UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
WASHINGTON, D.C. 20549

FORM 8-K

CURRENT REPORT

Pursuant to Section 13 or 15(d) of the Securities Exchange Act of 1934

Date of Report (Date of earliest event reported): November 05, 2026

Sarepta Therapeutics, Inc.

(Exact name of Registrant as Specified in Its Charter)

Delaware	**001-14895**	**93-0797222**
(State or Other Jurisdiction of Incorporation)	**(Commission File Number)**	**(IRS Employer Identification No.)**
215 First Street Cambridge, Massachusetts		**02142**
(Address of Principal Executive Offices)		**(Zip Code)**

Registrant's Telephone Number, Including Area Code: (617) 274-4000

(Former Name or Former Address, if Changed Since Last Report)

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:

☐ Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)

☐ Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)

☐ Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))

☐ Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

Securities registered pursuant to Section 12(b) of the Act:

Title of each class	Trading Symbol(s)	Name of each exchange on which registered
Common Stock, $0.0001 par value per share	SRPT	Nasdaq Global Select Market

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (§ 230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§ 240.12b-2 of this chapter).

Emerging growth company ☐

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act. ☐

Item 2.02 Results of Operations and Financial Condition.

On November 5, 2026, Sarepta Therapeutics, Inc. issued a press release announcing its financial results for the quarter ended September 30, 2026. The full text of the press release is furnished as Exhibit 99.1 to this Current Report on Form 8-K.

The information in Item 2.02 of this Form 8-K (including Exhibit 99.1) shall not be deemed "filed" for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the "Exchange Act"), or otherwise subject to the liabilities of that section, nor shall it be deemed incorporated by reference in any filing under the Securities Act of 1933, as amended, or the Exchange Act, except as expressly set forth by specific reference in such a filing.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits

Exhibit Number	Description
99.1	Press Release dated November 5, 2026
104	The cover page from this Current Report on Form 8-K of Sarepta Therapeutics, Inc., formatted in Inline XBRL and included as Exhibit 101

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

Sarepta Therapeutics, Inc.

Date: November 5, 2026

By: /s/ Michael Severino, M.D.
Michael Severino, M.D.
Chief Executive Officer

Sarepta Therapeutics Announces Third Quarter 2026 Financial Results and Recent Corporate Developments

- ***Upcoming multiple ascending dose data from SRP-1003 in DM1 and SRP-1001 in FSHD expected by year end; Company is actively preparing for Phase 3 development in both programs***
- ***Net product revenues for the third quarter 2026 totaled approximately $323.8 million, consisting of approximately $240.3 million of PMO net product revenues and $83.5 million of ELEVIDYS® net product revenue***
- ***Continued disciplined financial execution with positive cash flow and GAAP operating loss of ($9.0) million and non-GAAP operating income of $22.9 million for the third quarter 2026***
- ***Updates to 2026 guidance range include raising lower end of total net product revenue by $50.0 million, driven by strong PMO performance, and lowering higher end of expense by $25.0 million, reflecting operating discipline***
- ***Growing evidence base across ambulatory population reinforces ELEVIDYS benefit-risk profile, with new clinical, biomarker and real-world data supporting durability of benefit, robust dystrophin expression and approaches to liver safety risk mitigation***

CAMBRIDGE, Mass. -- (BUSINESS WIRE) -- Nov. 5, 2026 -- Sarepta Therapeutics, Inc. (NASDAQ:SRPT), the leader in precision genetic medicine for rare diseases, today reported financial results for the third quarter of 2026.

“After my first 90 days at Sarepta, I believe the fundamental strengths that attracted me to the company remain firmly intact: a durable Duchenne franchise, a promising pipeline with the potential to deliver differentiated medicines, and the financial strength to invest in our highest-value opportunities,” said Michael Severino, M.D., chief executive officer, Sarepta. “This quarter, we saw continued strength in our commercial execution. Our upcoming DM1 and FSHD datasets are important opportunities to demonstrate the depth of knockdown and downstream biological effects we believe our platform can achieve. We are already preparing both programs for potential late-stage development with the intent of moving efficiently if the results support advancement. We have work to do, and I want us to be measured by execution, but I believe Sarepta has the commercial foundation, pipeline opportunities and financial strength to deliver meaningful long-term value.”

Corporate Highlights:

- **DM1 and FSHD programs advancing toward MAD data in fourth quarter of 2026 and preparing for late-stage development:** Multiple ascending dose data from SRP-1003 and SRP-1001 are expected in the fourth quarter of 2026, providing important insights including DMPK knockdown and splicing correction in DM1 and DUX4 knockdown in FSHD. Sarepta is actively preparing for Phase 3 development in both programs, including beginning

manufacturing of clinical supply. The Company is preparing for both accelerated and traditional approval potential pathways, with the ultimate regulatory strategy to be informed by the data and discussions with FDA.

- **SRP-1003 demonstrates robust nuclear DMPK knockdown in DM1:** New preclinical data presented at the 2026 Oligonucleotide Therapeutics Society Annual Meeting demonstrated that SRP-1003 reduced pathogenic DMPK mRNA in both the nucleus and cytoplasm in a dose-dependent manner, achieving comparable knockdown in each compartment. Nuclear target engagement was supported by three independent methodologies, with correction of MBNL-dependent mis-splicing providing functional evidence that SRP-1003 is degrading disease-causing RNA within the nucleus.
- **Financial strength supports late-stage DM1 and FSHD execution:** Delivered another quarter of non-GAAP operating profit and positive cash flow, ending with approximately $998.7 million in cash, cash equivalents, restricted cash and investments. Our balance-sheet strength and commercial cash flows are expected to provide the resources to fund these programs through late-stage development while maintaining disciplined capital allocation.
- **2026 Guidance:** Company expects ELEVIDYS net product revenue in the fourth quarter to trend in line with the third quarter, consistent with previous guidance and subject to normal variability associated with one-time gene therapy. The Company expects the PMO franchise to remain strong and is updating its full-year guidance ranges, raising the low end of total net product revenue guidance by $50.0 million to a range of $1.25 billion to $1.3 billion. The Company also narrows its full-year combined non-GAAP R&D and SG&A expenses, lowering the high end of expense guidance by $25.0 million to a range of $800 million to $825 million.
- **Growing evidence base reinforces the ELEVIDYS benefit-risk profile:** At the 2026 World Muscle Society Annual Congress, Company presented ELEVIDYS data in ambulatory patients ages 8 to 12 at the time of treatment demonstrating durable and clinically meaningful benefit across the North Star Ambulatory Assessment and timed function measures over two years compared with a well-matched external control group. Company also presented data in patients treated at ages 2 and 3 demonstrating robust ELEVIDYS-dystrophin expression and sarcolemmal localization 12 weeks after infusion. At the Annual Neuromuscular Study Group Scientific Meeting, initial results from the Phase 4 ENDURE study provided preliminary real-world data evaluating approaches to liver safety risk mitigation.

