

Prothena Corporation

Further clarity needed on competitive positioning in Alzheimer's; D/G to Neutral

Rating Change: NEUTRAL | PO: 38.00 USD | Price: 33.76 USD

Next AD program update could come in 2025

We downgrade PRTA to Neutral from Buy with new \$38 PO (prev. \$68) as we think clarity on the potential clinical profile for PRX012 in AD is needed. Recall in January PRTA provided an initial update on the ongoing ph 1 trial evaluating '012 in AD highlighting initial encouraging amyloid reduction and ARIA-E rate at the 70mg dose but did not provide additional details on the magnitude of effect. When we caught up with mgmt to discuss, CEO Gene Kinney indicated that AD is a competitive space and that upon discussion with the PRTA board the company decided to wait until collection of more data to provide a fulsome update to the Street. The company noted the 45mg and 200mg cohorts are ongoing with the option to add additional dose cohorts up to 400mg, if needed. Mgmt did not provide specific guidance on the timing of the next update and commented it could move into 2025. They highlighted that results so far support monthly dosing and they remain confident on the potential for a competitive profile. While we think this update is a step in the right direction, we think it is prudent to assume a more conservative stance on '012 given the potential for a longer timeline to answer our and the Street's questions on how the drug compares to competition. In our DCF-based model, we roll the quarter and reduce our peak penetration for '012 in AD 5% (prev. 10%). We also push our estimated launch for birtamimab in AL to 2026 based on new timelines (see below) and assume a more conservative early ramp for the launch. These changes result in our new \$38 PO. '012 now contributes \$9/sh to our PO. While we still believe PRTA's science could allow for a differentiated product, we await clinical validation in what is an increasingly competitive AD space.

Ph 3 AFFIRM-AL topline now not expected before 4Q24

PRTA also noted they now expect the readout of the ph 3 AFFIRM-AL trial evaluating bir' in Mayo Stage IV AL amyloidosis between 4Q24-2Q25 (prev. 2024) based on their latest estimates. They highlighted this is an event-driven trial and the timing is still within expectations but noted a slower event rate compared to previous trials. The company noted an interim analysis for overwhelming efficacy is planned when ~50% of events have occurred and highlighted there is no futility analysis. Mgmt noted they will wait until the full readout to provide the next update. We note that a potential later readout is not uncommon for event-driven trials, but the delay could mean a lack of meaningful catalysts in the next 12 mos for the company. That said, when the AL amyloidosis study reads out, if positive, would provide meaningful upside to our current estimates given the potential to generate sales soon. Bir' now contributes \$15/sh to our PO.

Estimates (Dec) (US\$)	2021A	2022A	2023E	2024E	2025E
EPS	1.38	(2.47)	(2.35)	(4.51)	(5.62)
GAAP EPS	1.51	(2.47)	(2.35)	(4.51)	(5.62)
EPS Change (YoY)	NM	NM	4.9%	-91.9%	-24.6%
Consensus EPS (Bloomberg)			(2.79)	(4.32)	(4.39)
DPS	0	0	0	0	0
Valuation (Dec)					
P/E	24.5x	NM	NM	NM	NM
GAAP P/E	22.4x	NM	NM	NM	NM
EV / EBITDA*	18.0x	NM	NM	NM	NM
Free Cash Flow Yield*	5.1%	-6.0%	-5.5%	-12.8%	-17.8%

* For full definitions of *IQmethod*SM measures, see page 5.

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Timestamp: 30 January 2024 06:00AM EST

30 January 2024

Equity

Key Changes

(US\$)	Previous	Current
Inv. Opinion	C-1-9	C-2-9
Inv. Rating	BUY	NEUTRAL
Price Obj.	68.00	38.00
2025E Rev (m)	46.3	10.0
2025E EPS	-4.32	-5.62

Tazeen Ahmad
Research Analyst
BofAS
+1 646 855 4236
tazeen.ahmad@bofa.com

Daniel Giraldo
Research Analyst
BofAS
+1 646 855 0993
daniel.giraldoperez@bofa.com

Jeremiah Lorentz
Research Analyst
BofAS
+1 616 743 2514
jeremiah.lorentz@bofa.com

Stock Data

Price	33.76 USD
Price Objective	38.00 USD
Date Established	30-Jan-2024
Investment Opinion	C-2-9
52-Week Range	28.51 USD - 79.65 USD
Mkt Val (mn) / Shares Out (mn)	1,812 USD / 53.7
Free Float	73.8%
Average Daily Value (mn)	34.41 USD
BofA Ticker / Exchange	PRTA / NAS
Bloomberg / Reuters	PRTA US / PRTA.OQ
ROE (2023E)	-23.4%
Net Dbt to Eqty (Dec-2022A)	-114.2%
ESGMeter TM	Medium

