

Sarepta Therapeutics

Encouraging dystrophin expression from MOMENTUM Part B; Maintain Buy

Maintain Rating: BUY | PO: 164.00 USD | Price: 121.02 USD

'5051 could provide layer of protection to current revs

Sarepta (SRPT) announced positive Part B data from the phase 2 MOMENTUM trial evaluating SRP-5051 (next-gen exon skipping) in Duchenne muscular dystrophy (DMD) patients (DMD) amenable to exon 51 skipping. Data from 20 patients dosed at ~30mg/kg Q4W showed mean dystrophin expression of 5.17% representing a 4.53% change from baseline (p<0.0001) at week 28. These results suggest ~12.2-fold higher dystrophin expression compared to eteplirsen at week 24. Management commented they expect to have a pre-NDA meeting with the FDA in 3Q to discuss the data and potential path forward. On the potential for SRP-5051 in the treatment paradigm, the company commented that this will likely depend on the breadth of the Elevidys potential label expansion and feedback from regulators, but highlighted the favorable profile so far compared to standard of care. We are encouraged by the update suggesting significant improvement in dystrophin expression over eteplirsen but await more color on the regulatory path forward and the evolving competitive dynamics of the DMD treatment landscape. We already provide significant value to '5051 in our \$1.7bn pipeline value and await further details before assigning standalone value. We think the positive data could help position '5051 in the treatment paradigm and could serve as an additional revenue stream in case of a restricted Elevidys label. In our view, Elevidys continues to be the key value driver and we remain focused on the outcome of the label expansion application, which could provide additional upside to our estimates. We reiterate SRPT as one of our 2024 top picks. We maintain our Buy rating with \$164 PO.

Dystrophin expression independent of ambulatory status

Dystrophin expression correlated with mean exon skipping of 11.11% at \sim 30mg/kg. Additionally, dystrophin expression was consistent across ambulatory (4.76%, N=11) and non-ambulatory patients (5.67%, N=9), which suggests potential benefit to both DMD populations. Part B also evaluated the \sim 20mg/kg Q4W dose. At 20mg/kg, mean dystrophin expression change from baseline was 1.60% and mean exon skipping was 2.47%.

Hypomagnesemia can be managed prophylactically

On safety, the most common treatment-related adverse events were hypomagnesemia (95.2%) and hypokalemia (41.9%) in-line with previous reports. Management commented hypomagnesemia was well-managed with prophylactic magnesium supplementation and none of these events resulted in discontinuation. Glomerular filtration rate decreases were seen in 6.5% of patients but did not correlate with other biomarkers.

Estimates (Dec) (US\$)	2021A	2022A	2023E	2024E	2025E
EPS	(5.15)	(8.03)	(7.58)	0.67	13.37
EPS Change (YoY)	27.6%	-55.9%	5.6%	NM	NM
Consensus EPS (Bloomberg)			(1.22)	5.79	10.34
DPS	0	0	0	0	0
Valuation (Dec)					
P/E	NM	NM	NM	180.6x	9.1x
EV / EBITDA*	NM	NM	NM	86.5x	9.5x
Free Cash Flow Yield*	-4.3%	-3.1%	-3.7%	2.9%	13.6%
* For full definitions of <i>IQ</i> method SM measures, see page 5.					

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Refer to important disclosures on page 6 to 8. Analyst Certification on page 4. Price
Objective Basis/Risk on page 4.

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Stock Data

 Price
 121.02 USD

 Price Objective
 164.00 USD

 Date Established
 30-Oct-2023

 Investment Opinion
 C-1-9

 52-Week Range
 55.25 USD - 159.89 USD

 Mrkt Val (mn) / Shares Out
 11,321 USD / 93.5

(mn)

Free Float 95.6%
Average Daily Value (mn) 175.35 USD
BofA Ticker / Exchange SRPT / NAS
Bloomberg / Reuters SRPT US / SRPT.OQ
ROE (2023E) -350.9%
Net Dbt to Eqty (Dec-2022A) 150.0%