Conference Call

The event will be webcast live under the investor relations section of Sarepta's website at **https://investorrelations.sarepta.com/events-presentations** and following the event a replay will be archived there for one year. This event can be accessed using **this link.**

Q3 2026 Financial Highlights[1]

	For the Three Months Ended September 30		QTD Change	
	2026	2025		
	(in millions, except for per share amounts)		$	%
Total revenues	$ 381.4	$ 399.4	$ (18.0)	(5)%
Operating (loss) income:				
GAAP	$ (9.0)	$ (103.4)	$ 94.4	(91)%
Non-GAAP	$ 22.9	$ (35.7)	$ 58.6	*
Net (loss) income:				
GAAP[2]	$ (28.5)	$ (50.6)	$ 22.1	(44)%
Non-GAAP	$ 15.6	$ (12.9)	$ 28.5	*
Diluted (loss) earnings per share				
GAAP[2]	$ (0.27)	$ (0.50)	$ 0.23	(46)%
Non-GAAP	$ 0.13	$ (0.13)	$ 0.26	*

	For the Nine Months Ended September 30		YTD Change	
	2026	2025		
	(in millions, except for per share amounts)		$	%
Total revenues	$ 1,513.4	$ 1,755.3	$ (241.9)	(14)%
Operating income (loss):				
GAAP	$ 362.7	$ (288.2)	$ 650.9	*
Non-GAAP	$ 507.1	$ (122.5)	$ 629.6	*
Net income (loss):				
GAAP[2]	$ 297.6	$ (301.2)	$ 598.8	*
Non-GAAP	$ 479.6	$ (130.2)	$ 609.8	*
Diluted earnings (loss) per share				
GAAP[2]	$ 2.78	$ (3.06)	$ 5.84	*
Non-GAAP	$ 3.91	$ (1.32)	$ 5.23	*

[1] For an explanation of our use of non-GAAP financial measures, please refer to the "Use of Non-GAAP Financial Measures" section later in this press release, and for a reconciliation of each non-GAAP financial measure from the most comparable GAAP measures, see the table at the end of this press release.

[2] GAAP net loss and diluted loss per share for the three and nine months ended September 30, 2025 have been revised to reflect the correction of an immaterial error associated with the accounting for the August 2025 Exchange, as well as the adoption of ASU 2025-07. Please refer to the Company's Annual Report on Form 10-K for the year ended December 31, 2025 for additional information.

*Not meaningful

	As of September 30, 2026	As of December 31, 2025
	(in millions)	
Cash, cash equivalents, restricted cash and investments	$ 998.7	$ 953.8

Revenues

Total revenues were $381.4 million for the three months ended September 30, 2026, as compared to $399.4 million for the same period of 2025, a decrease of $18.0 million. This primarily reflects a lower volume of ELEVIDYS sales due to the continued impact of our updated label that only includes the ambulatory patient population for treatment. The decrease is partially offset by an increase of $31.4 million in contract manufacturing revenues associated with increased commercial ELEVIDYS supply delivered to Roche.

Total revenues were $1,513.4 million for the nine months ended September 30, 2026, as compared to $1,755.3 million for the same period of 2025, a decrease of $241.9 million. This primarily reflects a lower volume of ELEVIDYS sales due to our updated label that only includes the ambulatory patient population for treatment. The decrease is partially offset by an increase of $189.5 million in collaboration revenues related to the $365.0 million of collaboration revenue recognized related to Roche's declined option for certain program rights and the milestone recognized under the Roche collaboration agreement for the first commercial dosing of ELEVIDYS in Japan during the nine months ended September 30, 2026, as compared to $175.5 million of collaboration revenue in 2025 related to Roche’s expiration of an option to acquire a certain program and the Japan Approval Milestone. Furthermore, contract manufacturing revenues increased $72.5 million associated with increased commercial ELEVIDYS supply delivered to Roche.

Cost of sales (excluding amortization of in-licensed rights)

Cost of sales (excluding amortization of in-license rights) were $188.5 million for the three months ended September 30, 2026, as compared to $150.8 million for the same period of 2025, an increase of $37.7 million. Cost of sales (excluding amortization of in-license rights) were $446.7 million for the nine months ended September 30, 2026, as compared to $440.9 million for the same period of 2025, an increase of $5.8 million. The increases in both periods primarily reflect a take-or-pay charge of $62.0 million related to non-cancelable manufacturing obligations with Catalent, Inc. ("Catalent") for certain inventories that we elected not to manufacture. Additionally, there was an increase in cost of sales related to products sold to Roche, primarily related to increased volume of ELEVIDYS shipments as well as write-offs of certain batches of products not meeting quality specifications under the Roche collaboration agreement. The increases in both periods were partially offset by a lower volume of ELEVIDYS sales in the United States and corresponding royalty payments.

