEVIDYS infusion *[see Dosage and Administration (2.1)].***
- **Obtain prompt consultation with a specialist (e.g., gastroenterologist or hepatologist) if acute serious liver injury or impending acute liver failure is suspected *[see Dosage and Administration (2.2), Warnings and Precautions (5.1)].***

1 INDICATIONS AND USAGE

ELEVIDYS is indicated for the treatment of patients 4 years of age and older with Duchenne muscular dystrophy (DMD), who are ambulatory and have a confirmed mutation in the *DMD* gene *[see Clinical Pharmacology (12.2), Clinical Studies (14)]*.

Limitations of Use:

ELEVIDYS is not recommended in patients with:

- Preexisting liver impairment (defined as gamma-glutamyl transferase [GGT] > 2 x upper limit of normal or total bilirubin > the upper limit of normal not due to Gilbert's syndrome) or active hepatic viral infection due to the high risk of acute serious liver injury and acute liver failure.
- Recent vaccination (within 4 weeks of treatment) due to immunogenicity and potential safety concerns.
- Active or recent (within 4 weeks) infections due to safety concerns.

2 DOSAGE AND ADMINISTRATION

For single-dose intravenous infusion only.

2.1 Critical Dosing Information

- Instruct patients to maintain proximity to an appropriate healthcare facility, as determined by the healthcare provider, for at least 2 months following ELEVIDYS infusion.
- Prior to ELEVIDYS infusion:
  - Select patients for treatment with ELEVIDYS with anti-AAVrh74 total binding antibody titers <1:400. An FDA-authorized test for the detection of anti-AAVrh74 total binding antibodies is not currently available. Currently available tests may vary in accuracy and design.
  - Avoid ELEVIDYS administration in patients with elevated anti-AAVrh74 total binding antibody titers (≥1:400) *[Clinical Pharmacology (12.6)]*.

- Due to the increased risk of serious systemic immune response, postpone ELEVIDYS in patients with active or recent (within 4 weeks) infections *[see Warnings and Precautions (5.2)].*

- Assess liver function (clinical examination and laboratory testing including aspartate aminotransferase (AST), alanine aminotransferase (ALT), gamma-glutamyl transferase (GGT), albumin, activated partial thromboplastin time (aPTT), international normalized ratio (INR), and total bilirubin) *[see Dosage and Administration (2.4), Warnings and Precautions (5.1), Use in Specific Populations (8.6)]*.

- Obtain platelet count and troponin-I levels *[see Dosage and Administration (2.4), Warnings and Precautions (5.3)]*.

- Do not re-administer ELEVIDYS.

2.2 Recommended Dose

The recommended dose of ELEVIDYS is 1.33×10^{14} vector genomes per kilogram (vg/kg) of body weight (or 10 mL/kg body weight) for patients weighing less than 70 kg or 9.31×10^{15} vg total fixed dose for patients weighing 70 kg or greater.

For the number of vials required, refer to Table 10 *[see How Supplied/Storage and Handling (16.1)]*.

Calculate the dose as follows:

ELEVIDYS dose (in mL) = patient body weight (rounded to the nearest kilogram) x 10

The multiplication factor 10 represents the per kilogram dose (1.33×10^{14} vg/kg) divided by the amount of vector genome copies per mL of the ELEVIDYS suspension (1.33×10^{13} vg/mL).

Number of ELEVIDYS vials needed = ELEVIDYS dose (in mL) divided by 10.

Example: Calculation of volume needed for a 19.5 kg patient
19.5 kg rounded to the nearest kilogram = 20 kg
20 kg × 10 = 200 mL

Number of ELEVIDYS vials needed = 200 divided by 10, rounded to the nearest number of vials = 20 vials

Administer corticosteroids to reduce the risk of immune responses to the AAVrh74 vector after administration of ELEVIDYS *[see Clinical Pharmacology (12.6)]*. Start corticosteroids 1 day prior to ELEVIDYS infusion based on the schedule outlined in Table 1 below. Continue this regimen for a minimum of 60 days after the infusion, unless earlier tapering is clinically indicated.

Table 1: Recommended pre- and post-infusion oral corticosteroid dosing

Baseline corticosteroid dosing [a]	Peri-ELEVIDYS infusion corticosteroid dose (prednisone equivalent) [b]	Recommended maximum total daily oral dose (prednisone equivalent) [b]
Daily or intermittent dose	Start 1 day prior to infusion: 1 mg/kg/day (and continue baseline dose)	60 mg/day
High dose for 2 days per week	Start 1 day prior to infusion: 1 mg/kg/day taken on days without high-dose corticosteroid treatment (and continue baseline dose)	60 mg/day
Not on corticosteroids	Start 1 week prior to infusion: 1.5 mg/kg/day	60 mg/day

[a] Patient continues to receive this dose

[b] Corticosteroids other than prednisone and prednisolone have not been studied for use as a peri-ELEVIDYS infusion corticosteroid

Modify oral corticosteroid doses according to Table 2 for patients with liver function abnormalities (e.g., GGT >= 3 times baseline, total bilirubin > ULN) following ELEVIDYS infusion. Consider IV bolus corticosteroids instead of oral corticosteroids for GGT or bilirubin elevations that do not respond after 1 week of increased oral corticosteroids. Consult with a specialist experienced in immunosuppressive therapy for additional interventions as needed.

Obtain prompt consultation with a specialist (e.g., gastroenterologist or hepatologist) if acute serious liver injury or impending acute liver failure is suspected.

Taper the additional peri-ELEVIDYS corticosteroids for patients previously taking corticosteroids at baseline back to baseline dose over 2 weeks, or longer as needed. Taper the peri-ELEVIDYS corticosteroids for patients not previously taking corticosteroids at baseline back to no corticosteroids over 4 weeks, or longer as needed. Do not stop corticosteroids abruptly.

Table 2: Recommended oral corticosteroid regimen dose modification for liver function abnormalities following ELEVIDYS infusion[a]

Peri-ELEVIDYS infusion corticosteroid dosing	Modified oral corticosteroid dose following ELEVIDYS infusion (prednisone equivalent)[b c]	Recommended maximum total daily oral dose (prednisone equivalent)[b c]
Baseline + 1 mg/kg/day	Increase to 2 mg/kg/day (and continue baseline dose)	120 mg/day
Baseline + 1 mg/kg/day taken on days without high-dose corticosteroid treatment	Increase to 2 mg/kg/day taken on days without high-dose corticosteroid treatment (and continue baseline dose)	120 mg/day
1.5 mg/kg/day	Increase from 1.5 mg/kg/day to 2.5 mg/kg/day	120 mg/day

[a] GGT >= 3 times baseline and/or other clinically significant liver function abnormalities (e.g., total bilirubin > ULN) following infusion.

[b] Consider IV bolus corticosteroids instead of oral corticosteroids for GGT or bilirubin elevations that do not respond after 1 week of increased oral corticosteroids. Consult with a specialist experienced in immunosuppressive therapy for additional interventions as needed.