ESGMeter™ Medium

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Abbreviations

FDA: Food and Drug Administration

NDA: New drug application

Q4W: Every 4 weeks

iQprofile[™] Sarepta Therapeutics

iQ method [™] – Bus Performance*					
(US\$ Millions)	2021A	2022A	2023E	2024E	2025
Return on Capital Employed	-15.7%	-18.5%	-13.1%	3.8%	35.0%
Return on Equity	-49.6%	-107.2%	-350.9%	32.8%	106.8%
Operating Margin	-65.5%	-57.5%	-27.2%	5.4%	42.6%
Free Cash Flow	(482)	(356)	(421)	334	1,536
Q method SM – Quality of Earnings*					
(US\$ Millions)	2021A	2022A	2023E	2024E	2025
Cash Realization Ratio	NM	NM	NM	6.1x	1.3>
Asset Replacement Ratio	1.0x	0.7x	0.7x	0.7x	0.7
Tax Rate	0%	NM	NM	0.8%	0%
Net Debt-to-Equity Ratio	-109.8%	150.0%	NM	137.9%	-44.5%
Interest Cover	-6.7x	-15.3x	-9.6x	2.7x	36.4>
income Statement Data (Dec)					
US\$ Millions)	2021A	2022A	2023E	2024E	2025
Sales	702	933	1,234	1,792	2,999
% Change	30.0%	32.9%	32.3%	45.2%	67.3%
Gross Profit	605	793	1,078	1,611	2,723
% Change	26.9%	31.1%	35.9%	49.5%	69.0%
EBITDA	(422)	(494)	(290)	147	1,332
% Change	21.8%	-17.2%	41.3%	NM	806.8%
Net Interest & Other Income	41	(154)	(320)	(35)	(35)
Net Income (Adjusted)	(419)	(703)	(673)	61	1,241
% Change	24.4%	-68.0%	4.4%	NM	NM
US\$ Millions)	2021A	2022A	2023E	2024E	2025
Net Income from Cont Operations (GAAP)	(419)	(703)	(673)	61	1,241
Depreciation & Amortization	38	42	46	51	56
Change in Working Capital	(98)	(39)	(16)	(22)	(30)
Deferred Taxation Charge	NA	NA 276	NA	NA	NA 210
Other Adjustments, Net	36	376	256	282	310
Capital Expenditure Free Cash Flow	(38)	(31)	(34)	(37)	(41)
% Change	-482 NM	-356 26.1%	-421 -18.1%	334 NM	1,536 360.2%
Share / Issue Repurchase	549	20.170	-10.170	0	300.2%
Cost of Dividends Paid	NA	NA	NA	NA	N/
Change in Debt	0	202	0	0	(
	Ü	202	0	U	C
Balance Sheet Data (Dec)	20214	20224	20225	20245	2025
US\$ Millions)	2021A	2022A	2023E	2024E	2025
Cash & Equivalents	2,116	967	738	1,032	2,413
Trade Receivables Other Current Assets	153 335	215 1,376	258	309	371 1,586
			1,250	1,358	,
Property, Plant & Equipment Other Non-Current Assets	191 353	180 390	168 390	155 390	140 390
Total Assets	3,148	3,128	2,804	3,245	4,900
Short-Term Debt	3,140	3,126	2,004	0	4,500
Other Current Liabilities	453	620	682	750	825
Long-Term Debt	1,097	1,544	1,544	1,544	1,544
Other Non-Current Liabilities	670	580	580	580	580
Total Liabilities	2,220	2,743	2,805	2,874	2,949
i otal Liabilities	2,220	2,/ 43	•		•
Total Fouity	928	385	(1)	371	1 957
Total Equity Total Equity & Liabilities	928 3,148	385 3,128	(1) 2,804	371 3,245	1,952 4,900

Company Sector

Biotechnology

Company Description

Sarepta (SRPT) is a biopharmaceuticals company developing exon-skipping therapeutics and gene therapy for the treatment of Duchenne muscular dystrophy (DMD) and Limb Girdle muscular dystrophy (LGMD). The company's approved assets include Exondys, approved for DMD amenable to exon 51 skipping, Vyondys, approved for DMD amenable to exon 53 skipping, and Amondys, approved for DMD amenable to exon 45 skipping. SRPT also has a next-gen PPMO platform and gene therapy programs in DMD and LGMD.

