

HIGHLIGHTS OF PRESCRIBING INFORMATION

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

SPINRAZA (nusinersen) injection, for intrathecal use
Initial U.S. Approval: 2016

RECENT MAJOR CHANGES

Dosage and Administration (2.1, 2.2, 2.3,2.5)	03/2026
Warnings and Precautions (5.1)	03/2026

INDICATIONS AND USAGE

SPINRAZA is a survival motor neuron-2 (SMN2)-directed antisense oligonucleotide indicated for the treatment of spinal muscular atrophy (SMA) in pediatric and adult patients (1)

DOSAGE AND ADMINISTRATION

SPINRAZA is administered intrathecally (2.1)

Recommended Dosage (2.1)

- The recommended dosage is one of two options:
 - Low Dose Regimen: Administer one 12 mg loading dose every 14 days for three doses; then a fourth 12 mg loading dose 30 days after the third dose; then administer a 12 mg maintenance dose once every 4 months thereafter.
 - High Dose Regimen: Administer one 50 mg loading dose followed by a second 50 mg loading dose 14 days later; then administer a 28 mg maintenance dose once every 4 months thereafter.

Important Preparation and Administration Instructions (2.3)

- Allow to warm to room temperature prior to administration
- Administer within 4 hours of removal from vial
- Prior to administration, remove 5 mL of cerebrospinal fluid
- Administer as intrathecal bolus injection over 1 to 3 minutes

Laboratory Testing and Monitoring to Assess Safety (2.4)

- At baseline and prior to each dose, obtain a platelet count, coagulation laboratory testing, and quantitative spot urine protein testing

DOSAGE FORMS AND STRENGTHS

- Injection: 12 mg/5 mL (2.4 mg/mL) in a single-dose vial (3)
- Injection: 28 mg/5 mL (5.6 mg/mL) in a single-dose vial (3)
- Injection: 50 mg/5 mL (10 mg/mL) in a single-dose vial (3)

CONTRAINDICATIONS

None.

WARNINGS AND PRECAUTIONS

- Thrombocytopenia and Coagulation Abnormalities:** Increased risk for bleeding complications; testing required at baseline and before each dose and as clinically needed (5.1, 2.3)
- Renal Toxicity:** Quantitative spot urine protein testing required at baseline and prior to each dose (5.2, 2.3)

ADVERSE REACTIONS

The most common adverse reactions that occurred in at least 20% of SPINRAZA-treated patients who received Low Dose Regimen and occurred at least 5% more frequently than in control patients were:

- lower respiratory infection and constipation in patients with infantile-onset SMA (6.1)
- pyrexia, headache, vomiting, and back pain in patients with later-onset SMA (6.1)

The most common adverse reactions in at least 10% of SPINRAZA-treated patients who received High Dose Regimen and occurred at least 5% more frequently than in historic matched sham-control were:

- pneumonia, COVID-19, pneumonia aspiration, and malnutrition in patients with infantile-onset SMA (6.1)

To report SUSPECTED ADVERSE REACTIONS, contact Biogen at 1-844-477-4672 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

USE IN SPECIFIC POPULATIONS

Pregnancy: Based on animal data, may cause fetal harm (8.1)

See 17 for PATIENT COUNSELING INFORMATION.

Revised: 03/2026

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FULL PRESCRIBING INFORMATION

1 INDICATIONS AND USAGE

SPINRAZA is indicated for the treatment of spinal muscular atrophy (SMA) in pediatric and adult patients.

2 DOSAGE AND ADMINISTRATION

2.1 Recommended Dosage

SPINRAZA is administered intrathecally by, or under the direction of, healthcare professionals experienced in performing lumbar punctures.

Two dosing regimen options for SPINRAZA, which consist of loading followed by maintenance dosages, are presented in Table 1.

Table 1: Recommended Dosage for SPINRAZA

	Loading Dosages	Maintenance Dosage
Low Dose Regimen (Low dose with four loading doses)	Administer a total of four loading doses as follows: <ul style="list-style-type: none">• one 12 mg dose every 14 days for three doses, then• a fourth 12 mg dose 30 days after the third dose.	Administer 12 mg once every 4 months starting 4 months after the last loading dose.
High Dose Regimen (High dose with two loading doses)	Administer a total of two loading doses as follows: <ul style="list-style-type: none">• one 50 mg dose followed by a second 50 mg dose 14 days later.	Administer 28 mg once every 4 months starting 4 months after the last loading dose.

2.2 Missed Doses

Missed Dose of Low Dose Regimen

Missed Loading Dose

If a 12 mg loading dose (any of the 4 loading doses) is missed, administer the missed loading dose as soon as possible; adjust the date for the subsequent doses to maintain the recommended interval between doses.

Missed Maintenance Dose

Less than 8 months from last maintenance dose

Administer the missed 12 mg maintenance dose as soon as possible; then administer the next maintenance dose per the originally scheduled date, as long as these two doses are administered at least 14 days apart.

At least 8 months but less than 16 months from last maintenance dose

Administer the missed 12 mg maintenance dose as soon as possible, followed by one additional dose 14 days later, and then administer the next maintenance dose 4 months thereafter.

At least 16 months but less than 40 months from last maintenance dose

Administer the missed 12 mg maintenance dose as soon as possible, followed by two additional doses 14 days apart, and then administer the next maintenance dose 4 months thereafter.

At least 40 months from last dose

Restart Low Dose Regimen with 12 mg loading doses as described in Recommended Dosage.

Missed Dose of High Dose Regimen

Missed Second 50 mg Loading Dose

Administer the missed 50 mg loading dose as soon as possible; then administer 28 mg maintenance doses every 4 months thereafter.

Missed 28 mg Maintenance Dose

Less than 8 months from last maintenance dose

Administer the missed 28 mg maintenance dose as soon as possible; administer the next 28 mg maintenance dose per the originally scheduled date, as long as these two doses are administered at least 14 days apart; then administer 28 mg every 4 months thereafter.

At least 8 months to less than 40 months from last maintenance dose

Administer a 50 mg loading dose as soon as possible; then administer 28 mg maintenance doses every 4 months thereafter.

At least 40 months from last maintenance dose

Restart High Dose Regimen with two 50 mg loading doses as described in Recommended Dosage.

2.3 Important Preparation and Administration Instructions

SPINRAZA is for intrathecal use only.

Prepare and use SPINRAZA according to the following steps using aseptic technique. Each vial is intended for single dose only.

Use the vial strength that corresponds to the prescribed dose. Do not combine SPINRAZA vials of different strengths and concentrations or use partial vials to achieve the prescribed dose.

Preparation

- Store SPINRAZA in the carton in a refrigerator until time of use.
- Allow the SPINRAZA vial to warm to room temperature (25° C/77° F) prior to administration. Do not use external heat sources.
- Inspect the SPINRAZA vial for particulate matter and discoloration prior to administration. Do not administer SPINRAZA if visible particulates are observed or if the liquid in the vial is discolored. The use of external filters is not required.
- Withdraw 5 mL of SPINRAZA from the single-dose vial into a syringe and discard unused contents of the vial.
- Administer SPINRAZA within 4 hours of removal from vial.

