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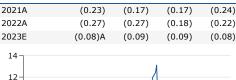
US Equity Research 16 May 2023

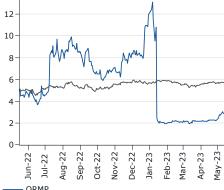
10 1107 2020	
Rating HOLD unchanged	Price Target US\$2.00↓ from US\$3.00
ORMP-NASDAQ	Price US\$3.67

Market Data

52-Week Range (US\$) :	1.81 - 13.73
Market Cap (US\$M) :	147.0
Shares Out., Basic (M) :	40.0
Enterprise Value (US\$M) :	(6.8)
Cash (US\$M):	153.7

FYE Dec	2021A	20	22A	2023E
Revenue (US\$M)	2.7	7	2.7	2.7
EPS (US\$)	(0.78)) (0.94)	(0.35)↑
Previous		-	-	(0.59)
Quarterly	Q1	Q2	Q3	Q4
Revenue				
2021A	0.7	0.7	0.7	0.7
2022A	0.7	0.7	0.7	0.7
2023E	0.7A	0.7	0.7	0.7
Quarterly EPS	Q1	Q2	Q3	Q4
20214	(0.23)	(0.17)	(0.17)	(0.24)





------ NASDAQ Biotechnology (NBI) (rebased) Source: FactSet

Priced intraday 16 May 2023

Lowering Target Price

Oramed Pharmaceuticals

Biotechnology

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Retrospective analysis looking for responding subpopulation; decreasing PT from \$3 to \$2

Oramed reported financial results for 1Q23. We updated our model to reflect the latest financial results and reduced our projected FY2023 G&A expense from \$12M to \$5.5M based on the 1Q23 result. We also anticipate several equity raises between now and 2036. Our 12-month price target decreased from \$3 to \$2 due to the increased share count from the raises. Currently, only cash and cash equivalents of the company are used to generate our price target.

Oramed Pharmaceuticals 1Q23snapshot

	1Q23A	CG 1Q23E	Var (A:E)	% Var	1Q23 Consensus	Var (A:C)	% Var	
GAAP EPS (diluted)	-0.08	-0.29	0.21	262.5%	-0.26	0.18	225.0%	
Product Sales Revenue (\$MM)	0.0	0.0	0.0		0.0	0.0		
R&D	4.4	7.0	-2.6	-59.1%	7.5	-3.1	-70.5%	
G&A	1.3	5.0	-3.7	-295.9%	3.9	-2.6	-208.8%	
Balance sheet	1Q23E							
Cash (\$MM)	\$153.7							
LTD (\$MM)	\$0.0							
Expected upcoming clinical/com	mercial mile	stones						
ORMD-0901	1H23E	Data read-o	ut on bioava	ilability stud	dy (T2D)			
Changes made to financial project	tions							
Updated model to reflct company	es latest fin	ancial result	s					
Reduced projected G&A expense for FY2023 from \$12M to \$5.5M based on 1Q23 results								
Projected several equity raises between 2023 and the out-year of our model 2036								

Source: Company reports and Canaccord Genuity estimates

Oral insulin submitted for marketing in China

Oramed announced that its Chinese partner, Hefei Tianhui (HTIT) has submitted a Marketing Authorization Application for the oral insulin product in China. The submission is supported by a positive Phase III trial in T2D conducted by HTIT in China. The study showed a statistically significant reduction of HbA1c. Data from the trial have not been released.

Post hoc analysis of failed U.S. Phase III searching for responding subgroups

Oramed also announced that a *post hoc* analysis of the U.S. Phase III trial for ORMD-0801 showed an over 1% placebo-adjusted, statistically significant reduction in HbA1c in subpopulations of T2D patients with pooled specific parameters, such as body mass index (BMI), baseline HbA1c, age, gender, and body weight.

A causal machine learning (ML) analysis was used to identify subgroup responders from the U.S. Phase III trial, and results showed that patient baseline BMI and age could be the two most impactful factors on patients' response to ORMD-0801 treatment. The company stated that patients over 54 years of age with BMI<31 tend to have the most reduction in A1c (p<0.002). Another analysis showed that age and BMI, as well as baseline levels of glucose, A1c, and C-peptide, may impact patients' response to the treatment of ORMD-0801.

In January, the company reported that the first of its two Phase III T2D trials missed its primary endpoint of reducing A1c at Week 26.The trial also missed the key secondary endpoint of change in fasting plasma glucose level. The trial, ORA-D-013-1, is the larger of the two trials for ORMD-0801 in T2D patients with inadequate glycemic control on two or three oral glucose-lowering medicines. Upon the news, management also announced the termination of the second T2D Phase III trial, ORA-D-013-2, for ORMD-0801 in T2D patients with only diet control or on metformin monotherapy.