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See inside for abbreviations

iQprofileSM Prothena Corporation

iQmethodSM – Bus Performance*

(US\$ Millions)	2021A	2022A	2023E	2024E	2025E
Return on Capital Employed	14.7%	-17.6%	-19.9%	-39.4%	-50.3%
Return on Equity	20.6%	-21.5%	-23.4%	-49.7%	-65.3%
Operating Margin	35.9%	-244.1%	-176.9%	-3,097.7%	-3,868.5%
Free Cash Flow	92	(109)	(100)	(232)	(323)

iQmethodSM – Quality of Earnings*

(US\$ Millions)	2021A	2022A	2023E	2024E	2025E
Cash Realization Ratio	1.4x	NM	NM	NM	NM
Asset Replacement Ratio	0.5x	0.6x	1.6x	2.4x	2.9x
Tax Rate	6.9%	6.9%	8.9%	4.7%	3.7%
Net Debt-to-Equity Ratio	-124.3%	-114.2%	-124.2%	-122.8%	-128.4%
Interest Cover	NM	NA	NA	NA	NA

Income Statement Data (Dec)

(US\$ Millions)	2021A	2022A	2023E	2024E	2025E
Sales	201	54	101	10	10
% Change	23,414.3%	-73.1%	87.5%	-90.1%	0%
Gross Profit	201	54	101	10	10
% Change	NM	-73.1%	87.5%	-90.1%	0%
EBITDA	72	(132)	(179)	(310)	0
% Change	NM	NM	-35.9%	-73.3%	100.0%
Net Interest & Other Income	0	6	36	37	39
Net Income (Adjusted)	67	(117)	(130)	(259)	(335)
% Change	NM	NM	-11.4%	-99.2%	-29.0%

Free Cash Flow Data (Dec)

(US\$ Millions)	2021A	2022A	2023E	2024E	2025E
Net Income from Cont Operations (GAAP)	67	(117)	(130)	(259)	(335)
Depreciation & Amortization	1	1	1	1	1
Change in Working Capital	(10)	(19)	(11)	(21)	(36)
Deferred Taxation Charge	4,573	(11,133)	0	0	0
Other Adjustments, Net	(4,538)	11,159	42	50	50
Capital Expenditure	(1)	0	(1)	(2)	(3)
Free Cash Flow	92	-109	-100	-232	-323
% Change	NM	NM	8.8%	-132.5%	-39.3%
Share / Issue Repurchase	175	224	0	300	250
Cost of Dividends Paid	0	0	0	0	0
Change in Debt	0	0	0	0	0

Balance Sheet Data (Dec)

(US\$ Millions)	2021A	2022A	2023E	2024E	2025E
Cash & Equivalents	579	710	611	679	606
Trade Receivables	0	0	0	0	0
Other Current Assets	6	9	10	12	13
Property, Plant & Equipment	2	2	2	2	2
Other Non-Current Assets	23	37	13	13	13
Total Assets	609	758	636	705	634
Short-Term Debt	0	0	0	0	0
Other Current Liabilities	33	50	49	49	48
Long-Term Debt	0	0	0	0	0
Other Non-Current Liabilities	110	86	94	104	114
Total Liabilities	143	136	144	152	162
Total Equity	466	622	492	553	472
Total Equity & Liabilities	609	758	636	705	634

* For full definitions of iQmethodSM measures, see page 5.

Company Sector

Biotechnology

Company Description

Prothena (PRTA) is a development-stage biotechnology company that was spun-off from Irish drug maker Elan in 2012. The company is based in Ireland, with US offices in South San Francisco, California. PRTA's pipeline includes both wholly-owned and partnered programs being developed for the potential treatment of diseases including AL amyloidosis, ATTR amyloidosis, Alzheimer's disease, Parkinson's disease and a number of other neurodegenerative diseases.

Investment Rationale

We rate PRTA shares Neutral. The company has PRX012 in development for Alzheimer's disease (AD) which benefit from read-through of recently approved Aduhelm (similar mechanism) data but we look for clarity on timelines for clinical validation in this increasingly competitive space. The company also has several programs in development including birtamimab for AL Amyloidosis and prasinezumab for PD.

Stock Data

Average Daily Volume 1,019,245

Quarterly Earnings Estimates

	2022	2023
Q1	-0.78A	-0.89A
Q2	-0.88A	-1.03A
Q3	-0.97A	0.38E
Q4	0.12A	-0.87E

Partner-led program readouts expected in 2024

The company also reported results from the phase 2b PADOVA study (led by Roche) evaluating prasinezumab in Parkinson's disease are expected in 2024. Management commented the next payments related to the program are related to regulatory milestones. Prasinezumab contributes \$7/sh to our PO. Topline phase 2 data for NNC6019 (PRX004, led by novo Nordisk) in transthyretin amyloidosis are also expected in 2024. Management commented they expect a potential clinical development milestone payment. Additionally, the company reported that BMS-986446 (PRX005) in Alzheimer's disease (led by Bristol Myers Squibb) is expected to move into phase 2 in 1H following phase 1 results.