Operating expenses and others

Research and development expenses were $82.5 million for the three months ended September 30, 2026, as compared to $218.9 million for the same period of 2025, a decrease of $136.4 million. The decrease primarily reflects a $100.0 million milestone payment made to Arrowhead Pharmaceutical, Inc. ("Arrowhead") during the three months ended September 30, 2025, triggered by Arrowhead's achievement of the first of two predetermined enrollment targets and subsequent authorization to dose escalate in a Phase 1/2 study for the DM1 program, with no similar activity during the same period of 2026. In addition, there were decreases in manufacturing and clinical expenses primarily due to our decision to reprioritize our pipeline and developmental priorities announced in July 2025, partially offset by an increase in compensation and other personnel expenses due to the reversal of previously recognized

expense related to certain incentive compensationour restructuring plan announced in July 2025 (the "Restructuring"). For the three months ended September 30, 2026, non-GAAP research and development expenses were $71.6 million, as compared to $206.5 million for the same period of 2025, a decrease of $134.9 million.

Research and development expenses were $327.8 million for the nine months ended September 30, 2026, as compared to $1,196.7 million for the same period of 2025, a decrease of approximately $868.9 million. The decrease primarily reflects the recognition of up-front and collaboration license fees of $583.6 million associated with the licensing, collaboration and stock purchase agreement with Arrowhead as well as the $100.0 million milestone payment made to Arrowhead during the nine months ended September 30, 2025, with no similar activity for the same period of 2026. In addition, there was a decrease in manufacturing and clinical expenses primarily due to our decision to reprioritize our pipeline and developmental priorities announced in July 2025, as well as a decrease in compensation, other personnel, and stock-based compensation expenses, all as a result of the Restructuring. This decrease was partially offset by the $50.0 million annual collaboration license fee incurred and paid to Arrowhead during the nine months ended September 30, 2026. For the nine months ended September 30, 2026, non-GAAP research and development expenses were $285.8 million, as compared to $1,137.4 million for the same period of 2025, a decrease of $851.6 million.

Selling, general and administrative expenses were $118.6 million for the three months ended September 30, 2026, as compared to $91.9 million for the same period of 2025, an increase of $26.7 million. The increase primarily reflects the reversal of previously recognized expense related to certain stock compensation awards and incentive compensation accruals in connection with the Restructuring during the three months ended September 30, 2025, with no similar activity during the same period of 2026. For the three months ended September 30, 2026, non-GAAP selling, general and administrative expenses were $97.6 million, as compared to $77.1 million for the same period of 2025, an increase of $20.5 million.

Selling, general and administrative expenses were $335.2 million for the nine months ended September 30, 2026, as compared to $363.4 million for the same period of 2025, a decrease of $28.2 million. The decrease primarily reflects a decrease in compensation, other personnel, and stock-based compensation expenses, all as a result of the Restructuring, as well as a decrease in professional services used related to ELEVIDYS commercialization efforts. For the nine months ended September 30, 2026, non-GAAP selling, general and administrative expenses were $271.7 million, as compared to $297.6 million for the same period of 2025, a decrease of $25.9 million.

Litigation settlement charge was $39.0 million for the nine months ended September 30, 2026, with no similar activity for the same period of 2025. We recorded a litigation settlement charge of $39.0 million in the second quarter of 2026 related to a settlement agreement that was executed in September 2026 between Sarepta Therapeutics, Inc., Sarepta Therapeutics Three, LLC, REGENXBIO, INC, the Trustees of the University of Pennsylvania and Catalent regarding the resolution of certain patent infringement litigations.

Restructuring charges were $40.5 million for the three and nine months ended September 30, 2025, with no similar activity for the same periods of 2026.

Other (expense) income, net for the three months ended September 30, 2026 and 2025 was approximately $(16.7) million and $35.0 million, respectively. Other expense, net for the nine months ended September 30, 2026 and 2025 was approximately $47.0 million and $10.1 million, respectively. The change primarily reflects a decrease in our strategic investments as a result of the sale of our investment in Arrowhead in August 2025, partially offset by an increase in interest expense due to our 2030 Notes carrying a higher interest rate than our 2027 Notes during the three and nine months ended September 30, 2026.

Gain on debt extinguishment was $3.8 million for the three and nine months ended September 30, 2025, with no similar activity for the same periods of 2026. This gain was the result of a partial refinancing of the 2027 Notes where we exchanged $700.0 million in aggregate principal amount of 2027 Notes for $602.0 million in aggregate principal amount of 2030 Notes and made cash payments of $127.3 million, including $4.0 million of accrued interest of the 2027 Notes and 5.9 million shares of our common stock with a fair market value of approximately $104.9 million, net of issuance costs of $2.4 million.

Income tax expense for the three and nine months ended September 30, 2026, was approximately $2.7 million and $18.1 million, respectively. Income tax (benefit) expense for the three and nine months ended September 30, 2025, was $(14.1) million and $6.6 million, respectively. Income tax expense for all periods presented primarily relates to state income taxes as a result of taxable profits in certain states requiring the capitalization of research and development costs and states which have suspended or limited the utilization of net operating loss carryforwards.

Use of Non-GAAP Financial Measures

In addition to the GAAP financial measures set forth in this press release, we have included the following non-GAAP measurements:

1. Non-GAAP net (loss) income is defined by us as GAAP net (loss) income excluding interest expense/income, net, depreciation and amortization expense, stock-based compensation expense, restructuring charges, other items, and the estimated income tax impact of each pre-tax non-GAAP adjustment.
2. Non-GAAP earnings per share is defined by us as non-GAAP net income, as defined previously, divided by the weighted-average number of shares of common stock and dilutive common stock equivalents outstanding, adjusted for the inclusion of additional shares under both the treasury stock method and the “if-converted” method, if applicable and not anti-dilutive. Non-GAAP net loss per share is defined by us as non-GAAP net loss, as defined above, divided by the weighted-average number of shares of common stock outstanding as the inclusion of dilutive common stock equivalents outstanding is anti-dilutive.
3. Non-GAAP operating income (loss) is defined by us as GAAP operating (loss) income excluding depreciation and amortization expense, stock-based compensation expense, restructuring charges and litigation settlement charges.