Administration

- Consider sedation as indicated by the clinical condition of the patient.
- Consider ultrasound or other imaging techniques to guide intrathecal administration of SPINRAZA, particularly in younger patients.
- Prior to administration, remove 5 mL of cerebrospinal fluid.
- Administer SPINRAZA as an intrathecal bolus injection over 1 to 3 minutes using a spinal anesthesia needle [see *Dosage and Administration (2.1)*]. Do not administer SPINRAZA in areas of the skin where there are signs of infection or inflammation [see *Adverse Reactions (6.3)*].

2.4 Laboratory Testing and Monitoring to Assess Safety

Conduct the following laboratory tests at baseline and prior to each dose of SPINRAZA and as clinically needed [see *Warnings and Precautions (5.1, 5.2)*]:

- Platelet count
- Prothrombin time; activated partial thromboplastin time
- Quantitative spot urine protein testing

2.5 Transition Between SPINRAZA Dose Regimens

If transitioning from SPINRAZA Low Dose Regimen to High Dose Regimen, administer a single 50 mg bolus dose at least four months (+/- 14 days) after the last 12 mg maintenance dose, followed by a 28 mg maintenance dose once every 4 months thereafter. Additional clinical benefit in patients who transition from the Low Dose Regimen to the High Dose Regimen has not been established in a controlled study.

3 DOSAGE FORMS AND STRENGTHS

Injection: nusinersen is a clear and colorless solution supplied in single-dose vials in the following strengths:

- 12 mg/5 mL (2.4 mg/mL)
- 28 mg/5 mL (5.6 mg/mL)
- 50 mg/5 mL (10 mg/mL)

4 CONTRAINDICATIONS

None.

5 WARNINGS AND PRECAUTIONS

5.1 Thrombocytopenia and Coagulation Abnormalities

Coagulation abnormalities and thrombocytopenia, including acute severe thrombocytopenia, have been observed after administration of some antisense oligonucleotides.

In the sham-controlled studies for patients with infantile-onset (Study 1) and later-onset (Study 2) SMA who received Low Dose Regimen [see *Clinical Studies (14.1, 14.2)*], 24 of 146 (16%) SPINRAZA-treated patients with high, normal, or unknown platelet count at baseline developed a platelet level below the lower limit of normal, compared to 10 of 72 (14%) sham-controlled patients.

In Study 2, two SPINRAZA-treated patients developed platelet counts less than 50,000 cells per microliter, with a lowest level of 10,000 cells per microliter recorded on study day 28.

In patients who received High Dose Regimen, decreases in platelet counts were also observed.

Because of the risk of thrombocytopenia and coagulation abnormalities from SPINRAZA, patients may be at increased risk of bleeding complications.

Perform a platelet count and coagulation laboratory testing at baseline and prior to each administration of SPINRAZA and as clinically needed.

5.2 Renal Toxicity

Renal toxicity, including potentially fatal glomerulonephritis, has been observed after administration of some antisense oligonucleotides.

SPINRAZA is present in and excreted by the kidney [see *Clinical Pharmacology (12.3)*]. In Study 1 and Study 2, 71 of 123 (58%) of SPINRAZA-treated patients had elevated urine protein, compared to 22 of 65 (34%) sham-controlled patients. Conduct quantitative spot urine protein testing (preferably using a first morning urine specimen) at baseline and prior to each dose of SPINRAZA. For urinary protein concentration greater than 0.2 g/L, consider repeat testing and further evaluation.

6 ADVERSE REACTIONS

The following serious adverse reactions are described in detail in other sections of the labeling:

- Thrombocytopenia and Coagulation Abnormalities [see *Warnings and Precautions (5.1)*]
- Renal Toxicity [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 SPINRAZA cannot be directly compared to rates in clinical trials of other drugs and may not reflect the rates observed in practice.

SPINRAZA Low Dose Regimen (12 mg loading doses/12 mg maintenance doses)

In clinical studies, 385 patients (47% male, 68% Caucasian, and 12% Asian) were treated with SPINRAZA Low Dose Regimen [see *Dosage and Administration* (2.1)], including 353 exposed for at least 6 months, 314 exposed for at least 1 year, and 256 exposed for at least 5 years.

Clinical Trial in Infantile-Onset SMA (Study 1)

In Study 1, baseline disease characteristics were largely similar in the SPINRAZA-treated patients and sham-control patients except that SPINRAZA-treated patients at baseline had a higher percentage compared to sham-control patients of paradoxical breathing (89% vs 66%), pneumonia or respiratory symptoms (35% vs 22%), swallowing or feeding difficulties (51% vs 29%), and requirement for respiratory support (26% vs 15%).

The most common adverse reactions that occurred in at least 20% of SPINRAZA-treated patients and occurred at least 5% more frequently than in control patients were lower respiratory infection and constipation. Serious adverse reactions of atelectasis were more frequent in SPINRAZA-treated patients (18%) than in control patients (10%). Because patients in Study 1 were infants, adverse reactions that are verbally reported could not be assessed in this study.

Table 2. Adverse Reactions that Occurred in at Least 5% of SPINRAZA Patients and Occurred at Least 5% More Frequently or At Least 2 Times as Frequently Than in Control Patients with Infantile-Onset SMA (Study 1)

Adverse Reactions	SPINRAZA 12 mg ¹	Sham-Procedure Control
	N = 80 %	N = 41 %
Lower respiratory infection ²	55	37
Constipation	35	22
Teething	18	7
Urinary tract infection	9	0
Upper respiratory tract congestion	8	2
Ear infection	6	2
Flatulence	5	2
Decreased weight	5	2

¹ Low Dose Regimen [see *Dosage and Administration* (2.1)]

² Includes adenovirus infection, bronchiolitis, bronchitis, bronchitis viral, corona virus infection, Influenza, lower respiratory tract infection, lower respiratory tract infection viral, lung infection, parainfluenzae virus

infection, pneumonia, pneumonia bacterial, pneumonia influenzal, pneumonia moraxella, pneumonia parainfluenzae viral, pneumonia pneumococcal, pneumonia pseudomonal, pneumonia respiratory syncytial viral, pneumonia viral, and respiratory syncytial virus bronchiolitis.

In an open-label clinical study in infants with symptomatic SMA, severe hyponatremia was reported in a patient treated with SPINRAZA requiring salt supplementation for 14 months.

Cases of rash were reported in patients treated with SPINRAZA. One patient, 8 months after starting SPINRAZA treatment, developed painless red macular lesions on the forearm, leg, and foot over an 8-week period. The lesions ulcerated and scabbed over within 4 weeks, and resolved over several months. A second patient developed red macular skin lesions on the cheek and hand ten months after the start of SPINRAZA treatment, which resolved over 3 months. Both cases continued to receive SPINRAZA and had spontaneous resolution of the rash.