Abbreviations

AD: Alzheimer's disease

AL: Light chain

ARIA-E: Amyloid-related Imaging Abnormalities -Effusion and Edema

mgmt: management

mos: months

ph: phase

PRTA: Prothena

Price objective basis & risk

Prothena Corporation (PRTA)

We use a sum of the parts DCF valuation approach to arrive at our NPV based PO of \$38/share. We assume a WACC of 11% and model a value of \$15/sh for birtamimab in AL amyloidosis, a WACC of 12% and model a value of \$9/sh for PRX012 in AD, and a WACC of 12% and model a value of \$7/sh for prasinezumab in Parkinson's. The balance of our value for PRTA comes from cash (\$10/sh), and pipeline value (-\$3). Our DCF goes out to 2040 and we assume zero terminal value (consistent with the way we model our other companies).

Upside risks to our PO are 1) earlier-than-expected approval and launch of prasinezumab or birtamimab, 2) success of pipeline clinical trials, and 3) faster-than-expected revenue ramp. Downside risks to our PO are 1) failure of one or more of PRTA's products to reach market, 2) higher-than-expected competition, and 3) negative news on competitor (BIIB) AB program in AD.

Analyst Certification

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US - Biotechnology Coverage Cluster

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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
	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
	Prothena Corporation	PRTA	PRTA 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

iQmethodSM Measures Definitions

Business Performance

Return On Capital Employed

Return On Equity

Operating Margin

Earnings Growth

Free Cash Flow

Quality of Earnings

Cash Realization Ratio

Asset Replacement Ratio

Tax Rate

Net Debt-To-Equity Ratio

Interest Cover

Valuation Toolkit

Price / Earnings Ratio

Price / Book Value

Dividend Yield

Free Cash Flow Yield

Enterprise Value / Sales

EV / EBITDA

Numerator

$\text{NOPAT} = (\text{EBIT} + \text{Interest Income}) \times (1 - \text{Tax Rate}) + \text{Goodwill Amortization}$

Net Income

Operating Profit

Expected 5 Year CAGR From Latest Actual

Cash Flow From Operations – Total Capex

Numerator

Cash Flow From Operations

Capex

Tax Charge

Net Debt = Total Debt – Cash & Equivalents

EBIT

Numerator

Current Share Price

Current Share Price

Annualised Declared Cash Dividend

Cash Flow From Operations – Total Capex

$\text{EV} = \text{Current Share Price} \times \text{Current Shares} + \text{Minority Equity} + \text{Net Debt} +$

Other LT Liabilities

Enterprise Value

Denominator

$\text{Total Assets} - \text{Current Liabilities} + \text{ST Debt} + \text{Accumulated Goodwill}$

Amortization

Shareholders' Equity

Sales

N/A

N/A

Denominator

Net Income

Depreciation

Pre-Tax Income

Total Equity

Interest Expense

Denominator

Diluted Earnings Per Share (Basis As Specified)

Shareholders' Equity / Current Basic Shares

Current Share Price

$\text{Market Cap} = \text{Current Share Price} \times \text{Current Basic Shares}$

Sales

Basic EBIT + Depreciation + Amortization

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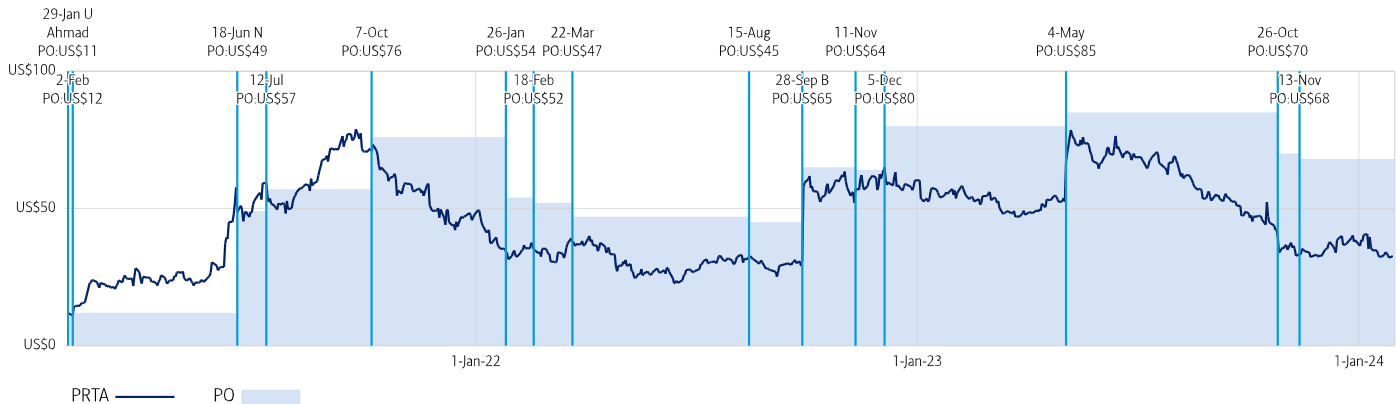
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Important Disclosures

Prothena (PRTA) 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%

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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%	≤ 70%
Neutral	≥ 0%	≤ 30%
Underperform	N/A	≥ 20%

^{R2} Ratings dispersions may vary from time to time where BofA Global Research believes it better reflects the investment prospects of stocks in a Coverage Cluster.

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