4. Non-GAAP research and development expenses are defined by us as GAAP research and development expenses excluding depreciation and amortization expense and stock-based compensation expense.
5. Non-GAAP selling, general and administrative expenses are defined by us as GAAP selling, general and administrative expenses excluding depreciation expense and stock-based compensation expense.
6. Non-GAAP effective tax rate is defined by us as the GAAP effective tax rate excluding the impact of our GAAP to non-GAAP adjustments.

The following components are used to adjust our GAAP financial measures into the previously defined non-GAAP measurements:

1. Interest, depreciation and amortization - Interest expense/income, net amounts can vary substantially from period to period due to changes in cash and debt balances and interest rates driven by market conditions outside of our operations. Depreciation expense can vary substantially from period to period as the purchases of property and equipment may vary significantly from period to period and without any direct correlation to our operating performance. Amortization expense primarily associated with patent costs are amortized over a period of several years after acquisition or patent application or renewal.
2. Stock-based compensation expenses - Stock-based compensation expenses represent non-cash charges related to equity awards we have granted. Although these are recurring charges to operations, we believe the measurement of these amounts can vary substantially from period to period and depend significantly on factors that are not a direct consequence of operating performance that is within our control. Therefore, we believe that excluding these charges facilitates comparisons of our operational performance in different periods.
3. Restructuring charges do not have a direct correlation to future business operations, nor do the resulting charges recorded reflect the performance of our ongoing operations for the period in which such charges are recorded. In addition, restructuring charges are not considered to be normal operating expenses due to the variability of amounts and lack of predictability as to occurrence and/or timing.
4. Other items - We evaluate other items of expense and income on an individual basis. We take into consideration quantitative and qualitative characteristics of each item, including (a) nature, (b) whether the items relate to our ongoing business operations, and (c) whether we expect the items to continue or occur on a regular basis. These other items include the loss (gain) on strategic investments, the impairment of strategic investments, litigation settlement charges, and gain on debt extinguishment and may include other items that fit the above characteristics in the future. We exclude from our non-GAAP results:
    a) The loss (gain) on strategic investments as the results of such gains and losses are not representative of our normal business operations, which accordingly would make it difficult to compare our results to peer companies that also provide non-GAAP disclosures.
    b) The impairment of strategic investments as such charges are not indicative of the performance of our core operations, which accordingly would make it difficult to compare our results to peer companies that also provide non-GAAP disclosures.
    c) Litigation settlement charge as such charges are considered to be an infrequent event as it is associated with a distinct, non-recurring litigation matter and is not indicative

of the performance of our core operations nor representative of our normal business operations, which accordingly would make it difficult to compare our results to peer companies that also provide non-GAAP disclosures.

d) The gain on debt extinguishment is considered to be an infrequent event as it is associated with a distinct financing decision and is not indicative of the performance of our core operations, which accordingly would make it difficult to compare our results to peer companies that also provide non-GAAP disclosures.

We use these non-GAAP measures as key performance measures for the purpose of evaluating operational performance and cash requirements internally. We also believe these non-GAAP measures increase comparability of period-to-period results and are useful to investors as they provide a similar basis for evaluating our performance as is applied by management. These non-GAAP measures are not intended to be considered in isolation or to replace the presentation of our financial results in accordance with GAAP. Use of the terms non-GAAP research and development expenses, non-GAAP selling, general and administrative expenses, non-GAAP operating income (loss), non-GAAP net income (loss), and non-GAAP diluted earnings (loss) per share may differ from similar measures reported by other companies, which may limit comparability, and are not based on any comprehensive set of accounting rules or principles. All relevant non-GAAP measures are reconciled from their respective GAAP measures in the attached table "Reconciliation of GAAP Financial Measures to Non-GAAP Financial Measures."

About EXONDYS 51

EXONDYS 51 uses Sarepta's proprietary phosphorodiamidate morpholino oligomer (PMO) chemistry and exon-skipping technology to bind to exon 51 of dystrophin pre-mRNA, resulting in exclusion, or "skipping", of this exon during mRNA processing in patients with genetic mutations that are amenable to exon 51 skipping. Exon skipping is intended to allow for production of an internally truncated dystrophin protein.

EXONDYS 51 is indicated for the treatment of Duchenne muscular dystrophy (DMD) in patients who have a confirmed mutation of the *DMD* gene that is amenable to exon 51 skipping. This indication is approved under accelerated approval based on an increase in dystrophin in skeletal muscle observed in some patients treated with EXONDYS 51. Continued approval for this indication may be contingent upon verification of a clinical benefit in confirmatory trials.

EXONDYS 51 has met the full statutory standards for safety and effectiveness and as such is not considered investigational or experimental.

Important Safety Information About EXONDYS 51

Hypersensitivity reactions, including bronchospasm, chest pain, cough, tachycardia, and urticaria have occurred in patients who were treated with EXONDYS 51. If a hypersensitivity reaction occurs, institute appropriate medical treatment and consider slowing the infusion or interrupting the EXONDYS 51 therapy.

Adverse reactions in DMD patients (N=8) treated with EXONDYS 51 30 mg or 50 mg/kg/week by intravenous (IV) infusion with an incidence of at least 25% more than placebo (N=4) (Study 1, 24 weeks) were (EXONDYS 51, placebo): balance disorder (38%, 0%), vomiting (38%, 0%) and contact dermatitis (25%, 0%). The most common adverse reactions were balance disorder and vomiting. Because of the small numbers of patients, these represent crude frequencies that may not reflect the frequencies observed in practice. The 50 mg/kg once weekly dosing regimen of EXONDYS 51 is not recommended.

The most common adverse reactions from observational clinical studies (N=163) seen in greater than 10% of patients were headache, cough, rash, and vomiting.

Other adverse events may occur.

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.

For further information, please see the full U.S. Prescribing Information for EXONDYS 51 (eteplirsen).