SPINRAZA may cause a reduction in growth as measured by height when administered to infants, as suggested by observations from the controlled study. It is unknown whether any effect of SPINRAZA on growth would be reversible with cessation of treatment.

Clinical Trial in Later-Onset SMA (Study 2)

In Study 2, baseline disease characteristics were largely similar in the SPINRAZA-treated patients and sham-control patients except for the proportion of SPINRAZA-treated patients who had ever achieved the ability to stand without support (13% vs 29%) or walk with support (24% vs 33%).

The most common adverse reactions that occurred in at least 20% of SPINRAZA-treated patients and occurred at least 5% more frequently than in control patients were pyrexia, headache, vomiting, and back pain.

Table 3. Adverse Reactions that Occurred in at Least 5% of SPINRAZA Patients and Occurred at Least 5% More Frequently or At Least 2 Times as Frequently Than in Control Patients with Later-Onset SMA (Study 2)

Adverse Reactions	SPINRAZA 12 mg ¹	Sham-Procedure Control
	N=84 %	N=42 %
Pyrexia	43	36
Headache	29	7
Vomiting	29	12
Back pain	25	0
Epistaxis	7	0

Fall	5	0
Respiratory tract congestion	5	2
Seasonal allergy	5	2

¹ Low Dose Regimen [see Dosage and Administration (2.1)]

Post-lumbar puncture syndrome has also been observed after administration of SPINRAZA.

SPINRAZA High Dose Regimen (50 mg loading doses/28 mg maintenance doses)

The safety of SPINRAZA High Dose Regimen was studied in 2 clinical trials in symptomatic patients with SMA (approximately 14 days to 65 years of age at first dose). In clinical studies, 128 patients (50% male, 63% Caucasian, and 22% Asian) were treated with SPINRAZA High Dose Regimen [see Dosage and Administration (2.1)], including 113 exposed for at least 6 months, 95 exposed for at least 1 year, and 67 exposed for at least 2 years.

Clinical Trial in Infantile-Onset SMA (Study 4)

The most common adverse reactions that occurred in at least 10% of SPINRAZA treated patients and occurred at least 5% more frequently than in historic-matched sham-control patients from Study 1 were pneumonia, COVID-19, pneumonia aspiration, and malnutrition in patients with infantile-onset SMA. COVID-19 was not discovered at the time of Study 1.

Table 4: Adverse Reactions that Occurred in at least 5% of SPINRAZA High Dose Regimen Patients and Occurred at Least 5% More Frequently or At Least 2 Times as Frequently Than Matched Sham Control in Patients with Infantile Onset SMA (Study 4)

	SPINRAZA High Dose Regimen (Study 4) (N=50) %	Matched Sham Control (Study 1) (N=20) %
Pneumonia	20	5
COVID-19	16	0
Pneumonia aspiration	14	5
Malnutrition	10	0
Procedural pain	6	0
Myocardial necrosis marker increased	6	0
Anemia	6	0

6.2 Postmarketing Experience

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

Serious infections associated with lumbar puncture, such as meningitis, have been reported. Hydrocephalus, aseptic meningitis, hypersensitivity reactions (e.g. angioedema, urticaria, rash), and arachnoiditis have also been reported.

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary

There are no adequate data on the developmental risk associated with the use of SPINRAZA in pregnant women. When nusinersen was administered by subcutaneous injection to mice throughout pregnancy and lactation, developmental toxicity (long-term neurobehavioral impairment) was observed at all doses tested (*see Data*). In the U.S. general population, the estimated background risk of major birth defects and miscarriage in clinically recognized pregnancies is 2 to 4% and 15 to 20%, respectively. The background risk of major birth defects and miscarriage for the indicated population is unknown.

Data

Animal Data

When nusinersen (0, 3, 10, or 25 mg/kg) was administered subcutaneously to male and female mice every other day prior to and during mating and continuing in females throughout organogenesis, no adverse effects on embryofetal development were observed. Subcutaneous administration of nusinersen (0, 6, 12.6, or 25 mg/kg) to pregnant rabbits every other day throughout organogenesis produced no evidence of embryofetal developmental toxicity.

When nusinersen (1.4, 5.8, or 17.2 mg/kg) was administered to pregnant female mice by subcutaneous injection every other day throughout organogenesis and continuing once every six days throughout the lactation period, adverse neurobehavioral effects (alterations in locomotor activity, learning and memory deficits) were observed when offspring were tested after weaning or as adults. A no-effect level for neurobehavioral impairment was not established.

8.2 Lactation

Risk Summary

There are no data on the presence of nusinersen in human milk, the effects on the breastfed infant, or the effects of the drug on milk production. Nusinersen was detected in the milk of lactating mice when administered by subcutaneous injection. The developmental and health benefits of breastfeeding should be considered along with the mother's clinical need for SPINRAZA and any potential adverse effects on the breastfed infant from SPINRAZA or from the underlying maternal condition.

8.4 Pediatric Use

The safety and effectiveness of SPINRAZA in pediatric patients from newborn to 17 years have been established [*see Clinical Studies (14.1)*].

Juvenile Animal Toxicity Data

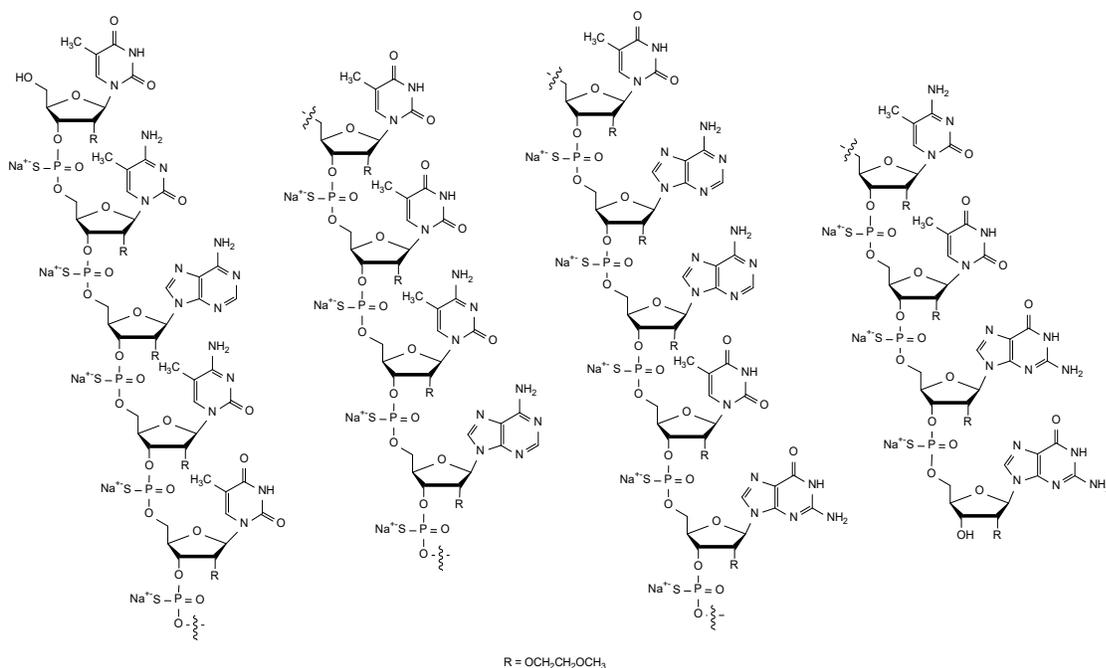
In intrathecal toxicity studies in juvenile monkeys, administration of nusinersen (0, 0.3, 1, or 3 mg/dose for 14 weeks and 0, 0.3, 1, or 4 mg/dose for 53 weeks) resulted in brain histopathology (neuronal vacuolation and necrosis/cellular debris in the hippocampus) at the mid and high doses and acute, transient deficits in lower spinal reflexes at the high dose in each study. In addition, possible neurobehavioral deficits were observed on a learning and memory test at the high dose in the 53-week monkey study. In a combined 6- and 13-week toxicity study in juvenile monkeys, intrathecal administration of nusinersen at higher doses (0, 5, 10, or 15 mg/dose) resulted in additional acute, transient effects, including limited use of limbs at the mid and high dose and uncoordinated movement at the high dose. The no-effect dose for neurohistopathology in monkeys (0.3 mg/dose) is approximately equivalent to and lower than the recommended clinical maintenance doses of 12 and 28 mg, respectively, when calculated on annual dose basis and corrected for species differences in CSF volume.