[c] Corticosteroids other than prednisone and prednisolone have not been studied for use as a peri-ELEVIDYS infusion corticosteroid.

2.3 Preparation

General precautions

- Prepare ELEVIDYS using aseptic technique.
- Verify the required dose of ELEVIDYS based on the patient's body weight.

- Confirm that the kit contains sufficient number of vials to prepare the ELEVIDYS infusion for the patient.
- Visually inspect parenteral drug products for particulate matter and discoloration prior to administration, whenever suspension and container permit. ELEVIDYS may contain white to off-white particles.

Recommended supplies and materials:

- 60 mL siliconized polypropylene syringes
- 21-gauge maximum or smaller stainless steel needles

Preparing ELEVIDYS infusion

1. Thaw ELEVIDYS before use.
    - When thawed in the refrigerator, ELEVIDYS vials are stable for up to 14 days in the refrigerator (2°C to 8°C [36º F to 46º F]) when stored in the upright position.
    - Frozen ELEVIDYS vials will thaw in approximately 2 hours when placed at room temperature (up to 25°C [77ºF]) when removed from original packaging.
    - Thawed ELEVIDYS is stable for up to 24 hours at room temperature (up to 25°C [77ºF]).
2. Inspect vials to ensure no ice crystals are present prior to preparation.
3. When thawed, swirl gently.
    - Do not shake.
    - Do not refreeze.
    - Do not place back in the refrigerator.
4. Visually inspect each vial of ELEVIDYS. ELEVIDYS is a clear, colorless liquid that may have some opalescence. ELEVIDYS may contain white to off-white particles.
    - Do not use if the suspension in the vials is cloudy or discolored.
5. Remove the plastic flip-off cap from the vials and disinfect the rubber stopper with a sterilizing agent (e.g., alcohol wipes).
6. Withdraw 10 mL of ELEVIDYS from each vial provided in the customized ELEVIDYS kit (refer to Table 10).
    - Do not use filter needles during preparation of ELEVIDYS.
    - Multiple syringes will be required to withdraw the required volume.
    - Remove air from the syringes and cap the syringes.
7. Maintain syringes at room temperature prior to and during administration.

2.4 Administration

Recommended supplies and materials:

- Syringe infusion pump
- 0.2-micron PES* in-line filter with a large surface area. To avoid the risk of occlusions, the use of smaller pediatric in-line filters (e.g., less than 10 cm^2 surface area) is not recommended.
- PVC* (non-DEHP*) IV infusion tubing, and polyurethane catheter

*PVC = Polyvinyl chloride, DEHP = Di(2-ethylhexyl) phthalate, PES = Polyether sulfone

Administer ELEVIDYS as a single-dose intravenous infusion through a peripheral venous catheter:

ELEVIDYS should be administered in a setting where treatment for infusion-related reactions is immediately available *[see Warnings and Precautions (5.4)]*. Do not infuse ELEVIDYS at a rate of 10 mL/kg/hour or faster.

Consider application of a topical anesthetic to the infusion site prior to administration of IV insertion.

Recommend inserting a back-up catheter.

1. Flush the intravenous access line with 0.9% Sodium Chloride Injection prior to the ELEVIDYS infusion at the same infusion rate.
2. Administer ELEVIDYS via intravenous infusion using a syringe infusion pump with an in-line 0.2-micron filter at a duration of approximately 1 to 2 hours, or longer at care team discretion, through a peripheral limb vein.
3. Infuse at a rate of less than 10 mL/kg/hour.
    - Do not administer ELEVIDYS as an intravenous push.
    - Do not infuse ELEVIDYS in the same intravenous access line with any other product.
    - Use ELEVIDYS within 12 hours after drawing into syringe. Discard the ELEVIDYS-containing syringe(s) if infusion of the drug has not been completed within the 12-hour timeframe.
4. In the event of an infusion-related reaction during administration *[see Warnings and Precautions (5.4)]:*
    - Slow or stop the infusion based on patient's clinical presentation.
    - Discontinue infusion for anaphylaxis.
    - Administer treatment as needed to manage infusion-related reaction.
    - ELEVIDYS infusion may be restarted at a lower rate after the infusion-related reaction has resolved at the discretion of the physician, based on severity of patient's clinical presentation.
    - If the ELEVIDYS infusion needs to be stopped and restarted, ELEVIDYS should be infused within 12 hours after drawing into the syringe *[see How Supplied/Storage and Handling (16.2)]*.
5. Flush the intravenous access line with 0.9% Sodium Chloride Injection after the ELEVIDYS infusion.
    - Discard unused ELEVIDYS *[see How Supplied/Storage and Handling (16.2)].*
    - Dispose of the needle and syringe *[see How Supplied/Storage and Handling (16.2)].*

Monitoring Post-ELEVIDYS Administration

- Assess liver function (clinical exam, AST, ALT, GGT, albumin, aPTT, INR, and total bilirubin) weekly for the first 3 months. Continue monitoring if clinically indicated, until results are unremarkable (e.g., normal clinical exam, GGT and total bilirubin levels return to near baseline levels) *[see Warnings and Precautions (5.1), Specific Populations (8.6)]*.
- Obtain platelet counts weekly for the first two weeks *[see Adverse Reactions (6.1)]*. Continue monitoring if clinically indicated.
- Measure troponin-I weekly for the first month *[see Warnings and Precautions (5.3)]*. Continue monitoring if clinically indicated, until results return to near baseline levels or stabilize.

3 DOSAGE FORMS AND STRENGTHS

ELEVIDYS is a preservative-free, sterile, clear, colorless liquid that may have some opalescence and may contain white to off-white particles.

ELEVIDYS is a suspension for intravenous infusion with a nominal concentration of 1.33×10^{13} vg/mL.

ELEVIDYS is provided in a customized kit containing ten to seventy 10 mL single-dose vials, with each kit constituting a dosage unit based on the patient's body weight *[see How Supplied/Storage and Handling (16.1)]*.

4 CONTRAINDICATIONS

ELEVIDYS is contraindicated in patients with any deletion in exon 8 and/or exon 9, including a deletion of any portion or the entirety of these exons, in the *DMD* gene *[see Warnings and Precautions (5.5)]*.

5 WARNINGS AND PRECAUTIONS

5.1 Acute Serious Liver Injury and Acute Liver Failure

Acute serious liver injury marked by elevations of liver enzymes (e.g., GGT, ALT) and total bilirubin and acute liver failure, has occurred with ELEVIDYS. Onset of the liver injury typically begins within 8 weeks after administration of ELEVIDYS. In non-ambulatory patients treated with ELEVIDYS, acute liver failure with fatal outcome has occurred in the clinical and post-marketing settings.

Life-threatening mesenteric vein thrombosis, complicated by bowel ischemia and necrosis, and portal hypertension has been reported following acute liver injury associated with ELEVIDYS in a non-ambulatory patient *[see Adverse Reactions (6.2)]*.