About VYONDYS 53

VYONDYS 53 (golodirsen) uses Sarepta's proprietary phosphorodiamidate morpholino oligomer (PMO) chemistry and exon-skipping technology to bind to exon 53 of dystrophin pre-mRNA, resulting in exclusion, or "skipping," of this exon during mRNA processing in patients with genetic mutations that are amenable to exon 53 skipping. Exon skipping is intended to allow for production of an internally truncated dystrophin protein.

VYONDYS 53 is indicated for the treatment of Duchenne muscular dystrophy (DMD) in patients who have a confirmed mutation of the *DMD* gene that is amenable to exon 53 skipping. This indication is approved under accelerated approval based on an increase in dystrophin production in skeletal muscle observed in patients treated with VYONDYS 53. Continued approval for this indication may be contingent upon verification of a clinical benefit in confirmatory trials.

VYONDYS 53 has met the full statutory standards for safety and effectiveness and as such is not considered investigational or experimental.

Important Safety Information for VYONDYS 53

CONTRAINDICATIONS: VYONDYS 53 is contraindicated in patients with a serious hypersensitivity reaction to golodirsen or to any of the inactive ingredients in VYONDYS 53. Anaphylaxis has occurred in patients receiving VYONDYS 53.

WARNINGS AND PRECAUTIONS

Hypersensitivity Reactions: Hypersensitivity reactions, including anaphylaxis, rash, pyrexia, pruritus, urticaria, dermatitis, and skin exfoliation have occurred in VYONDYS 53-treated patients, some requiring treatment. If a hypersensitivity reaction occurs, institute appropriate medical treatment and consider slowing the infusion, interrupting, or discontinuing the VYONDYS 53 therapy and monitor until the condition resolves. VYONDYS 53 is contraindicated in patients with a history of a serious hypersensitivity reaction to golodirsen or to any of the inactive ingredients in VYONDYS 53.

Kidney Toxicity: Kidney toxicity was observed in animals who received golodirsen. Although kidney toxicity was not observed in the clinical studies with VYONDYS 53, the clinical experience with VYONDYS 53 is limited, and kidney toxicity, including potentially fatal glomerulonephritis, has been observed after administration of some antisense oligonucleotides. Kidney function should be monitored in patients taking VYONDYS 53. Because of the effect of reduced skeletal muscle mass on creatinine measurements, creatinine may not be a reliable measure of kidney function in DMD patients. Serum cystatin C, urine dipstick, and urine protein-to-creatinine ratio should be measured before starting VYONDYS 53. Consider also measuring glomerular filtration rate using an exogenous filtration marker before starting VYONDYS 53. During treatment, monitor urine dipstick every month, and serum cystatin C and urine protein-to-creatinine ratio every three months. Only urine expected to be free of excreted VYONDYS 53 should be used for monitoring of urine protein. Urine obtained on the day of VYONDYS 53 infusion prior to the infusion, or urine obtained at least 48 hours after the most recent infusion, may be used. Alternatively, use a laboratory test that does not use the reagent pyrogallol red, as this reagent has the potential to cross react with any VYONDYS 53 that is excreted in the urine and thus lead to a false positive result for urine protein.

If a persistent increase in serum cystatin C or proteinuria is detected, refer to a pediatric nephrologist for further evaluation.

ADVERSE REACTIONS: Adverse reactions observed in at least 20% of treated patients and greater than placebo were (VYONDYS 53, placebo): headache (41%, 10%), pyrexia (41%, 14%), fall (29%, 19%), abdominal pain (27%, 10%), nasopharyngitis (27%, 14%), cough (27%, 19%), vomiting (27%, 19%), and nausea (20%, 10%).

Other adverse reactions that occurred at a frequency greater than 5% of VYONDYS 53-treated patients and at a greater frequency than placebo were: administration site pain, back pain, pain, diarrhea, dizziness, ligament sprain, contusion, influenza, oropharyngeal pain, rhinitis, skin abrasion, ear infection, seasonal allergy, tachycardia, catheter site related reaction, constipation, and fracture.

Other adverse events may occur.

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.

For further information, please see the full U.S. Prescribing Information for VYONDYS 53 (golodirsen).

About AMONDYS 45

AMONDYS 45 (casimersen) uses Sarepta's proprietary phosphorodiamidate morpholino oligomer (PMO) chemistry and exon-skipping technology to bind to exon 45 of dystrophin pre-mRNA, resulting in exclusion, or "skipping," of this exon during mRNA processing in patients with genetic mutations that are amenable to exon 45 skipping. Exon skipping is intended to allow for production of an internally truncated dystrophin protein.

AMONDYS 45 is indicated for the treatment of Duchenne muscular dystrophy (DMD) in patients who have a confirmed mutation of the *DMD* gene that is amenable to exon 45 skipping. This indication is approved under accelerated approval based on an increase in dystrophin production in skeletal muscle observed in patients treated with AMONDYS 45. Continued approval for this indication may be contingent upon verification of a clinical benefit in confirmatory trials.

AMONDYS 45 has met the full statutory standards for safety and effectiveness and as such is not considered investigational or experimental.

Important Safety Information for AMONDYS 45

CONTRAINDICATION: AMONDYS 45 is contraindicated in patients with a known serious hypersensitivity to casimersen or any of the inactive ingredients in AMONDYS 45. Instances of hypersensitivity including angioedema and anaphylaxis have occurred.

WARNINGS AND PRECAUTIONS

Hypersensitivity: Hypersensitivity reactions, including angioedema and anaphylaxis, have occurred in patients who were treated with AMONDYS 45. If a hypersensitivity reaction occurs, institute appropriate medical treatment, and consider slowing the infusion, interrupting, or discontinuing the AMONDYS 45 infusion and monitor until the condition resolves. AMONDYS 45 is contraindicated in patients with known serious hypersensitivity to casimersen or to any of the inactive ingredients in AMONDYS 45.