8.5 Geriatric Use

Clinical studies of SPINRAZA did not include sufficient numbers of subjects aged 65 and over to determine whether they respond differently from younger subjects.

11 DESCRIPTION

Nusinersen is a modified antisense oligonucleotide, where the 2'-hydroxy groups of the ribofuranosyl rings are replaced with 2'-O-2-methoxyethyl groups and the phosphate linkages are replaced with phosphorothioate linkages. Nusinersen binds to a specific sequence in the intron downstream of exon 7 of the *SMN2* transcript. The structural formula is:



SPINRAZA is supplied as a sterile, preservative-free, colorless solution for intrathecal use in a single-dose glass vial in the following strengths:

- 12 mg/5 mL (2.4 mg/mL)
- 28 mg/5 mL (5.6 mg/mL)
- 50 mg/5 mL (10 mg/mL)

Each 1 mL solution of the 12 mg/5 mL strength contains 2.4 mg of nusinersen (equivalent to 2.53 mg of nusinersen sodium salt). Each 1 mL also contains calcium chloride dihydrate (0.21 mg) USP, magnesium chloride hexahydrate (0.16 mg) USP, potassium chloride (0.22 mg) USP, sodium chloride (8.77 mg) USP, sodium phosphate dibasic anhydrous (0.10 mg) USP, sodium phosphate monobasic dihydrate (0.05 mg) USP, and Water for Injection USP.

Each 1 mL solution of the 28 mg/5 mL strength contains 5.6 mg of nusinersen (equivalent to 5.90 mg of nusinersen sodium salt). Each 1 mL also contains calcium chloride dihydrate (0.21 mg) USP, magnesium chloride hexahydrate (0.16 mg) USP, potassium chloride (0.22 mg) USP, sodium chloride (8.39 mg) USP, sodium phosphate dibasic anhydrous (0.10 mg) USP, sodium phosphate monobasic dihydrate (0.05 mg) USP, and Water for Injection USP.

Each 1 mL solution of the 50 mg/5 mL strength contains 10 mg of nusinersen (equivalent to 10.54 mg of nusinersen sodium salt). Each 1 mL also contains calcium chloride dihydrate (0.21 mg) USP, magnesium chloride hexahydrate (0.16 mg) USP, potassium chloride (0.22 mg) USP, sodium chloride (8.11 mg) USP, sodium phosphate dibasic anhydrous (0.10 mg) USP, sodium phosphate monobasic dihydrate (0.05 mg) USP, and Water for Injection USP.

For all strengths the product may contain hydrochloric acid or sodium hydroxide to adjust pH. The pH is ~7.2.

The molecular formula of SPINRAZA is $C_{234}H_{323}N_{61}O_{128}P_{17}S_{17}Na_{17}$ and the molecular weight is 7501.0 daltons.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

SPINRAZA is an antisense oligonucleotide (ASO) designed to treat SMA caused by mutations in chromosome 5q that lead to SMN protein deficiency. Using in vitro assays and studies in transgenic animal models of SMA, SPINRAZA was shown to increase exon 7 inclusion in *SMN2* messenger ribonucleic acid (mRNA) transcripts and production of full-length SMN protein.

12.2 Pharmacodynamics

Autopsy samples from patients (n=3) had higher levels of *SMN2* messenger ribonucleic acid (mRNA) containing exon 7 in the thoracic spinal cord compared to untreated SMA infants.

Cardiac Electrophysiology

Across the sham-controlled studies in 247 patients with spinal muscular atrophy who received either SPINRAZA Low Dose Regimen or sham-control, QTcF values > 500 ms and change from baseline values > 60 ms were observed in 4 (2.4%) patients receiving SPINRAZA. Compared to the sham-control, there was no increase in the incidence of cardiac adverse reactions associated with delayed ventricular repolarization in patients treated with SPINRAZA.

12.3 Pharmacokinetics

Absorption

Intrathecal injection of SPINRAZA into the cerebrospinal fluid (CSF) allows nusinersen to be distributed from the CSF to the target central nervous system (CNS) tissues. Following intrathecal administration, trough plasma concentrations of nusinersen were relatively low, compared to the trough CSF concentration. Median plasma T_{max} values ranged from 1.7 to 6.0 hours. Mean plasma C_{max} and AUC values increased approximately dose-proportionally up to a dose of 12 mg.

Distribution

Autopsy data from patients (n=3) showed that SPINRAZA administered intrathecally was distributed within the CNS and peripheral tissues, such as skeletal muscle, liver, and kidney.

Elimination

Metabolism

Nusinersen is metabolized via exonuclease (3'- and 5')-mediated hydrolysis and is not a substrate for, or inhibitor or inducer of CYP450 enzymes.

Excretion

The mean terminal elimination half-life is estimated to be 135 to 177 days in CSF, and 63 to 87 days in plasma. The primary route of elimination is likely by urinary excretion for nusinersen and its chain-shortened metabolites. At 24 hours, only 0.5% of the administered dose was recovered in the urine.

12.6 Immunogenicity

The observed incidence of anti-drug antibodies is highly dependent on the sensitivity and specificity of the assay. Differences in assay methods preclude meaningful comparisons of the incidence of anti-drug antibodies in the studies described below with the incidence of anti-drug antibodies in other studies, including those of SPINRAZA or of other nusinersen products.