Patients with preexisting liver impairment, chronic hepatic condition or acute liver disease (e.g., acute hepatic viral infection) may be at higher risk of acute serious liver injury or acute liver failure. Postpone ELEVIDYS administration in patients with acute liver disease until resolved or controlled. Patients with hepatic impairment, acute liver disease, chronic hepatic condition or elevated GGT have not been studied in clinical trials with ELEVIDYS *[see Specific Populations (8.6)]*.

In clinical studies, liver function test increased (including increases in GGT, ALT, AST, or total bilirubin) was commonly reported typically within 8 weeks following ELEVIDYS infusion, with the majority of cases being asymptomatic *[see Adverse Reactions (6.1)]*. Most cases resolved spontaneously or with systemic corticosteroids and resolved without clinical sequelae within 2 months.

Prior to ELEVIDYS administration, perform liver enzyme test *[see Dosage and Administration (2.1)]*. Monitor liver function (clinical examination, AST, ALT, GGT, albumin, aPTT, INR, and total bilirubin) weekly for the first 3 months following ELEVIDYS infusion. Continue monitoring if clinically indicated, until results are unremarkable (e.g., normal clinical exam, GGT and total bilirubin levels return to near baseline levels) *[see Dosage and Administration (2.4)]*.

Systemic corticosteroid treatment is recommended for patients before and after ELEVIDYS infusion *[see Dosage and Administration (2.2)]*. Adjust corticosteroid regimen when indicated *[see Dosage and Administration (2.2)]*. Obtain prompt consultation with a specialist (e.g., gastroenterologist or hepatologist) if acute serious liver injury or impending acute liver failure is suspected.

5.2 Serious Infections

Increased susceptibility to serious infections may occur due to concomitant administration of corticosteroid regimen and additional immunosuppressants, and ELEVIDYS. Serious respiratory infections, including with fatal outcomes, have occurred in patients taking immunosuppressant corticosteroids required for ELEVIDYS administration *[see Adverse Reactions (6.2)]*.

Monitor patients for signs and symptoms of infection before and after ELEVIDYS administration and treat appropriately. Administer immunizations according to best clinical practices and immunization guidelines prior to initiation of the corticosteroid regimen required before ELEVIDYS infusion *[see Drug Interactions (7)]*.

Avoid administration of ELEVIDYS to patients with active infections.

5.3 Myocarditis

Acute, serious, life-threatening myocarditis and troponin-I elevations have been observed within 24 hours to more than 1 year following ELEVIDYS infusion *[see Adverse Reactions (6.1)*].

If a patient experiences myocarditis, those with pre-existing left ventricle ejection fraction (LVEF) impairment may be at higher risk of adverse outcomes. Patients with moderate to severe LVEF impairment have not been studied in clinical trials with ELEVIDYS.

Monitor troponin-I before ELEVIDYS infusion and weekly for the first month following infusion *[see Dosage and Administration (2.4)]*. Continue monitoring if clinically indicated, until results return to near baseline levels or stabilize. More frequent monitoring may be warranted in the presence of cardiac symptoms, such as chest pain or shortness of breath.

Advise patients to contact a physician immediately if they experience cardiac symptoms.

5.4 Infusion-related Reactions

Infusion-related reactions, including hypersensitivity reactions and anaphylaxis, have occurred during or up to several hours following ELEVIDYS administration. Closely monitor patients during and for at least 3 hours after the end of infusion for signs and symptoms of infusion-related reactions including tachycardia, tachypnea, lip swelling, difficulty breathing, nasal flaring, urticaria, flushing, lip pruritus, rash, cheilitis, vomiting, nausea, rigors and pyrexia.

ELEVIDYS should be administered in a setting where treatment for infusion-related reactions is immediately available.

In the event of an infusion-related reaction, administration of ELEVIDYS may be slowed or stopped based on the severity of the patient's clinical presentation. Administer treatment as needed to manage infusion-related reactions based on the severity of patient's signs and symptoms. *[see Dosage and Administration (2.4)]*. If the infusion was stopped, ELEVIDYS infusion may be restarted at a lower rate once patient's symptoms have resolved, at the discretion of the physician. Discontinue infusion for anaphylaxis.

5.5 Immune-mediated Myositis

Immune-mediated myositis, including serious and life-threatening events have occurred approximately 1 month following ELEVIDYS infusion *[see Adverse Reactions (6)*]. Signs and symptoms include severe muscle weakness, including dysphagia, dyspnea, dysphonia, and hypophonia.

These immune reactions may be due to a T-cell based response against specific regions of the micro-dystrophin transgenic protein. Severe to life-threatening immune-mediated myositis has been reported in patients with deletions including portions of exons 1-17 and/or exons 59-71 of the *DMD* gene. ELEVIDYS is contraindicated in patients with any deletion in exon 8 and/or exon 9, including a deletion of any portion or the entirety of these exons, in the *DMD* gene due to the increased risk for a severe immune-mediated myositis reaction *[see Contraindications (4)]*.

Regardless of genetic mutation, advise patients to contact a physician immediately if they experience any unexplained increased muscle pain, tenderness, or weakness, including dysphagia, dyspnea, dysphonia or hypophonia as these may be symptoms of myositis. Consider additional immunomodulatory treatment based on patient's clinical presentation and medical history if these symptoms occur.

5.6 Pre-existing Immunity against AAVrh74

In AAV-vector based gene therapies, preexisting anti-AAV antibodies may impede transgene expression at desired therapeutic levels. Following treatment with ELEVIDYS all patients developed anti-AAVrh74 antibodies. Perform baseline testing for the presence of anti-AAVrh74 total binding antibodies prior to ELEVIDYS administration *[see Dosage and Administration (2.1)]*.

ELEVIDYS administration is not recommended in patients with elevated anti-AAVrh74 total binding antibody titers (≥1:400).

6 ADVERSE REACTIONS

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.

The safety data described in this section reflect exposure to a one-time intravenous infusion of ELEVIDYS in 156 male patients with a confirmed mutation of the *DMD* gene in four clinical studies, including one completed open-label study, one ongoing open-label study, and two studies that included a double-blind, placebo-controlled period. Prior to ELEVIDYS infusion, patients in the ELEVIDYS treatment group had a mean age of 6.7 years (range: 3 to 20) and mean weight of 24.6 kg (range: 12.5 to 80.1). 144 patients received the recommended dose of 1.33×10^{14} vg/kg, and 12 received a lower dose. Table 3 below presents adverse reactions from these four clinical studies.

The most common adverse reactions (incidence ≥5%) across all studies are summarized in Table 3.

Adverse reactions were typically seen within the first 2 weeks (nausea, vomiting, thrombocytopenia, pyrexia), the first month (myocarditis, troponin-I increased) or within the first 2 months (immune-mediated myositis, liver injury). Vomiting may occur as early as on the day of the infusion.