Kidney Toxicity: Kidney toxicity was observed in animals who received casimersen. Although kidney toxicity was not observed in the clinical studies with AMONDYS 45, kidney toxicity, including potentially fatal glomerulonephritis, has been observed after administration of some antisense oligonucleotides. Kidney function should be monitored in patients taking AMONDYS 45. Because of the effect of reduced skeletal muscle mass on creatinine measurements, creatinine may not be a reliable measure of kidney function in DMD patients. Serum cystatin C, urine dipstick, and urine protein-to-creatinine ratio should be measured before starting AMONDYS 45. Consider also measuring glomerular filtration rate using an exogenous filtration marker before starting AMONDYS 45. During treatment, monitor urine dipstick every month, and serum cystatin C and urine protein to-creatinine ratio (UPCR) every three months. Only urine expected to be free of excreted AMONDYS 45 should be used for monitoring of urine protein. Urine obtained on the day of AMONDYS 45 infusion prior to the infusion, or urine obtained at least 48 hours after the most recent infusion, may be used. Alternatively, use a laboratory test that does not use the reagent pyrogallol red, as this reagent has the potential to cross react with any AMONDYS 45 that is excreted in the urine and thus lead to a false positive result for urine protein.

If a persistent increase in serum cystatin C or proteinuria is detected, refer to a pediatric nephrologist for further evaluation.

Adverse Reactions: Adverse reactions occurring in at least 20% of patients treated with AMONDYS 45 and at least 5% more frequently than in the placebo group were (AMONDYS 45, placebo): upper respiratory infections (65%, 55%), cough (33%, 26%), pyrexia (33%, 23%), headache (32%, 19%), arthralgia (21%, 10%), and oropharyngeal pain (21%, 7%).

Other adverse reactions that occurred in at least 10% of patients treated with AMONDYS 45 and at least 5% more frequently than in the placebo group were: ear pain, nausea, ear infection, post-traumatic pain, and dizziness and light-headedness.

Other adverse events may occur.

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.

For further information, please see the full U.S. Prescribing Information for AMONDYS 45 (casimersen).

About ELEVIDYS (delandistrogene moxeparvovec-rokl)

ELEVIDYS (delandistrogene moxeparvovec-rokl) is a single-dose, adeno-associated virus (AAV)-based gene transfer therapy for intravenous infusion designed to address the underlying genetic cause of Duchenne muscular dystrophy – mutations or changes in the DMD gene that result in the lack of dystrophin protein – through the delivery of a transgene that codes for the targeted production of ELEVIDYS micro-dystrophin in skeletal muscle.

ELEVIDYS is indicated for the treatment of ambulatory patients 4 years of age and older with Duchenne muscular dystrophy (DMD) who have a confirmed mutation in the *DMD* gene.

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.

IMPORTANT SAFETY INFORMATION

BOXED WARNING: Acute Serious Liver Injury and Acute Liver Failure

Acute serious liver injury, including life-threatening and fatal acute liver failure, has occurred. Patients with preexisting liver impairment may be at higher risk.

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.

Instruct patients to maintain proximity to an appropriate healthcare facility, as determined by the healthcare provider, for at least 2 months following ELEVIDYS infusion.

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

CONTRAINDICATION: 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.

WARNINGS AND PRECAUTIONS:

Acute Serious Liver Injury and Acute Liver Failure

See *Boxed Warning.*

- 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 of ELEVIDYS administration. 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 have been reported following acute liver injury associated with ELEVIDYS in a non-ambulatory patient.
- 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.
- Systemic corticosteroid treatment is recommended for patients before and after ELEVIDYS infusion. Adjust corticosteroid regimen when indicated.

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.
- 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.
- Avoid administration of ELEVIDYS to patients with active infections.

Myocarditis

- Acute, serious, life-threatening myocarditis and troponin-I elevations have been observed within 24 hours to more than 1 year following ELEVIDYS infusion.
- If a patient experiences myocarditis, those with pre-existing left ventricle ejection fraction (LVEF) impairment may be at higher risk of adverse outcomes.
- Monitor troponin-I before ELEVIDYS infusion and weekly for the first month following infusion and 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.

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. If symptoms of infusion-related reactions occur, slow or stop the infusion and give appropriate treatment. Once symptoms resolve, the infusion may be restarted at a lower rate.
- ELEVIDYS should be administered in a setting where treatment for infusion-related reactions is immediately available.
- Discontinue infusion for anaphylaxis.

Immune-mediated Myositis

- Immune-mediated myositis, including serious and life-threatening events, has occurred approximately 1 month following ELEVIDYS infusion. Signs and symptoms include severe muscle weakness, including dysphagia, dyspnea, dysphonia, and hypophonia.
- 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.
- 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.

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

ADVERSE REACTIONS

- The most common adverse reactions (incidence ≥5%) reported in clinical studies were vomiting, nausea, liver injury, pyrexia, thrombocytopenia, and troponin-I increased.

Report negative side effects of prescription drugs to the FDA. Visit www.fda.gov/medwatch or call 1-800-FDA-1088. You may also report side effects to Sarepta Therapeutics at 1-888-SAREPTA (1-888-727-3782).

Please see the full Prescribing Information for ELEVIDYS, including Boxed Warning and Medication Guide.

About Sarepta Therapeutics

Sarepta is on an urgent mission: engineer precision genetic medicine for rare diseases that devastate lives and cut futures short. We hold a leadership position in Duchenne muscular dystrophy (Duchenne) and are

building a robust portfolio of programs across muscle, central nervous system, and cardiac diseases. For more information, please visit www.sarepta.com or follow us on LinkedIn, X, Instagram and Facebook.

Forward-Looking Statements

In order to provide Sarepta's investors with an understanding of its current results and future prospects, this press release contains statements that are forward-looking. Any statements contained in this press release that are not statements of historical fact may be deemed to be forward-looking statements. Words such as "believes," "anticipates," "plans," "expects," "will," "may," "intends," "prepares," "looks," "potential," "possible" and similar expressions are intended to identify forward-looking statements. These forward-looking statements include statements relating to our future operations, financial performance, projections and guidance, business plans, market opportunities and potential growth, priorities and research and development programs and technologies; the potential benefits of our technologies and scientific approaches, including our siRNA programs; the timing of our ongoing and planned clinical trials; and our expected plans and milestones, including upcoming data readouts for DM1 and FSHD by the end of 2026.