The immunogenic response to nusinersen in patients who received SPINRAZA Low Dose Regimen was evaluated in 367 patients with post-baseline plasma samples for anti-drug antibodies (ADAs). Thirty eight patients (10%) developed treatment-emergent ADAs, of which 16 were transient and 22 were considered to be persistent. Persistent was defined as having one positive test followed by another one more than 100 days after the first positive test. In addition, “persistent” is also defined as having one or more positive samples and no sample more than 100 days after the first positive sample. Transient was defined as having one or more positive results and not confirmed to be persistent.

The immunogenic response to nusinersen in patients who received SPINRAZA High Dose Regimen was evaluated in 117 patients with post-baseline plasma samples for ADAs. Eleven patients (9%) developed treatment-emergent ADAs, of which 5 were transient and 6 were persistent.

There are insufficient data to evaluate an effect of ADAs on clinical response, adverse events, or the pharmacokinetic profile of nusinersen.

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

Carcinogenesis

Administration of nusinersen (0, 5, 15, or 50 mg/kg) to male and female mice by subcutaneous injection, once every two weeks for 2 years, resulted in an increase in the incidence of vascular tumors (combined hemangioma and hemangiosarcoma) at the highest dose tested.

Mutagenesis

Nusinersen demonstrated no evidence of genotoxicity in in vitro (Ames and chromosomal aberration in CHO cells) and in vivo (mouse micronucleus) assays.

Impairment of Fertility

When nusinersen (0, 3, 10, or 25 mg/kg) was administered by subcutaneous injection to mice every other day prior to and during mating and continuing in females throughout organogenesis, no adverse effects on male or female fertility were observed.

14 CLINICAL STUDIES

The efficacy of SPINRAZA Low Dose Regimen was demonstrated in two double-blind, sham-procedure controlled clinical trials in patients with symptomatic infantile-onset and later-onset SMA (Study 1 and Study 2) and was supported by open-label clinical trials conducted in patients

with presymptomatic (Study 3) and symptomatic SMA [see *Dosage and Administration (2.1)*]. The efficacy of SPINRAZA High Dose Regimen was demonstrated in a double-blind, randomized, external-controlled trial in patients with symptomatic infantile-onset and later-onset SMA (Study 4). The overall findings from these trials support the effectiveness of SPINRAZA administered with either the Low Dose Regimen or the High Dose Regimen across the range of ages and severities in patients with SMA, and support the early initiation of treatment with SPINRAZA.

14.1 Infantile-Onset SMA

Study 1 (NCT02193074) was a multicenter, randomized, double-blind, sham-procedure controlled study in 121 symptomatic infants ≤ 7 months of age at the time of first dose, diagnosed with SMA (symptom onset before 6 months of age). Patients were randomized 2:1 to receive either SPINRAZA Low Dose Regimen or sham injection as a series of loading doses administered intrathecally followed by maintenance doses administered every 4 months. Patients in this study were deemed most likely to develop Type 1 SMA.

A planned interim efficacy analysis was conducted based on patients who died, withdrew, or completed at least 183 days of treatment. Of the 82 patients included in the interim analysis (52 patients in the SPINRAZA-treated group and 30 in the sham-control group), 44% were male, 87% were Caucasian, 2% were Black, and 4% were Asian. Age at first treatment ranged from 30 to 262 days (median 181). Length of treatment ranged from 6 to 442 days (median 261 days). Baseline demographics were balanced between the SPINRAZA and control groups with the exception of age at first treatment (median age 175 vs. 206 days, respectively). The SPINRAZA and control groups were balanced with respect to gestational age, birth weight, disease duration, and SMN2 copy number. Median disease duration was 14 weeks. There was some imbalance in age at symptom onset with 88% of subjects in the SPINRAZA group and 77% in the control group experiencing symptoms within the first 12 weeks of life.

The primary endpoint assessed at the time of interim analysis was the proportion of responders: patients with an improvement in motor milestones according to Section 2 of the Hammersmith Infant Neurologic Exam (HINE). This endpoint evaluates seven different areas of motor milestone development, with a maximum score between 2-4 points for each, depending on the milestone, and a total maximum score of 26. A treatment responder was defined as any patient with at least a 2-point increase (or maximal score of 4) in ability to kick (consistent with improvement by at least 2 milestones), or at least a 1-point increase in the motor milestones of head control, rolling, sitting, crawling, standing or walking (consistent with improvement by at least 1 milestone). To be classified as a responder, patients needed to exhibit improvement in more categories of motor milestones than worsening. Of the 82 patients who were eligible for the interim analysis, a statistically significantly greater percentage of patients achieved the definition of a motor milestone responder in the SPINRAZA group (40%) compared to the sham-control group (0%). Results from the final analysis were consistent with those from the interim analysis (Table 5). Fifty-one percent of patients in the SPINRAZA group achieved the definition of a motor milestone responder compared to 0% of patients in the sham-control group. Figure 1 is a descriptive display of the distribution of net change from baseline in the total motor milestone score for Section 2 of the HINE for patients in the final efficacy set who did not die or withdraw from the study.

The primary endpoint assessed at the final analysis was time to death or permanent ventilation (\geq 16 hours ventilation/day continuously for $>$ 21 days in the absence of an acute reversible event or tracheostomy). Statistically significant effects on event-free survival and overall survival were observed in patients in the SPINRAZA group compared to those in the sham-control group (Table 6). A 47% reduction in the risk of death or permanent ventilation was observed in the SPINRAZA group ($p=0.005$) (Figure 2). Median time to death or permanent ventilation was not reached in SPINRAZA group and was 22.6 weeks in the sham-control group. A statistically significant 63% reduction in the risk of death was also observed ($p=0.004$).

At the final analysis, the study also assessed treatment effects on the Children’s Hospital of Philadelphia Infant Test of Neuromuscular Disorders (CHOP-INTEND), which is an evaluation of motor skills in patients with infantile-onset SMA. The CHOP-INTEND results are displayed in Table 5.

Table 5. Motor Milestone Response and CHOP-INTEND Results of the Final Analysis of Patients with Infantile-Onset SMA (Study 1)

Endpoint	SPINRAZA- treated Patients ¹ (n=73)	Sham-control Patients (n=37)
Motor function		
Motor milestones²		
Proportion achieving pre-defined motor milestone responder criteria (HINE section 2) ^{3,4}	37 (51%) P<0.0001	0 (0%)
CHOP-INTEND²		
Proportion achieving a 4-point improvement	52 (71%) p<0.0001	1 (3%)
Proportion achieving a 4-point worsening ⁵	2 (3%)	17 (46%)

¹ Low Dose Regimen [see Dosage and Administration (2.1)]

² At the final analysis, CHOP-INTEND and motor milestone analyses were conducted using the Efficacy Set (SPINRAZA n=73; Sham-control n=37).

³ Assessed at the later of Day 183, Day 302, and Day 394 Study Visit

⁴ According to HINE section 2: ≥ 2 point increase [or maximal score] in ability to kick, OR ≥ 1 point increase in the motor milestones of head control, rolling, sitting, crawling, standing or walking, AND improvement in more categories of motor milestones than worsening), defined as a responder for this primary analysis.