Table 3. Adverse reactions (Incidence ≥5%) following treatment with ELEVIDYS in Clinical Studies

Adverse reactions	ELEVIDYS (N=156) %
Vomiting	65
Nausea	43
Liver injury[a]	40
Pyrexia	28
Thrombocytopenia[b c]	8
Troponin-I increased[d]	8

[a] Includes: AST increased, ALT increased, GGT increased, GLDH increased, GLDH level abnormal, Hepatotoxicity, Hepatic enzyme increased, Hypertransaminasemia, Liver function test increased, Liver injury, Transaminases increased, Blood bilirubin increased

[b] Includes: Thrombocytopenia, Platelet count decreased

[c] Transient, mild, asymptomatic decrease in platelet counts

[d] Includes: Troponin I increased, Troponin increased, Troponin I abnormal

In clinical trials, immune-mediated myositis was observed in 2 of 6 patients with deletion mutations involving exon 8 and/or 9 in the *DMD* gene *[see Contraindications (4), and Warnings and Precautions (5.5)]*.

In the double-blind, placebo-controlled trial, Study 3 Part 1, patients 4 to 7 years of age (N=125) received either ELEVIDYS (N=63) at the recommended dose of 1.33×10^{14} vg/kg or placebo (N=62). Table 4 below presents the most frequent adverse reactions from Study 3 Part 1.

Table 4. Adverse reactions occurring in ELEVIDYS-treated patients and at least twice more frequently than with placebo in Study 3 Part 1

Adverse reactions	ELEVIDYS (N=63) %	Placebo (N=62) %
Vomiting	64	19
Nausea	40	13

Liver injury[a]	41	8
Pyrexia	32	24
Thrombocytopenia[bc]	3	0

[a] Includes: AST increased, ALT increased, GGT increased, GLDH increased, GLDH level abnormal, Hepatotoxicity, Hepatic enzyme increased, Hypertransaminasemia, Liver function test increased, Liver injury, Transaminases increased.
[b] Includes: platelet count decreased, thrombocytopenia
[c] Transient, mild, asymptomatic decrease in platelet counts

6.2 Postmarketing Experience

The following adverse reactions have been identified during post-approval use of ELEVIDYS. 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.

Hepatobiliary Disorders: Acute liver injury, acute liver failure, including fatal outcome and life-threatening mesenteric vein thrombosis *[see Warnings and Precautions (5.1)]*

Infections and Infestations: Bacterial and viral respiratory infections, including fatal outcome *[see Warnings and Precautions (5.2)]*

Immune System Disorders: Infusion-related reactions, including hypersensitivity reactions and anaphylaxis, have occurred during or up to several hours following ELEVIDYS administration *[see Warnings and Precautions (5.4)]*.

Musculoskeletal and connective tissue disorders: Immune-mediated myositis *[see Warnings and Precautions (5.5)].*

7 DRUG INTERACTIONS

Prior to initiating the corticosteroid regimen required before ELEVIDYS administration, consider the patient's vaccination status. Patients should, if possible, be brought up to date with all immunizations in agreement with current immunization guidelines. Vaccinations should be completed at least 4 weeks prior to initiation of the corticosteroid regimen.

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary
ELEVIDYS is not intended for use in pregnant women.

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

8.2 Lactation

Risk Summary
There is no information available on the presence of ELEVIDYS in human milk, the effects on the breastfed infant, or the effects on milk production.

8.4 Pediatric Use

The safety and effectiveness of ELEVIDYS for the treatment of Duchenne muscular dystrophy has been established in pediatric patients at least 4 years of age with a confirmed mutation in the *DMD* gene. The use of ELEVIDYS in pediatric patients was supported by evidence from three adequate and well controlled clinical studies which included 144 pediatric patients aged 4 years of age and older *[see Adverse Reactions (6), Clinical Pharmacology (12.2), Clinical Studies (14)]*.

8.5 Geriatric Use

The safety and efficacy of ELEVIDYS in geriatric patients with DMD have not been studied.

8.6 Hepatic Impairment

The safety and efficacy of ELEVIDYS in patients with hepatic impairment or elevated GGT have not been studied.

Postpone ELEVIDYS administration in patients with acute liver disease until resolved or controlled. Treatment with ELEVIDYS should be carefully considered in patients with preexisting liver impairment or chronic hepatic viral infection. These patients may be at increased risk of acute serious liver injury or acute liver failure *[see Warnings and Precautions (5.1)]*.

In clinical trials, liver function test increase was commonly reported in patients following ELEVIDYS infusion *[see Warnings and Precautions (5.1), Adverse Reactions (6.1)]*.

11 DESCRIPTION

ELEVIDYS (delandistrogene moxeparvovec-rokl) is a recombinant gene therapy designed to deliver the gene encoding the ELEVIDYS micro-dystrophin protein. ELEVIDYS is a non-replicating, recombinant, adeno-associated virus serotype rh74 (AAVrh74) based vector containing the ELEVIDYS micro-dystrophin transgene under the control of the MHCK7 promoter. The genome within the ELEVIDYS AAVrh74 vector contains no viral genes and consequently is incapable of replication or reversion to a replicating form. The micro-dystrophin protein expressed by ELEVIDYS is a shortened version (138 kDa, compared to 427 kDa size of dystrophin expressed in normal muscle cells) that contains selected domains of dystrophin expressed in normal muscle cells.

ELEVIDYS is a preservative-free, sterile, clear, colorless liquid that may have some opalescence and may contain white to off-white particles. ELEVIDYS is a suspension for intravenous infusion with a nominal concentration of 1.33×10^{13} vg/mL and supplied in a single-dose 10 mL vial. Each vial contains an extractable volume of 10 mL and the following excipients: 200mM sodium chloride, 13 mM tromethamine HCl, 7 mM tromethamine, 1mM magnesium chloride, 0.001% poloxamer 188, with a pH of 8.0 ± 0.3.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

ELEVIDYS is the recombinant gene therapy product that is comprised of a non-replicating, recombinant, adeno-associated virus (AAV) serotype rh74 (AAVrh74) capsid and a single-stranded DNA expression cassette flanked by inverted terminal repeats (ITRs) derived from AAV2. The cassette contains: 1) an MHCK7 gene regulatory component comprising a creatine kinase 7 promoter and an α-myosin heavy chain enhancer, and 2) the DNA transgene encoding the engineered micro-dystrophin protein.

Vector/Capsid: Clinical and nonclinical studies have demonstrated AAVrh74 serotype transduction in skeletal muscle cells. Additionally, in nonclinical studies, AAVrh74 serotype transduction has been demonstrated in cardiac and diaphragm muscle cells.

Promoter: The MHCK7 promoter/enhancer drives transgene expression and has been shown in animal models to drive transgenic micro-dystrophin protein expression predominantly in skeletal muscle (including diaphragm) and cardiac muscle. In clinical studies, muscle biopsy analyses have confirmed micro-dystrophin expression in skeletal muscle.