These forward-looking statements involve risks and uncertainties, many of which are beyond Sarepta's control. Actual results could materially differ from those stated or implied by these forward-looking statements as a result of such risks and uncertainties. Known risk factors include the following: different methodologies, assumptions and applications we use to assess particular safety or efficacy parameters may yield different statistical results, and even if we believe the data collected from clinical trials are positive, the results of future research may not be consistent with past positive results, or may fail to meet regulatory approval requirements for the safety and efficacy of our products; success in preclinical and clinical trials, especially if based on a small patient sample, does not ensure that later clinical trials will be successful; our products or product candidates may be perceived as insufficiently effective, unsafe or may result in unforeseen adverse events; we may observe adverse reactions in our clinical trials or in patients who receive our approved products; our products may not be widely adopted by patients, payors or healthcare providers, which would adversely impact our business; our products or product candidates may cause undesirable side effects that result in significant negative consequences following any marketing approval; certain programs may never advance in the clinic or may be discontinued for a number of reasons, including regulators imposing a clinical hold and us suspending or terminating clinical research or trials; if the actual number of patients suffering from the diseases we aim to treat is smaller than estimated, our revenue and ability to achieve profitability may be adversely affected; we may not be able to execute on our business plans, including meeting our expected or planned regulatory milestones and timelines, research and clinical development plans, and bringing our product candidates to market, for various reasons, some of which may be outside of our control, including possible limitations of company financial and other resources, manufacturing limitations that may not be anticipated or resolved for in a timely manner, and regulatory, court or agency decisions, such as decisions by the United States Patent and Trademark Office with respect to patents that cover our product candidates; and those risks identified under the heading "Risk Factors" in our most recent Quarterly Report on Form 10-Q filed with the Securities and Exchange Commission (SEC) as well as other SEC filings made by the Company which you are encouraged to review.

Internet Posting of Information

We routinely post information that may be important to investors in the 'For Investors' section of our website at www.sarepta.com. We encourage investors and potential investors to consult our website regularly for important information about us.

Sarepta Therapeutics, Inc.
Condensed Consolidated Statements of (Loss) Income
(unaudited, in thousands, except per share amounts)

	For the Three Months Ended September 30,		For the Nine Months Ended September 30,	
	2026	**2025**	**2026**	**2025**
Revenues:				
Products, net	$ 323,763	$ 370,043	$ 982,967	$ 1,494,689
Collaboration and other	57,588	29,313	530,438	260,614
Total revenues	381,351	399,356	1,513,405	1,755,303
Cost and expenses:				
Cost of sales (excluding amortization of in-licensed rights)	188,512	150,775	446,665	440,897
Research and development	82,549	218,890	327,767	1,196,730
Selling, general and administrative	118,612	91,893	335,163	363,419
Litigation settlement charge	—	—	39,000	—
Restructuring charge	—	40,510	—	40,510
Amortization of in-licensed rights	718	677	2,126	1,945
Total cost and expenses	390,391	502,745	1,150,721	2,043,501
Operating (loss) income	(9,040)	(103,389)	362,684	(288,198)
Other (loss) income, net:				
Other (expense) income, net[1]	(16,726)	34,958	(47,025)	(10,113)
Gain on debt extinguishment[1]	—	3,761	—	3,761
Total other (loss) income, net	(16,726)	38,719	(47,025)	(6,352)
(Loss) income before income tax expense (benefit)	(25,766)	(64,670)	315,659	(294,550)
Income tax expense (benefit)	2,695	(14,102)	18,051	6,634
Net (loss) income	$ (28,461)	$ (50,568)	$ 297,608	$ (301,184)
(Loss) earnings per share:				
Basic	$ (0.27)	$ (0.50)	$ 2.82	$ (3.06)
Diluted	$ (0.27)	$ (0.50)	$ 2.78	$ (3.06)
Weighted average number of shares of common stock used in computing (loss) earnings per share:				
Basic	105,930	100,237	105,453	98,545
Diluted	105,930	100,237	107,646	98,545

[1] Other (expense) income, net and gain on debt extinguishment for the three and nine months ended September 30, 2025 have been revised to reflect the correction of an immaterial error associated with the accounting for the August 2025 Exchange, as well as the adoption of ASU 2025-07. Please refer to the Company's Annual Report on Form 10-K for the year ended December 31, 2025 for additional information.

Sarepta Therapeutics, Inc.
Reconciliation of GAAP Financial Measures to Non-GAAP Financial Measures
(unaudited, in thousands, except per share amounts)

	For the Three Months Ended September 30,		For the Nine Months Ended September 30,	
	2026	2025	2026	2025
GAAP net (loss) income[1]	$ (28,461)	$ (50,568)	$ 297,608	$ (301,184)
Interest expense (income), net[1]	16,093	1,078	42,386	(8,768)
Depreciation and amortization expense	5,182	12,221	24,777	31,771
Stock-based compensation expense	26,743	14,927	80,614	93,380
Loss (gain) on strategic investments	523	(36,739)	2,683	17,268
Restructuring charge	—	40,510	—	40,510
Impairment of strategic investment	—	—	1,000	—
Litigation settlement charge	—	—	39,000	—
Gain on debt extinguishment[1]	—	(3,761)	—	(3,761)
Income tax effect of adjustments	(4,452)	9,477	(8,475)	607
Non-GAAP net income (loss)	$ 15,628	$ (12,855)	$ 479,593	$ (130,177)
GAAP (loss) earnings per share - diluted:[1, 2]	$ (0.27)	$ (0.50)	$ 2.78	$ (3.06)
Add: impact of GAAP to Non-GAAP adjustments[1]	0.40	0.37	1.13	1.74
Non-GAAP earnings (loss) per share - diluted[2]	$ 0.13	$ (0.13)	$ 3.91	$ (1.32)
Weighted average number of shares of common stock used in computing diluted earnings (loss) per share:[3]				
GAAP	105,930	100,237	107,646	98,545
Non-GAAP	123,223	100,237	122,609	98,545