⁵ Not statistically controlled for multiple comparisons

Table 6. Survival Results of Patients with Infantile-Onset SMA (Study 1)

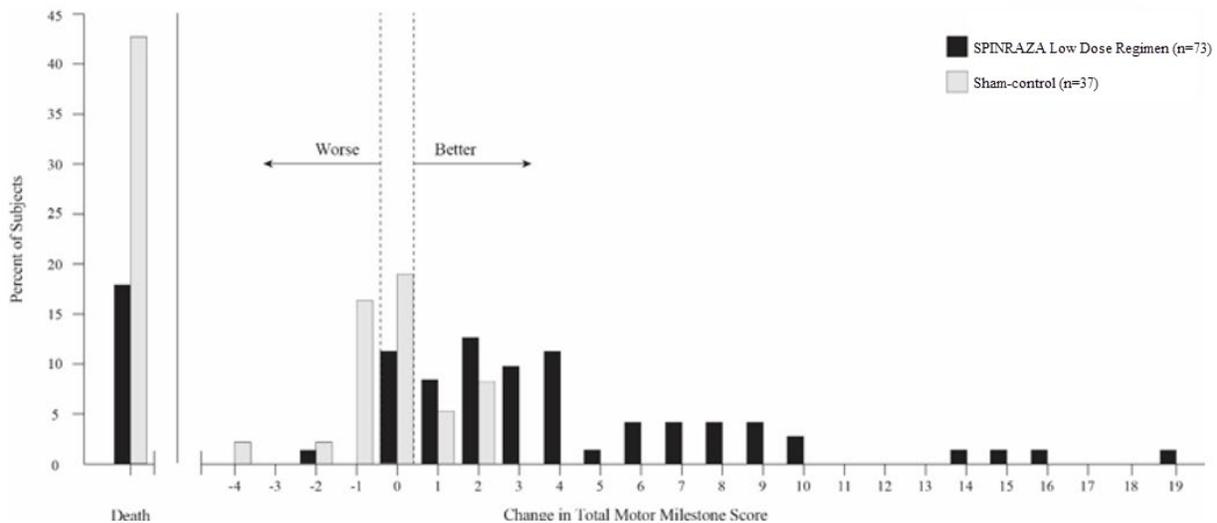
Endpoint	SPINRAZA-treated Patients ¹ (n=80)	Sham-control Patients (n=41)
Survival		
Event-free survival²		
Number of patients who died or received permanent ventilation	31 (39%)	28 (68%)
Hazard ratio (95% CI)	0.53 (0.32 -0.89)	
p-value ³	p=0.005	
Overall survival²		
Number of patients who died	13 (16%)	16 (39%)
Hazard Ratio (95% CI)	0.37 (0.18 – 0.77)	
p-value ³	p=0.004	

¹Low Dose Regimen [see Dosage and Administration (2.1)]

²At the final analysis, event-free survival and overall survival were assessed using the Intent to Treat population (ITT SPINRAZA n=80; Sham-control n=41).

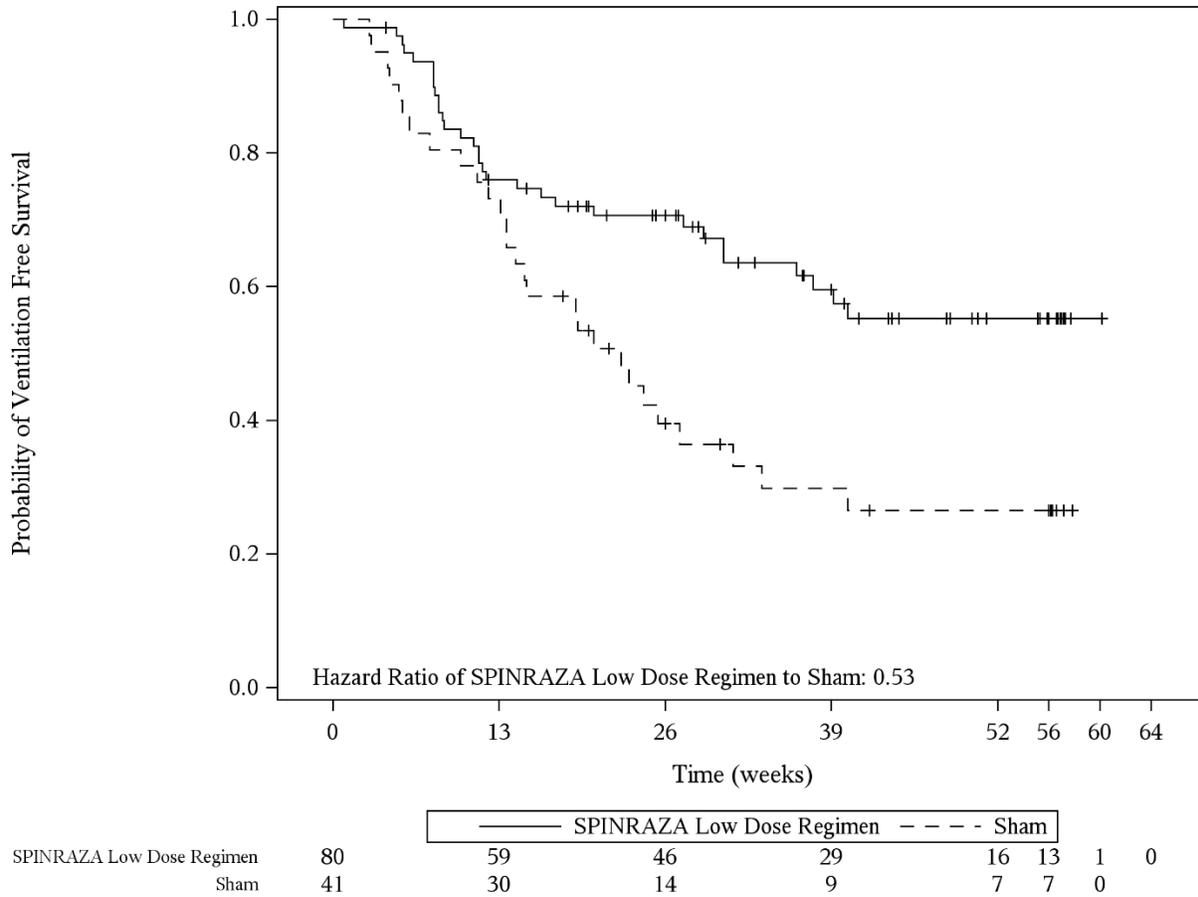
³Based on log-rank test stratified by disease duration

Figure 1. Percent of Patients Who Died and Net Change from Baseline in Total Motor Milestone Score (HINE) Among Patients Alive in the Final Efficacy Set of Study 1 *



*For subjects who were alive and ongoing in the study, the change in total motor milestone score was calculated at the later of Day 183, Day 302, or Day 394.