Transgene: DMD is caused by a mutation in the *DMD* gene resulting in lack of functional dystrophin protein. ELEVIDYS carries a transgene encoding a micro-dystrophin protein consisting of selected domains of dystrophin expressed in normal muscle cells.

ELEVIDYS micro-dystrophin has been demonstrated to localize to the sarcolemma.

12.2 Pharmacodynamics

In 92 patients who received ELEVIDYS in clinical studies, micro-dystrophin protein expression from muscle biopsies (gastrocnemius or biceps brachii) was quantified by western blot and localized by immunofluorescence staining (fiber intensity and percentage micro-dystrophin).

Micro-dystrophin expression (expressed as change from baseline) in ELEVIDYS-treated patients as measured by western blot was the primary objective of Study 1 and Study 2, and a key secondary objective for Study 3. Muscle biopsies were obtained at baseline prior to ELEVIDYS infusion and at Week 12 after ELEVIDYS infusion in all patients. The absolute quantity of micro-dystrophin was measured by western blot assay, adjusted by muscle content and expressed as a percent of control (levels of wild-type dystrophin in patients without DMD or Becker muscular dystrophy) in muscle biopsy samples. Study 1 and 2 results of patients receiving 1.33×10^{14} vg/kg ELEVIDYS are presented in Table 5.

Table 5. Micro-Dystrophin Expression in Study 1 and Study 2 at Week 12 from Baseline (Western Blot Assay)[abc]

Western blot (% of micro-dystrophin compared to control)	Study 1 Part 1 (n=6)	Study 1 Part 2 (n=21)	Study 2 Ambulatory (n=40)
Mean change from baseline (SD)	43.4 (48.6)	40.7 (32.3)	51.0 (47)
Median change from baseline (Min, Max)	24.3 (1.6, 116.3)	40.8 (0.0, 92.0)	46.9 (1.9, 197.3)

[a] All patients received 1.33 x 10^{14} vg/kg, as measured by ddPCR
[b] Change from baseline was statistically significant
[c] Adjusted for muscle content. Control was level of wild-type (normal) dystrophin in normal muscle.

In Study 3 Part 1, muscle biopsies were obtained at Week 12 in 31 patients. For the ELEVIDYS-treated patients, the mean micro-dystrophin expression at Week 12 was 34.3% (N=17, SD: 41.0%), compared to placebo patients of 0% (N=14, SD: 0%).

Assessment of micro-dystrophin levels can be meaningfully influenced by differences in sample processing, analytical technique, reference materials, and quantitation methodologies. Therefore, valid comparisons of micro-dystrophin measurements obtained from different assays cannot be made.

12.3 Pharmacokinetics

Vector Distribution and Vector Shedding

Nonclinical Data

Biodistribution of ELEVIDYS was evaluated in tissue samples collected from healthy mice and DMD^{mdx} mice following intravenous administration in toxicology studies. At 12 weeks following ELEVIDYS administration at dose levels of 1.33×10^{14} to 4.02×10^{14} vg/kg, vector DNA was detected in all major organs with the highest quantities detected in the liver, followed by lower levels in the heart, adrenal glands, skeletal muscle, and aorta. ELEVIDYS was also detected at low levels in the spinal cord, sciatic nerve and gonads (testis). Protein expression of micro-dystrophin was highest in cardiac tissue, exceeding physiologic dystrophin expression

levels in healthy mice, with lower levels in the skeletal muscle and diaphragm. In some studies, micro-dystrophin was also detected at low levels in the liver.

Clinical Data

Following IV administration, ELEVIDYS vector genome undergoes distribution via systemic circulation and distributes into target muscle tissues followed by elimination in the urine and feces. ELEVIDYS biodistribution and tissue transduction are detected in the target muscle tissue groups and quantified in the gastrocnemius or biceps femoris biopsies obtained from patients with mutations in the *DMD* gene. Evaluation of ELEVIDYS vector genome exposure in clinical muscle biopsies at Week 12 post-dose expressed as copies per nucleus revealed ELEVIDYS drug distribution and transduction with a mean change from baseline of 2.91 and 3.44 copies per nucleus at the recommended dose of 1.33×10^{14} vg/kg for Study 1 and Study 2 Cohort 1, respectively.

In Study 2 Cohorts 1-3, the biodistribution and vector shedding of ELEVIDYS in the serum and excreta were quantified, respectively. The mean maximum concentration (C_{max}) in the serum was 0.0055×10^{13} copies/mL and 2.78×10^{6} copies/mL in the urine, 7.86×10^{7} copies/mL in the saliva, and 4.87×10^{7} copies/µg in the feces. The median time to achieve maximum concentration (T_{max}) was 5.8 hours post-dose in the serum, followed by 6.7 hours, 6.5 hours and 13 days post-dose in the saliva, urine, and feces, respectively. The median time to achieve first below limit of quantification (BLOQ) sample followed by 2 consecutive BLOQ samples was 55 days post-dose for serum. The median time to achieve complete elimination as the first below limit of detection (BLOD) sample followed by 2 consecutive BLOD samples were 49.8 days, 78 days and 162 days post-dose for saliva, urine and feces, respectively. The estimated elimination half-life of ELEVIDYS vector genome in the serum is approximately 12 hours, and the majority of the drug is expected to be cleared from the serum by 1-week post-dose. In the excreta, the estimated elimination half-life of ELEVIDYS vector genome is approximately 40 hours, 55 hours, and 60 hours in the urine, feces, and saliva, respectively. As an AAV-based gene therapy that consists of a protein capsid containing the transgene DNA genome of interest, ELEVIDYS capsid proteins are broken down through proteasomal degradation following AAV entry into target cells. As such, ELEVIDYS is not likely to exhibit the drug-drug interaction potential mediated by known drug metabolizing enzymes (cytochrome P450-based) and drug transporters.

12.6 Immunogenicity

The observed incidence of anti-AAVrh74 antibodies is highly dependent on the sensitivity and specificity of the assay. Differences in assay methods preclude meaningful comparisons of the incidence of anti-AAVrh74 antibodies in the studies described below with the incidence of anti-AAVrh74 antibodies in other studies.

In ELEVIDYS clinical studies, patients were required to have baseline anti-AAVrh74 total binding antibodies of <1:400, measured using an investigational total binding antibody enzyme-linked immunosorbent assay (ELISA), and only patients with baseline anti-AAVrh74 total binding antibodies <1:400 were enrolled in those studies.

Across clinical studies evaluating a total of 156 patients, elevated anti-AAVrh74 total binding antibodies titers were observed in all patients following a one-time ELEVIDYS infusion. Anti-AAVrh74 total binding antibody titers reached at least 1:3200 in every patient, and the maximum titers exceeded 1:26,214,400 in certain patients. The safety of re-administration of ELEVIDYS or any other AAVrh74 vector-based gene therapy in the presence of high anti-AAVrh74 total binding antibody titer has not been evaluated in humans *[see Warnings and Precautions (5.6)]*.