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Oramed Pharmaceuticals (ORMP) income statement

	2021	FY tran	sition	2022			2023E															
	Aug-21	Nov-21 3 months	Dec-21 4 months	Year	1Q	2QE	3QE	4QE	Year	2024E	2025E	2026E	2027E	2028E	2029E	2030E	2031E	2032E	2033E	2034E	2035E	203
ORMD-0801 T2D product sales	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0
Collaboration revenue	2.7	0.7	0.9	2.7	0.7	0.7	0.7	0.7	2.7	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0
Total Revenue	2.7	0.7	0.9	2.7	0.7	0.7	0.7	0.7	2.7	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0
Cost of U.S. product sales (10%)	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0
Gross profit	2.7	0.7	0.9	2.7	0.7	0.7	0.7	0.7	2.7	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0
R&D	21.0	6.4	9.0	27.6	4.4	3.0	3.0	2.0	12.4	6.0	6.0	6.0	6.0	6.0	6.0	6.0	6.0	6.0	6.0	6.0	6.0	6.
G&A	5.9	1.7	3.3	13.8	1.3	1.5	1.5	1.2	5.5	6.0	7.0	8.0	8.0	8.0	8.0	8.0	8.0	8.0	8.0	8.0	8.0	8.
Sales expense	0.0	0.6	0.9	1.9	0.2	0.0	0.0	0.1	0.3	18.8	31.3	37.5	38.6	39.8	41.0	42.2	43.5	44.8	46.1	47.5	48.9	50
Operating expense	26.9	8.7	13.2	43.3	5.9	4.5	4.5	3.4	18.2	30.8	44.3	51.5	52.6	53.8	55.0	56.2	57.5	58.8	60.1	61.5	62.9	64
Operating Profit (Loss)	(24.2)	(8.1)	(12.3)	(40.6)	(5.2)	(3.8)	(3.8)	(2.7)	(15.5)	(30.8)	(44.3)	(51.5)	(52.6)	(53.8)	(55.0)	(56.2)	(57.5)	(58.8)	(60.1)	(61.5)	(62.9)	(64.
Interest income	1.2	(0.1)	0.2	3.8	1.6	0.2	0.2	(1.2)	0.8	0.8	0.8	0.8	(0.5)	(0.5)	-0.5	-0.6	-0.6	-0.6	-0.6	-0.6	-0.6	-0
Interest expense	(0.0)	(0.2)	(0.1)	(0.8)	0.0	0.0	0.0		0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.
Loss (gain) fair value of investments	0.0	0.2	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0
Net loss attributable to non-controlling interests	(0.8)	0.0	(0.6)	1.2	0.2	0.0	0.0	(0.2)	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0
Pretax profit (loss)	(22.2)	(7.9)	(11.7)	(36.5)	(3.4)	(3.6)	(3.6)	(4.1)	(14.7)	(30.0)	(43.5)	(50.7)	(53.2)	(54.3)	-55.5	-56.8	-58.0	-59.4	-60.7	-62.1	-63.6	-65
Income Tax	0.0	0.0	0.0	0.1	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	(14.4)	(14.7)	-15.0	-15.3	-15.7	-16.0	-16.4	-16.8	-17.2	-17
Tax rate	0.0%	0.0%	0.0%	-0.3%	0.0%	0.0%	0.0%	0.0%	0.0%	0.0%	0.0%	0.0%	27.0%	27.0%	27.0%	27.0%	27.0%	27.0%	27.0%	27.0%	27.0%	27.0
Net Income (Loss)	(22.2)	(7.9)	(11.7)	(36.6)	(3.4)	(3.6)	(3.6)	(4.1)	(14.7)	(30.0)	(43.5)	(50.7)	(38.8)	(39.7)	(40.5)	(41.4)	(42.4)	(43.3)	(44.3)	(45.3)	(46.4)	(47.
EPS - basic	(\$0.78)	(\$0.22)	(\$0.31)	(\$0.94)	(\$0.08)	(\$0.09)	(\$0.09)	(\$0.08)	(\$0.35)	(\$0.49)	(\$0.66)	(\$0.77)	(\$0.56)	(\$0.54)	(\$0.53)	(\$0.53)	(\$0.54)	(\$0.55)	(\$0.55)	(\$0.53)	(\$0.53)	(\$0.5
EPS - diluted	(\$0.78)	(\$0.22)		(\$0.94)	(\$0.08)	(\$0.09)	(\$0.09)	(\$0.08)	(\$0.35)	(\$0.49)	(\$0.66)	(\$0.77)	(\$0.56)	(\$0.54)	(\$0.53)	(\$0.53)	(\$0.54)	(\$0.55)	(\$0.55)	(\$0.53)	(\$0.53)	(\$0.5
Weighted average basic shares	28.5	36.7	37.1	39.0	40.0	40.0	40.0	48.3	42.0	61.2	66.2	66.2	69.8	73.4	76.2	78.9	78.9	78.9	81.2	85.7	88.0	88
Weighted average diluted shares	28.5	36.7		39.0	40.0	40.0	40.0	48.3	42.0	61.2	66.2	66.2	69.8	73.4	76.2	78.9	78.9	78.9	81.2	85.7	88.0	88