Figure 2. Event-Free Survival in the Intent to Treat Set



Study 4 (NCT04089566) Part B was a multicenter, double-blind, randomized, controlled study, which included 75 patients with infantile-onset SMA (2 SMN2 copies; symptom onset before 6 months of age). Part B was powered to assess efficacy in infantile-onset patients by evaluating the change in CHOP-INTEND at Day 183 in the patients receiving SPINRAZA High Dose Regimen (n=50) as compared to a prespecified matched sham group from Study 1 (n=20; matched on baseline disease duration and baseline CHOP-INTEND score).

Baseline demographic characteristics of the SPINRAZA group and matched sham group were balanced. Of the 70 patients 50% were male, 64% were Caucasian, and 16% were Asian. Relative to the infantile-onset population in Study 1, patients enrolled in Study 4 had shorter disease duration (time from symptom onset to screening) and lower baseline CHOP-INTEND scores, suggesting they were progressing more quickly and further into their disease course.

A statistically significant improvement in the mean change from baseline in CHOP-INTEND at Day 183 was observed in the SPINRAZA High Dose Regimen group (15.1 point improvement)

compared to the matched sham group (11.1 point worsening) (LS mean difference:26.19 points (95% CI: 20.7, 31.7)) p = <0.0001 (Table 7).

A statistically significantly greater percentage of patients in the SPINRAZA High Dose Regimen group met the HINE-2 responder definition at Day 183 as compared to the matched sham group (58% vs. 0%; p<0.0001).

Table 7. Motor Milestone Response and CHOP-INTEND Results of the Final Analysis of Patients with Infantile-Onset SMA (Study 4)

Efficacy Parameter	SPINRAZA ¹ (n = 50)	Matched Sham from Study 1 (n = 20)	Differences between arms (95% CI)
<u>CHOP-INTEND</u>			
LS mean (95% CI) for ranked score of change from baseline to Day 183	42.9 (38.7, 47.2)	16.9 (10.1, 23.7)	26.06 (17.94, 34.17) p<0.0001 ³
LS mean change (95% CI) from baseline to Day 183 ^{2,3}	15.1 (12.4, 17.8)	-11.1 (-15.9, -6.2)	26.2 (20.7, 31.7) ²
<u>HINE-2 Responder</u>			
Proportion achieving motor milestone responder criteria at Day 183	29 (58%)	0 (0%)	58% (39.5, 71.8) ⁴ p<0.0001

¹ High Dose Regimen [see Dosage and Administration (2.1)]

² ANCOVA and Multiple Imputation applied

³ Least Square Mean difference

⁴ Fisher Exact Test

Similar to Study 1, the SPINRAZA High Dose Regimen group experienced a nominally statistically significant 67% reduction in the risk of death or permanent ventilation relative to the matched sham group (p = 0.0006). The median time to death or permanent ventilation was not reached in the SPINRAZA group and was 19.1 weeks in the matched sham group. Similar observations were seen for overall survival.

14.2 Later-Onset SMA

Study 2 (NCT02292537) was a multicenter, randomized, double-blind, sham-procedure controlled study in 126 symptomatic children with later-onset SMA (symptom onset after 6 months of age). Patients were randomized 2:1 to either SPINRAZA Low Dose Regimen or sham injection as a series of loading doses administered intrathecally followed by maintenance doses administered every 6 months.

The median age at screening was 3 years (range 2-9 years), and the median age of onset of clinical signs and symptoms of SMA was 11 months (range 6-20 months). Of the 126 patients included in the study, 47% were male, 75% were Caucasian, 2% were Black, and 18% were Asian. Length of treatment ranged from 324 to 482 days (median 450 days). At baseline, patients had a mean Hammersmith Functional Motor Scale – Expanded (HFMSSE) score of 21.6, all had

achieved independent sitting, and no patients had achieved independent walking. Patients in this study were deemed most likely to develop Type 2 or 3 SMA.

The primary endpoint assessed was the change from baseline score at Month 15 on the HFMSE. The HFMSE evaluates motor function in patients with SMA who have limited ambulation, comprising of 33 scored activities that give objective information on motor ability and clinical progression, such as the ability to sit unassisted, stand, or walk. Each item is scored from 0-2, with a maximum total score of 66. Higher scores indicate better motor function. The primary analysis was conducted in the Intent to Treat (ITT) population, which included all subjects who were randomized and received at least 1 dose of SPINRAZA or at least one sham procedure. At the final analysis, a statistically significant improvement in HFMSE scores from baseline to Month 15 was observed in the group treated with SPINRAZA Low Dose Regimen compared to the sham-control group (Table 8).

Table 8. HFMSE Results in Patients with Later-Onset SMA (Study 2)

Endpoint	SPINRAZA-treated Patients¹ (n=84)	Sham-control Patients (n=42)
HFMSE score		
Change from baseline in total HFMSE score at 15 months ^{2,3,4}	3.9 (95% CI: 3.0, 4.9) p=0.0000001	-1.0 (95% CI: -2.5, 0.5)
Proportion of patients who achieved at least a 3-point improvement from baseline to Month 15 ²	56.8% (95% CI: 45.6, 68.1) p=0.0006 ⁵	26.3% (95% CI: 12.4, 40.2)

¹Low Dose Regimen [see Dosage and Administration (2.1)]