Investment Rationale

We rate SRPT shares Buy. We believe SRPT's assets address the current under-met need in the treatment of rare muscular dystrophies. The exon skipping assets, in our view, should continue to provide floor valuation and generate sustained revenue streams. We believe the company is well positioned to drive upside potential and deliver shareholder value over time given existing assets and strategic partnerships in gene therapy programs with Nationwide, Lacerta, and Lysogene.

Stock Data

Average Daily Volume

1,448,970

Quarterly Earnings Estimates

	2022	2023
Q1	-1.20A	-5.85A
Q2	-2.65A	-0.27A
Q3	-2.94A	-0.46A
04	-1.24A	-1.03E



Price objective basis & risk

Sarepta Therapeutics (SRPT)

Our \$164 PO is based on a probability-adjusted net present value (NPV) analysis that includes \$17/share for Exondys, \$8/share for Vyondys, \$22/share for Amondys, \$78/share for micro-dystrophin gene therapy, \$38/share for LGMD assets. The remainder of our valuation comes from pipeline and cash. Our discounted cash flow (DCF) analysis assumes sales out to 2035, with weighted average cost of capital (WACC) of 9-13%.

Upside risks to our PO are 1) better-than-expected market uptake from its exon skipping assets, 2) SRP-9001 US label expansion, and 3) EU approval of SRP-9001 and other assets.

Downside risks to our PO are 1) failure and delay in approval for exon skipping assets, 2) unexpected safety or durability findings in gene therapy programs, 3) other micro dystrophin competitors, and 4) higher-than-expected royalty payments.

Analyst Certification

I, Tazeen Ahmad, hereby certify that the views expressed in this research report accurately reflect my personal views about the subject securities and issuers. I also certify that no part of my compensation was, is, or will be, directly or indirectly, related to the specific recommendations or view expressed in this research report.

US - Biotechnology Coverage Cluster

Investment rating	Company	BofA Ticker	Bloomberg symbol	Analyst
BUY				
	4D Molecular Therapeutics, Inc.	FDMT	FDMT US	Tazeen Ahmad
	Alnylam Pharmaceuticals	ALNY	ALNY US	Tazeen Ahmad
	Amicus Therapeutics	FOLD	FOLD US	Tazeen Ahmad
	Annexon Biosciences	ANNX	ANNX US	Tazeen Ahmad
	Apellis Pharmaceuticals	APLS	APLS US	Tazeen Ahmad
	Argenx SE	ARGX	ARGX US	Tazeen Ahmad
	Arvinas	ARVN	ARVN US	Tazeen Ahmad
	Ascendis Pharma	ASND	ASND US	Tazeen Ahmad
	Biocryst Pharmaceuticals Inc	BCRX	BCRX US	Tazeen Ahmad
	BioNTech	BNTX	BNTX US	Tazeen Ahmad
	Denali Therapeutics	DNLI	DNLI US	Tazeen Ahmad
	Inozyme Pharma, Inc.	INZY	INZY US	Tazeen Ahmad
	Merus	MRUS	MRUS US	Tazeen Ahmad
	Neurocrine Biosciences	NBIX	NBIX US	Tazeen Ahmad
	PepGen Inc	PEPG	PEPG US	Tazeen Ahmad
	Prothena Corporation	PRTA	PRTA US	Tazeen Ahmad
	Rhythm Pharmaceuticals	RYTM	RYTM US	Tazeen Ahmad
	Sarepta Therapeutics	SRPT	SRPT US	Tazeen Ahmad
	Ultragenyx Pharmaceuticals	RARE	RARE US	Tazeen Ahmad
NEUTRAL				
	Acadia Pharmaceuticals	ACAD	ACAD US	Tazeen Ahmad
	Incyte Corporation	INCY	INCY US	Tazeen Ahmad
	SAGE Therapeutics	SAGE	SAGE US	Tazeen Ahmad
UNDERPERFORM				
	Achilles Therapeutics	ACHL	ACHL US	Tazeen Ahmad
	Fate Therapeutics	FATE	FATE US	Tazeen Ahmad
	Fulcrum Therapeutics	FULC	FULC US	Tazeen Ahmad
	Pharvaris	PHVS	PHVS US	Tazeen Ahmad
	PTC Therapeutics	PTCT	PTCT US	Tazeen Ahmad