There is insufficient data to assess whether the observed anti-AAVrh74 antibodies titers have clinically significant effect on pharmacokinetics, pharmacodynamics, safety, and efficacy of ELEVIDYS.

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

No animal studies have been performed to evaluate the effects of ELEVIDYS on carcinogenicity, mutagenesis, or impairment of fertility.

14 CLINICAL STUDIES

The efficacy of ELEVIDYS was evaluated in two double-blind, placebo-controlled studies (Study 1 [NCT 03769116] and Study 3 [NCT 05096221]) and one open-label study (Study 2 [NCT 04626674]) in which a total of 214 male patients with a confirmed disease-causing mutation in the *DMD* gene were dosed.

Study 1

Study 1 is a completed multi-center study including:

- Part 1: a 48-week, randomized, double-blind, placebo-controlled period
- Part 2: a 48-week period that began following completion of Part 1. Patients who received placebo during Part 1 were treated with ELEVIDYS, and patients treated with ELEVIDYS during Part 1 received placebo.

The study population consisted of male ambulatory DMD patients (N=41) aged 4 through 7 years with either a confirmed frameshift mutation, or a premature stop codon mutation between exons 18 to 58 in the *DMD* gene.

Patients were randomized 1:1 to receive either ELEVIDYS (N=20) or placebo (N=21), as a single intravenous infusion via a peripheral limb. Randomization was stratified by age (i.e., aged 4 to 5 years vs. aged 6 to 7 years). In the 4 through 5-year-old subgroup, the mean age, mean weight and mean NSAA total score (range) for the ELEVIDYS-treated patients (n=8) were 4.98 years, 20.1 kg and 20.1 (17-23), and for the placebo patients (n=8) were 5.15 years, 19.8 kg and 20.4 (15-24). In the ELEVIDYS group, eight patients received 1.33×10^{14} vg/kg of ELEVIDYS, and 12 patients received lower doses. Key demographic and baseline characteristics are presented in Table 6.

Table 6: Key Demographic and Baseline Characteristics (Study 1 Part 1)

Characteristic	All (n=41)	ELEVIDYS (n=20)	Placebo (n=21)
Race (%) Asian/Black or African American/White/Other	12/0/73/15	20/0/65/15	5/0/81/14
Ethnicity (%) Hispanic or Latino/ Other	12/88	5/95	19/81
Mean age [range] (years)	6.3 [4.3 to 7.9]	6.3 [4.5 to 7.9]	6.2 [4.3 to 7.9]
Mean weight [range] (kg)	22.4 [15.0 to 34.5]	23.3 [18.0 to 34.5]	21.6 [15.0 to 30.0]
Mean NSAA total score [range]	21.2 [13 to 29]	19.8 [13 to 26]	22.6 [15 to 29]
Mean time to rise from floor [range] (seconds)	4.3 [2.7 to 10.4]	5.1 [3.2 to 10.4]	3.6 [2.7 to 4.8]

All patients were on a stable dose of corticosteroids for DMD for at least 12 weeks prior to ELEVIDYS infusion. All randomized patients had baseline anti-AAVrh74 antibody titers <1:400 as determined by an investigational total binding antibody ELISA.

One day prior to treatment with ELEVIDYS or placebo, the patient's background dose of corticosteroid for DMD was increased to at least 1 mg/kg of a corticosteroid (prednisone equivalent) daily and was continued at this level for at least 60 days after the infusion, unless earlier tapering was clinically indicated.

The efficacy outcomes of Study 1 were to evaluate expression of micro-dystrophin in skeletal muscle, and to evaluate the effect of ELEVIDYS on the North Star Ambulatory Assessment (NSAA) total score.

Results of micro-dystrophin measured by western blot are presented in Table 5 *[see Clinical Pharmacology (12.2)]*.

The change in NSAA total score was assessed from baseline to Week 48 after infusion of ELEVIDYS or placebo. The difference between the ELEVIDYS and placebo groups was not statistically significant (p=0.37). The least squares (LS) mean changes in NSAA total score from baseline to Week 48 was 1.7 (standard error [SE]: 0.6) points for the ELEVIDYS group and 0.9 (SE: 0.6) points for the placebo group.

Exploratory subgroup analyses showed that for patients aged 4 through 5 years, the LS mean changes (SE) in NSAA total score from baseline to Week 48 were 4.3 (0.7) points for the ELEVIDYS group, and 1.9 (0.7) points for the placebo group, a numerical advantage for ELEVIDYS. For patients aged 6 through 7 years, the LS mean changes (SE) in NSAA total score from baseline to Week 48 were -0.2 (0.7) points for the ELEVIDYS group and 0.5 (0.7) points for the placebo group, a numerical disadvantage for ELEVIDYS.

Study 2

Study 2 is an ongoing, open-label, multi-center study which includes 5 cohorts of 48 male DMD patients.

Patients in cohorts 1, 2 and 3 have a confirmed frameshift, splice site or premature stop codon mutation anywhere in the *DMD* gene, while patients in cohort 4 included patients with mutations in the *DMD* gene starting at or after exon 18. All patients in cohort 5 had mutations that partially or fully overlap with exons 1-17 in the *DMD* gene. Patients received corticosteroids for DMD before infusion according to Table 1 *[see Dosage and Administration (2.2)]*. All patients had baseline anti-AAVrh74 antibodies titers ≤1:400 as determined by the investigational total binding antibody ELISA. Patients received a single intravenous infusion of 1.33×10^{14} vg/kg ELEVIDYS if they weighed less than 70 kg or 9.31×10^{15} vg/kg total fixed dose if they weighed 70 kg or greater.

Cohorts 1, 2, 4 and 5a enrolled 40 ambulatory patients 3 to 12 years of age, with weights ranging from 12.5 to 50.5 kg, baseline mean NSAA total score of 20.3 (11 to 30), and mean time to rise from floor of 4.7 seconds (2.4 to 9.7). Cohorts 3 and 5b include 8 non-ambulatory patients 10 to 20 years of age, with weights ranging from 36.1 to 80.1 kg. Overall key demographics and key baseline characteristics by Cohort are presented in Table 7.