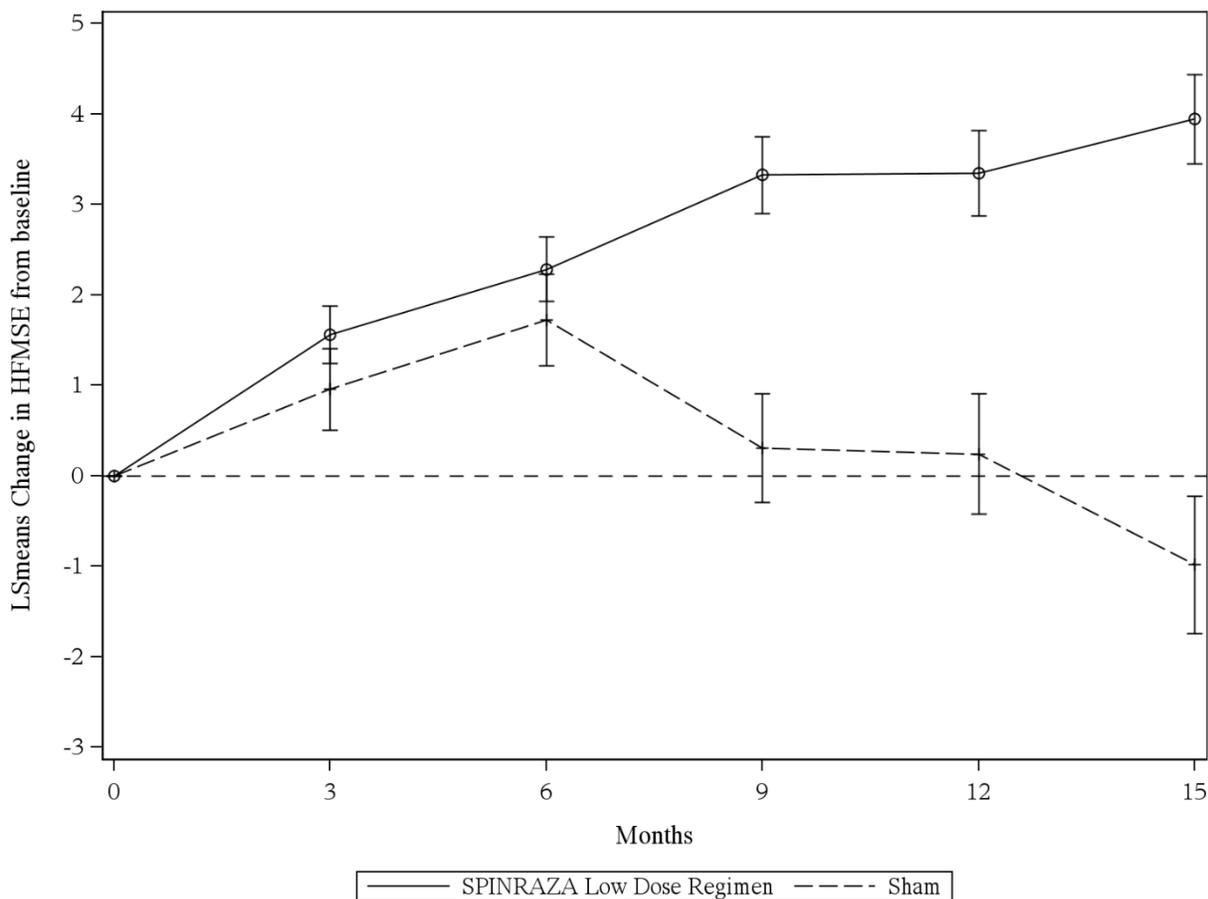
²Assessed using the Intent to Treat population who received at least one dose of SPINRAZA or at least one sham procedure (SPINRAZA n=84; Sham-control n=42); data for patients without a Month 15 visit were imputed using the multiple imputation method

³Least squares mean

⁴Negative value indicates worsening, positive value indicates improvement.

⁵Based on logistic regression with treatment effect and adjustment for each subject's age at screening and HFMSE score at baseline

Figure 3. Mean Change from Baseline in HFMSE Score Over Time in the Intent to Treat Set^{1,2}(Study 2)



¹Data for patients without a Month 15 visit were imputed using the multiple imputation method

²Error bars denote +/- standard error

Study 4 (NCT04089566) Part B was a double-blind, randomized, controlled study, which included 24 patients with later-onset SMA (symptom onset after six months of age) who were randomized 2:1 to receive SPINRAZA High Dose Regimen (n=16) or SPINRAZA Low Dose Regimen treatment (n=8) [see *Dosage and Administration (2.1)*].

Baseline demographic characteristics between all groups were generally balanced.

The efficacy endpoints for the later-onset population in Study 4 were the change from baseline to Day 302 in the HFMSE and in the Revised Upper Limb Module (RULM) score. The RULM test assesses functional ability of the upper limbs in patients with SMA and is comprised of 19 items scored on a 3-point scale. The maximum score is 37 points. The change from baseline to Day 302 in HFMSE score and RULM score showed a slight numerical difference favoring the

SPINRAZA High Dose Regimen, though the comparison was not powered to achieve statistical significance.

14.3 Presymptomatic SMA

The results of the sham-controlled trial in patients with infantile-onset (Study 1) (NCT02193074) and later-onset (Study 2) (NCT02292537) SMA were supported by an open-label uncontrolled trial conducted in 25 patients with presymptomatic SMA who had a genetic diagnosis of 5q SMA and 2 or 3 copies of SMN2 (Study 3) (NCT02386553). In Study 3, 15 patients (60%) who had 2 SMN2 copies, and 10 patients (40%) who had 3 SMN2 copies; 48% were male, 56% were Caucasian, 12% were Asian, 4% were American Indian or Alaska Native, and 28% were of another race, or had no race reported. Patients ranged in age from 3 days to 42 days (median 22 days) at the time of first dose. Patients received SPINRAZA Low Dose Regimen [see *Dosage and Administration (2.1)*]. Patients were assessed with the World Health Organization (WHO) motor milestones, a set of 6 milestones in motor development that would be expected to be attained by 24 months of age in healthy children. An interim analysis was performed after all patients had received SPINRAZA for at least 14 months (median 25 months, range 14 to 34 months). Patients ranged in age from 14 to 34 months (median age of 26 months) at the time of the analysis. At the time of the interim analysis (data cutoff May 2018), all patients receiving SPINRAZA before the onset of SMA symptoms survived without requiring permanent ventilation, and beyond what would be expected based on their SMN2 copy number. All 25 patients (100%) had achieved the WHO motor milestone of sitting without support, and 22 patients (88%) had achieved the milestone of walking with assistance. Of the 22 patients who were older than the age expected to have achieved the ability to walk independently (as defined by the 95th percentile of the WHO expected age of achievement), 17 (77%) achieved the milestone of walking alone (i.e., walking independently).

16 HOW SUPPLIED/STORAGE AND HANDLING

16.1 How Supplied

SPINRAZA injection is a sterile, clear and colorless, preservative-free solution in single-dose glass vials supplied as one vial per carton in the following strengths:

- 12 mg/5 mL (2.4 mg/mL) (NDC 64406-058-01)
- 28 mg/5 mL (5.6 mg/mL) (NDC 64406-036-01)
- 50 mg/5 mL (10 mg/mL) (NDC 64406-037-01)

16.2 Storage and Handling

Store in a refrigerator between 2°C to 8°C (36°F to 46°F) in the original carton to protect from light. Do not freeze.

SPINRAZA should be protected from light and kept in the original carton until time of use.

If no refrigeration is available, SPINRAZA may be stored in its original carton, protected from light at or below 30°C (86°F) for up to 14 days.

Prior to administration, unopened vials of SPINRAZA can be removed from and returned to the refrigerator, if necessary. If removed from the original carton, the total combined time out of refrigeration should not exceed 30 hours at a temperature that does not exceed 25°C (77°F).

17 PATIENT COUNSELING INFORMATION

Thrombocytopenia and Coagulation Abnormalities

Inform patients and caregivers that SPINRAZA could increase the risk of bleeding. Inform patients and caregivers of the importance of obtaining blood laboratory testing at baseline and prior to each dose to monitor for signs of increased potential for bleeding. Instruct patients and caregivers to seek medical attention if unexpected bleeding occurs [*see Warnings and Precautions (5.1)*].

Renal Toxicity

Inform patients and caregivers that SPINRAZA could cause renal toxicity. Inform patients and caregivers of the importance of obtaining urine testing at baseline and prior to each dose to monitor for signs of potential renal toxicity [*see Warnings and Precautions (5.2)*].

60110-01

Manufactured for:

Biogen

Cambridge, MA 02142

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