*IQ*method[™] Measures Definitions

Business Performance	Numerator	Denominator
Return On Capital Employed	NOPAT = (EBIT + Interest Income) × (1 – Tax Rate) + Goodwill Amortization	Total Assets – Current Liabilities + ST Debt + Accumulated Goodwill Amortization
Return On Equity	Net Income	Shareholders' Equity
Operating Margin	Operating Profit	Sales
Earnings Growth	Expected 5 Year CAGR From Latest Actual	N/A
Free Cash Flow	Cash Flow From Operations — Total Capex	N/A
Quality of Earnings	Numerator	Denominator
Cash Realization Ratio	Cash Flow From Operations	Net Income
Asset Replacement Ratio	Capex	Depreciation
Tax Rate	Tax Charge	Pre-Tax Income
Net Debt-To-Equity Ratio	Net Debt = Total Debt - Cash & Equivalents	Total Equity
Interest Cover	EBIT	Interest Expense
Valuation Toolkit	Numerator	Denominator
Price / Earnings Ratio	Current Share Price	Diluted Earnings Per Share (Basis As Specified)
Price / Book Value	Current Share Price	Shareholders' Equity / Current Basic Shares
Dividend Yield	Annualised Declared Cash Dividend	Current Share Price
Free Cash Flow Yield	Cash Flow From Operations – Total Capex	Market Cap = Current Share Price × Current Basic Shares
Enterprise Value / Sales	EV = Current Share Price × Current Shares + Minority Equity + Net Debt + Other LT Liabilities	Sales

EV / EBITDA Enterprise Value Basic EBIT + Depreciation + Amortization

Menethod 3*is the set of BofA Global Research standard measures that serve to maintain global consistency under three broad headings: Business Performance, Quality of Earnings, and validations. The key features of iQmethod are: A consistently structured, detailed, and transparent methodology. Guidelines to maximize the effectiveness of the comparative valuation process, and to identify some common pitfalls.

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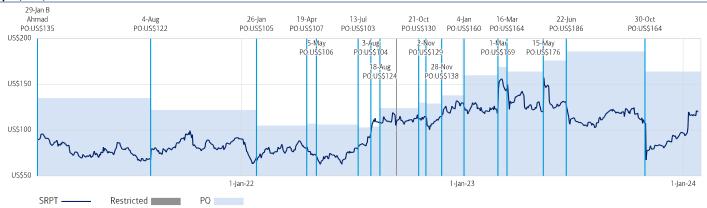
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Sarepta (SRPT) Price Chart



B: Buy, N: Neutral, U: Underperform, PO: Price Objective, NA: No longer valid, NR: No Rating

The Investment Opinion System is contained at the end of the report under the heading "Fundamental Equity Opinion Key". Dark grey shading indicates the security is restricted with the opinion suspended. Medium grey shading indicates the security is under review with the opinion withdrawn. Light grey shading indicates the security is not covered. Chart is current as of a date no more than one trading day prior to the date of the report.

Equity Investment Rating Distribution: Health Care Group (as of 31 Dec 2023)

Coverage Universe	Count	Percent	Inv. Banking Relationships R1	Count	Percent
Buy	234	60.94%	Buy	115	49.15%
Hold	80	20.83%	Hold	36	45.00%
Sell	70	18.23%	Sell	29	41.43%

Equity Investment Rating Distribution: Global Group (as of 31 Dec 2023)

Coverage Universe	Count	Percent	Inv. Banking Relationships R1	Count	Percent
Buy	1895	53.62%	Buy	1083	57.15%
Hold	832	23.54%	Hold	454	54.57%
Sell	807	22.84%	Sell	383	47.46%

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Investment rating Total return expectation (within 12-month period of date of initial rating) Ratings dispersion guidelines for coverage cluster^{R2}

Buy	≥ 10%	≤ /0%
Neutral	≥ 0%	≤ 30%
Jnderperform	N/A	≥ 20%

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