Table 7: Key Demographic and Baseline Characteristics for Study 2

Characteristics	All (n=48)	Cohort 1 (n=20)	Cohort 2 (n=7)	Cohort 3[a] (n=6)	Cohort 4 (n=7)	Cohort 5a (n=6)	Cohort 5b[a] (n=2)
Race (%) Asian/Black or African American/White/Other	8/6/77/8	5/5/75/15	14/0/71/14	0/0/100/0	14/0/86/0	0/33/67/0	50/0/50/0
Ethnicity (%) Hispanic or Latino/ Not Hispanic or Latino	15/85	25/75	14/86	0/100	14/86	0/100	0/100
Mean age [range] (years)	7.7 [3.2 to 20.2]	5.8 [4.4 to 7.9]	10.1 [8.0 to 12.1]	15.3 [9.9 to 20.2]	3.5 [3.2 to 3.9]	6.7 [4.7 to 8.6]	13.4 [12.3 to 14.6]
Mean weight [range] (kg)	30.1 [12.5 to 80.1]	21.2 [15.2 to 33.1]	37.1 [28.0 to 50.5]	59.9 [36.1 to 80.1]	15.2 [12.5 to 16.5]	32.1 [19.1 to 47.4]	51.2 [43.4 to 59.0]
Mean NSAA total score [range]	20.3 [11 to 30]	22.1 [18 to 26]	20.7 [17 to 26]	N/A	12.9 [11 to 17]	22.5 [18 to 30]	N/A
Mean time to rise from floor [range] (seconds)	4.7 [2.4 to 9.7]	4.2 [2.4 to 8.2]	5.9 [3.8 to 9.7]	N/A	5.2 [3.8 to 6.7]	4.6 [2.5 to 7.7]	N/A
Mean Performance of Upper Limb v. 2.0 score [range]	30.7 [18 to 42]	NA	38.9 [33 to 42]	22.2 [18 to 31]	NA	NA	27.5 [21 to 34]

[a] NSAA and Time to rise from floor were not evaluated in non-ambulatory patients

The efficacy outcome measure of the study was to evaluate the effect of micro-dystrophin expression as measured by western blot. Results are presented in Table 5 *[see Clinical Pharmacology (12.2)]*.

Study 3

Study 3 is a multi-center, randomized, double-blind, placebo-controlled study in which 125 ambulatory male patients aged 4 through 7 years, with a confirmed frameshift, splice site, premature stop codon, or other disease-causing mutation in the *DMD* gene starting at or after exon 18, were dosed. Patients with exon 45 (inclusive), or in-frame deletions, in-frame duplications, and variants of uncertain significance ("VUS"), were excluded. Patients received corticosteroids for DMD before infusion according to Table 1 *[see Dosage and Administration (2.2)]*. All patients had baseline anti-AAVrh74 antibodies titers <1:400 as determined by the investigational total binding antibody ELISA and received a single intravenous infusion of 1.33×10^{14} vg/kg ELEVIDYS. Key demographic and baseline characteristics are presented in Table 8.

The efficacy outcome measure of the study was to evaluate the effect of ELEVIDYS on physical function as assessed by the NSAA total score. Key secondary outcome measures were to evaluate expression of micro-dystrophin in skeletal muscle, time to rise from floor, and time of 10-meter walk/run. Additional efficacy outcome measures included time of 100-meter walk/run, and time to ascend 4 steps. Results of micro-dystrophin measured by western blot are presented in Table 5 *[see Clinical Pharmacology (12.2)]*.

Table 8: Key Demographic and Baseline Characteristics for Study 3

Characteristic	ELEVIDYS (n=63)	Placebo (n=62)
Race (%) Asian/Black or African American/ White/Multiple/Other/Not Reported	13/0/78/2/3/5	18/3/74/0/2/3
Ethnicity (%) Hispanic or Latino/Not Hispanic or Latino/ Not Reported/Unknown	24/75/0/2	13/86/2/0
Mean age [range] (years)	6.0 [4.1 to 7.9]	6.1 [4.0 to 7.9]
Mean weight [range] (kg)	21.3 [13.5 to 38.5]	22.4 [14.4 to 41.6]
Mean NSAA total score [range]	23.10 [14 to 32]	22.82 [15.5 to 30]
Mean time to rise from floor [range] (seconds)	3.52 [1.9 to 5.8]	3.60 [2.3 to 5]
Mean time of 10-meter walk/run [range] (seconds)	4.82 [3.2 to 6.9]	4.92 [3.7 to 7]
Mean time of 100-meter walk/run [range] (seconds)	60.67 [38.0 to 129.2]	63.01 [38.7 to 118.1]
Mean time to ascend 4 steps [range] (seconds)	3.17 [1.6 to 7.1]	3.37 [1.5 to 7.1]

The change in NSAA total score was assessed from baseline to Week 52 after infusion of ELEVIDYS or placebo. The difference between the ELEVIDYS (n=63) and placebo groups (n=61) was not statistically significant (p=0.24). The least squares (LS) mean changes in NSAA total score from baseline to Week 52 was 2.57 (95% confidence interval [CI]: 1.80, 3.34) points for the ELEVIDYS group and 1.92 (95% CI: 1.14, 2.70) points for the placebo group, with a LS mean difference from placebo of 0.65 (95% CI: -0.45, 1.74). Changes of clinical relevance were noted in three secondary efficacy endpoints, including time to rise from the floor, 10-meter walk/run and time to ascend 4 steps.

Table 9: Change from Baseline to Week 52 of Timed Function Tests in Study 3 Part 1

	ELEVIDYS	Placebo	LS Mean Difference from placebo (95% CI)
Time to rise from the floor (seconds)	n=63	n=61	-
LS mean Change (95% CI)	-0.27 (-0.56, 0.02)	0.37 (0.08, 0.67)	-0.64 (-1.06, -0.23)
Time of 10-meter walk/run (seconds)	n=63	n=61	-
LS mean Change (95% CI)	-0.34 (-0.55, -0.14)	0.08 (-0.13, 0.29)	-0.42 (-0.71, -0.13)
Time of 100-meter walk/run (seconds)	n=59	n=57	-
LS mean Change (95% CI)	-6.57 (-10.05, -3.09)	-3.28 (-6.86, 0.29)	-3.29 (-8.28, 1.70)
Time to ascend 4 steps (seconds)	n=62	n=60	-
LS mean Change (95% CI)	-0.44 (-0.69, -0.20)	-0.08 (-0.33, 0.17)	-0.36 (-0.71, -0.01)

16 HOW SUPPLIED/STORAGE AND HANDLING

16.1 How Supplied

ELEVIDYS is shipped frozen (≤ -60ºC [-76ºF]) in 10 mL vials.

ELEVIDYS is supplied as a customized kit to meet dosing requirements for each patient *[see Dosage and Administration (2.2)]*. Each kit contains:

- Ten (10) to seventy (70) single-dose vials of ELEVIDYS
- One alcohol wipe per vial

Each ELEVIDYS pack may contain a maximum of two different drug product lots.

The total number of vials in each kit corresponds to the dosing requirement for the individual patient, based on the patient's body weight, and is specified on the package *[see Dosage and Administration (2.2)]*. Each kit includes a specified number of ELEVIDYS vials (with a minimum of 10 vials for a patient with 10.0 – 10.4 kg body weight range, and a maximum of 70 vials for a patient with body weight of 69.5 kg and above).
Kit sizes and National Drug Codes (NDC) are provided in Table 10.