Source: Company Reports, Canaccord Genuity estimates. A more detailed financial model, including balance sheet, income statement, and cash flow projections, if available, may be obtained by contacting your Canaccord Genuity Sales Person or the Authoring Analyst, whose contact information appears on the front page of this report.



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Investment Recommendation

Date and time of first dissemination: May 16, 2023, 13:23 ET Date and time of production: May 16, 2023, 13:23 ET

Target Price / Valuation Methodology:

Oramed Pharmaceuticals - ORMP

We value shares of Oramed by employing a sum-of-the-parts analysis that includes programs where we believe clinical data is available to fairly determine the overall probability of success, as well as net cash on hand. Currently, only cash and cash equivalents of the company are used to generate our \$2 12-month price target.

Risks to achieving Target Price / Valuation:

Oramed Pharmaceuticals - ORMP

Clinical risk: Although ORMD-0801 has demonstrated clinical proof-of-concept in patients with T2D, the molecule is now being dosed in larger Phase III trials, and as more patients are exposed to drug over time there is the chance that issues could appear relating to efficacy, safety or both. While there is a significant amount of literature speaking to insulin use in treating T2D, ORMD-0801 is a unique orally delivered version of insulin and only through additional clinical trials can the molecule be further de-risked.

Regulatory risk: Given that there are currently no oral-insulin options approved for the treatment of T2D, the FDA is in new territory regarding the review of this type of molecule administration. The FDA continues to be unpredictable even with the review pathways, designations and outside panel reviews that can be employed during the review process.

Commercial risk: Given that there are currently no oral insulin therapeutics approved for the treatment of T1D and T2D, grasping the true commercial opportunity post-launch is difficult. The outcome of the two-Phase III trials and final labeling will be key to understanding the true market potential for ORMD-0801.

Competitive risk: The competitive landscape for T2D drug development is crowded. The opportunity to tap into a mature multibillion-dollar market opportunity will result in the space remaining competitive. In addition to the overall T2D clinical landscape being competitive, specific to Oramed there are multiple players attempting to develop an oral delivery option for insulin.

Management risk: For a clinical stage biotech, stability in the C-suite roles is key as it is with any company. Turnover, especially in regulatory agency-facing roles such as CEO and CSO, could negatively impact share performance.

Distribution of Ratings:

Global Stock Ratings (as of 05/16/23)

Rating	Coverag	Coverage Universe					
	#	%	%				
Buy	602	64.66%	22.26%				
Hold	140	15.04%	10.71%				
Sell	16	1.72%	0.00%				
Speculative Buy	159	17.08%	46.54%				
	931*	100.0%					

*Total includes stocks that are Under Review

Canaccord Genuity Ratings System

BUY: The stock is expected to generate risk-adjusted returns of over 10% during the next 12 months.



HOLD: The stock is expected to generate risk-adjusted returns of 0-10% during the next 12 months.

SELL: The stock is expected to generate negative risk-adjusted returns during the next 12 months.

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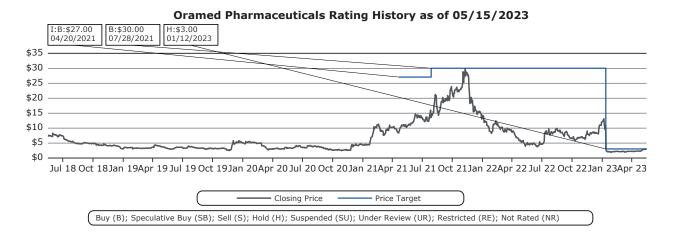
12-Month Recommendation History (as of date same as the Global Stock Ratings table)

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