[1] GAAP net (loss) income, interest expense (income), net, gain on debt extinguishment, GAAP diluted loss per share and the corresponding impact of these GAAP to non-GAAP adjustments for the three and nine months ended September 30, 2025 have been revised to reflect the correction of an immaterial error associated with the accounting for the August 2025 Exchange, as well as the adoption of ASU 2025-07. Please refer to the Company's Annual Report on Form 10-K for the year ended December 31, 2025 for additional information.
[2] GAAP and non-GAAP earnings per share is calculated using diluted shares whereas GAAP and non-GAAP net loss per share is calculated using basic shares as all other instruments are anti-dilutive.
[3]The difference between the weighted average number of shares of common stock used in computing diluted GAAP and non-GAAP earnings per share for the nine months ended September 30, 2026 is a result of the exclusion of the potential share settlement of the 2030 Convertible Notes from the GAAP earnings per share as the inclusion of such shares was anti-dilutive.

Sarepta Therapeutics, Inc.
Reconciliation of GAAP Financial Measures to Non-GAAP Financial Measures
(unaudited, in thousands, except per share amounts)

	For the Three Months Ended September 30,		For the Nine Months Ended September 30,	
	2026	**2025**	**2026**	**2025**
GAAP research and development expenses	$ 82,549	$ 218,890	$ 327,767	$ 1,196,730
Stock-based compensation expense	(8,509)	(4,139)	(27,295)	(36,733)
Depreciation and amortization expense	(2,412)	(8,227)	(14,641)	(22,601)
Non-GAAP research and development expenses	$ 71,628	$ 206,524	$ 285,831	$ 1,137,396

	For the Three Months Ended September 30,		For the Nine Months Ended September 30,	
	2026	**2025**	**2026**	**2025**
GAAP selling, general and administrative expenses	$ 118,612	$ 91,893	$ 335,163	$ 363,419
Stock-based compensation expense	(18,234)	(10,788)	(53,319)	(56,647)
Depreciation expense	(2,770)	(3,994)	(10,136)	(9,170)
Non-GAAP selling, general and administrative expenses	$ 97,608	$ 77,111	$ 271,708	$ 297,602

	For the Three Months Ended September 30,		For the Nine Months Ended September 30,	
	2026	**2025**	**2026**	**2025**
GAAP operating (loss) income	$ (9,040)	$ (103,389)	$ 362,684	$ (288,198)
Stock-based compensation expense	26,743	14,927	80,614	93,380
Depreciation and amortization expense	5,182	12,221	24,777	31,771
Restructuring charge	—	40,510	—	40,510
Litigation settlement charge	—	—	39,000	—
Non-GAAP operating income (loss)	$ 22,885	$ (35,731)	$ 507,075	$ (122,537)

	For the Three Months Ended September 30,		For the Nine Months Ended September 30,	
	2026	**2025**	**2026**	**2025**
Total effective tax rate, GAAP[1]	(10.5) %	21.8 %	5.7 %	(2.3) %
Less: impact of GAAP to non-GAAP adjustments[1]	41.9	42.9	(0.5)	(2.6)
Total effective tax rate, Non-GAAP	31.4 %	64.7 %	5.2 %	(4.9) %

[1] Total effective tax rate, GAAP and impact of GAAP to non-GAAP adjustments for the three and nine months ended September 30, 2025 have been revised to reflect the correction of an immaterial error associated with the accounting for the August 2025 Exchange, as well as the adoption of ASU 2025-07. Please refer to the Company's Annual Report on Form 10-K for the year ended December 31, 2025 for additional information.

Sarepta Therapeutics, Inc.
Condensed Consolidated Balance Sheets
(unaudited, in thousands, except share and per share amounts)

	As of September 30, 2026	As of December 31, 2025
Assets		
Current assets:		
Cash and cash equivalents	$ 472,760	$ 801,282
Short-term investments	406,016	138,368
Accounts receivable, net	369,178	398,233
Inventory	876,118	914,744
Manufacturing-related deposits and prepaids	62,206	113,455
Other current assets	86,611	171,856
Total current assets	2,272,889	2,537,938
Property and equipment, net	319,288	345,125
Right of use assets	117,709	125,495
Non-current inventory	200,579	184,543
Non-current investments	108,815	1,048
Other non-current assets	146,786	155,554
Total assets	$ 3,166,066	$ 3,349,703
Liabilities and Stockholders' Equity		
Current liabilities:		
Accounts payable	$ 45,704	$ 280,841
Accrued expenses	321,764	359,659
Deferred revenue, current portion	104,887	443,397
Current portion of long-term debt	157,971	—
Other current liabilities	26,027	11,393
Total current liabilities	656,353	1,095,290
Long-term debt	699,393	828,974
Lease liabilities, net of current portion	195,663	199,378
Deferred revenue, net of current portion	83,910	83,910
Other non-current liabilities	1,894	1,529
Total liabilities	1,637,213	2,209,081
Stockholders' equity:		
Preferred stock, $0.0001 par value, 3,333,333 shares authorized; none issued and outstanding	—	—
Common stock, $0.0001 par value, 198,000,000 shares authorized; 107,888,367 and 107,232,725 issued and outstanding, respectively, at September 30, 2026 and 105,615,096 and 104,964,220 issued and outstanding, respectively, at December 31, 2025	11	11
Treasury stock, at cost, 655,642 and 650,876 shares at September 30, 2026 and December 31, 2025, respectively	(25,013)	(25,263)
Additional paid-in capital	6,134,817	6,042,586
Accumulated other comprehensive (loss) income, net of tax	(1,586)	272
Accumulated deficit	(4,579,376)	(4,876,984)
Total stockholders' equity	1,528,853	1,140,622
Total liabilities and stockholders' equity	$ 3,166,066	$ 3,349,703

Source: Sarepta Therapeutics, Inc.

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