Table 10: ELEVIDYS Multi-vial Kits

Patient Weight (kg)	Total Vials per Kit	Total Dose Volume per Kit (mL)	NDC Number
10.0 – 10.4	10	100	60923-501-10
10.5 – 11.4	11	110	60923-502-11
11.5 – 12.4	12	120	60923-503-12
12.5 – 13.4	13	130	60923-504-13
13.5 – 14.4	14	140	60923-505-14
14.5 – 15.4	15	150	60923-506-15
15.5 – 16.4	16	160	60923-507-16
16.5 – 17.4	17	170	60923-508-17
17.5 – 18.4	18	180	60923-509-18
18.5 – 19.4	19	190	60923-510-19
19.5 – 20.4	20	200	60923-511-20
20.5 – 21.4	21	210	60923-512-21
21.5 – 22.4	22	220	60923-513-22
22.5 – 23.4	23	230	60923-514-23
23.5 – 24.4	24	240	60923-515-24
24.5 – 25.4	25	250	60923-516-25
25.5 – 26.4	26	260	60923-517-26
26.5 – 27.4	27	270	60923-518-27

Patient Weight (kg)	Total Vials per Kit	Total Dose Volume per Kit (mL)	NDC Number
27.5 – 28.4	28	280	60923-519-28
28.5 – 29.4	29	290	60923-520-29
29.5 – 30.4	30	300	60923-521-30
30.5 – 31.4	31	310	60923-522-31
31.5 – 32.4	32	320	60923-523-32
32.5 – 33.4	33	330	60923-524-33
33.5 – 34.4	34	340	60923-525-34
34.5 – 35.4	35	350	60923-526-35
35.5 – 36.4	36	360	60923-527-36
36.5 – 37.4	37	370	60923-528-37
37.5 – 38.4	38	380	60923-529-38
38.5 – 39.4	39	390	60923-530-39
39.5 – 40.4	40	400	60923-531-40
40.5 – 41.4	41	410	60923-532-41
41.5 – 42.4	42	420	60923-533-42
42.5 – 43.4	43	430	60923-534-43
43.5 – 44.4	44	440	60923-535-44
44.5 – 45.4	45	450	60923-536-45
45.5 – 46.4	46	460	60923-537-46
46.5 – 47.4	47	470	60923-538-47
47.5 – 48.4	48	480	60923-539-48
48.5 – 49.4	49	490	60923-540-49
49.5 – 50.4	50	500	60923-541-50
50.5 – 51.4	51	510	60923-542-51
51.5 – 52.4	52	520	60923-543-52
52.5 – 53.4	53	530	60923-544-53
53.5 – 54.4	54	540	60923-545-54
54.5 – 55.4	55	550	60923-546-55
55.5 – 56.4	56	560	60923-547-56
56.5 – 57.4	57	570	60923-548-57

Patient Weight (kg)	Total Vials per Kit	Total Dose Volume per Kit (mL)	NDC Number
57.5 – 58.4	58	580	60923-549-58
58.5 – 59.4	59	590	60923-550-59
59.5 – 60.4	60	600	60923-551-60
60.5 – 61.4	61	610	60923-552-61
61.5 – 62.4	62	620	60923-553-62
62.5 – 63.4	63	630	60923-554-63
63.5 – 64.4	64	640	60923-555-64
64.5 – 65.4	65	650	60923-556-65
65.5 – 66.4	66	660	60923-557-66
66.5 – 67.4	67	670	60923-558-67
67.5 – 68.4	68	680	60923-559-68
68.5 – 69.4	69	690	60923-560-69
69.5 and above	70	700	60923-561-70

A 10 mL single-dose vial carton for ELEVIDYS (NDC 60923-562-01) is not sold individually.

16.2 Storage and Handling

- ELEVIDYS is shipped and delivered at ≤ −60ºC [−76ºF].
- ELEVIDYS can be refrigerated for up to 14 days when stored at 2°C to 8°C (36º F to 46º F) in the upright position.
- Do not refreeze.
- Do not shake.
- Do not place back in the refrigerator once brought to room temperature.
- Follow local guidelines on handling of biological waste.

17 PATIENT COUNSELING INFORMATION

Advise the patient to read the FDA-approved Medication Guide.

Provide a copy of the Medication Guide and review the contents with the patient.

Inform patients or caregivers that:

- ELEVIDYS can increase certain liver enzyme levels and cause acute serious liver injury or acute liver failure, and death. Management of acute serious liver injury may require hospitalization. Patients will receive oral corticosteroid medication before and after infusion with ELEVIDYS. Weekly blood tests will be required to monitor liver enzyme levels for 3 months after treatment. Contact a healthcare provider immediately if the patient's skin and/or whites of the eyes appear yellowish, if the patient misses a dose of corticosteroid or vomits it up, or if the patient experiences a change in mental status *[see Warnings and Precautions (5.1)]*.
- Due to the concomitant administration of corticosteroids, an infection (e.g., cold, flu, gastroenteritis, otitis media, bronchiolitis, pneumonia, etc.) before or after ELEVIDYS infusion could lead to more

serious complications, including death. Contact a healthcare provider immediately if symptoms suggestive of infection are observed (e.g., coughing, wheezing, sneezing, runny nose, sore throat, or fever) *[see Warnings and Precautions (5.2)]*.

- Myocarditis (inflammation of the heart) has been observed within days to more than a year following ELEVIDYS infusion. Weekly monitoring of troponin-I for the first month after treatment is required. Contact a healthcare provider immediately if the patient begins to experience chest pain and/or shortness of breath *[see Warnings and Precautions (5.3)]*.
- Infusion-related reactions including hypersensitivity and anaphylaxis have occurred during and after ELEVIDYS infusion. Possible symptoms of infusion-related reactions are fast heart rate, fast breathing, swollen lips, being short of breath, nostrils widening, hives, red and blotchy skin, itchy or inflamed lips, rash, vomiting, nausea, chills and fever. Contact a healthcare provider immediately if the patient experiences such a reaction *[see Warnings and Precautions (5.4)].*
- Immune-mediated myositis (an immune response affecting muscles) was observed in some patients following ELEVIDYS infusion. Contact a physician immediately if the patient experiences any unexplained increased muscle pain, tenderness, or weakness, including difficulty swallowing, difficulty breathing or difficulty speaking, as these may be symptoms of myositis *[see Warnings and Precautions (5.5)]*.
- Patient's immunizations should be up to date with current immunization guidelines prior to initiation of the corticosteroid regimen required before ELEVIDYS infusion. Vaccinations should be completed at least 4 weeks prior to initiation of the corticosteroid regimen *[see Drug Interactions (7)]*.
- Vector shedding of ELEVIDYS occurs primarily through body waste. Practice proper hand hygiene, such as hand washing, when coming into direct contact with patient body waste. Place potentially contaminated materials that may have the patient's bodily fluids/waste in a sealable bag and dispose into regular trash. These precautions should be followed for one month after ELEVIDYS infusion.

Manufactured for: Sarepta Therapeutics, Inc.
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Elevidys
delandistrogene
moxeparvovec-rokl
suspension for intravenous infusion