

	Annual EPS	Annual Revenue	Rating/Target
Today's Changes	2014E \$(0.21) from \$(0.09) 2015E \$(1.89) from \$(1.84)	No changes	No changes

BUY

GlycoMimetics

GLYC: NASDAO: US\$9.13

Target: US\$17.00

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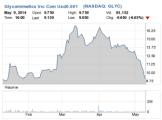
COMPANY STATISTICS:

Forecast Return:	86%
Shares Out (M):	18.6
Market Cap (M):	US\$169.9
52-week Range:	US\$8.40 - 18.99

FARNINGS SUMMARY:

	•••••			
FYE Dec		2013A	2014E	2015E
Revenue:		21.9	23.5	0.0
EPS:		0.61	(0.21)	(1.89)
Revenue:	Q1	1.3A	0.0A	-
	Q2	1.3A	13.5	-
	Q3	1.3A	0.0	-
	Q4	18.0	10.0	-
Total		21.9	23.5	0.0
EPS:	Q1	(0.18)A	(0.30)A	-
	Q2	(0.19)A	0.42	-
	Q3	(0.20)A	(0.42)	-
	Q4	0.76	0.09	-
Total		0.61	(0.21)	(1.89)

SHARE PRICE PERFORMANCE:



Source: Interactive Data Corporation

COMPANY DESCRIPTION:

GlycoMimetics is a clinical-stage biotech company focused on novel glycomimetic (carbohydrate imitating) drugs. The disease-related functions of carbohydrates can include roles in inflammation, cancer cell survival/behavior and infection. GlycoMimetics' GMI-1070 is a very promising therapy for sickle cell disease.

All amounts in US\$ unless otherwise noted.

Life Sciences -- Biotechnology

Q1/14: PFIZER ON TRACK TO START PH3 SICKLE CELL IN H2, AS '1271 PH1 TRIALS GET SET TO START

Investment recommendation

Reiterate BUY, \$17 target on GMI-1070's potential in sickle cell crisis. GLYC's lead candidate, Pfizer-partnered GMI-1070, is a pan-selectin inhibitor for sickle cell vaso-occlusive crisis (VOC). We expect positive data from the upcoming Ph3 trial to show reduced length of sickle cell vasoocclusive crisis based on strong Ph2 data. We think GMI-1070 could gain significant market share and model potential US peak sales of \$1.2B. Our \$17.00 target is based on a pNPV analysis.

Investment highlights

- Q1/14 EPS \$(0.30) vs consensus \$(0.28), our estimate \$(0.29).
- Pfizer to finalize SPA shortly with Ph3 start in H2/14. GLYC anticipates it will book a \$15M milestone from Pfizer in Q2 with an additional \$20M to be paid on trial start. Based on our conversations with KOLs, we expect high interest from clinical institutions in the trial, and expect site sign-up to be rapid. Given the nature of SCD crisis trials, enrollment time will be lengthy though.
- GMI-1271 set to enter the clinical for AML shortly. The E-selectin '1271 antagonist has the potential to address metastasis and thromboembolic complications. AML is the first indication for clinical development of the drug, and clinical trials are expected to begin next quarter. We hope for endpoint or potential biomarker around the end of 2014 or early 2015 depending on enrollment.

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Figure 1: GLYC P&L

	2011A	2012A	2013A	Q1/14A	Q2/14E	Q3/14E	Q4/14E	2014E	2015E	2016E	2017E	2018E	2019E	2020E
0.00.40=0														
GMI-1070	-	-	-	-	-	-	-	-	-	-	17.3	56.0	100.7	130.3
Product revenues	-	-	-	-	-	-	-	-	-	-	17.3	56.0	100.7	130.3
Collaboration revenue	3.8	15.3	21.9	-	13.50		10.00	23.5		-	60.0	50.0		50.0
Total revenues	3.8	15.3	21.9	-	13.5	-	10.0	23.5	-	-	77.3	106.0	100.7	180.3
Cost of goods sold		-	-	-	_	_	_							
Gross Profit	3.8	15.3	21.9	-	13.50	-	10.00	23.5	-	-	77.3	106.0	100.7	180.3
R&D expense	7.8	9.4	12.4	3.9	5.0	6.0	7.0	21.9	30.0	50.0	50.5	51.0	51.5	52.0
SG&A expense	2.1	2.2	2.1	1.2	1.3	1.4	1.4	5.3	3.0	3.5	3.7	3.9	4.1	4.3
Other operating expense		0.0	-	-	-			-	-	-	-	-	-	-
Total operating expense	9.9	11.6	14.5	5.1	6.3	7.4	8.4	27.2	33.0	53.5	54.2	54.9	55.6	56.3
Operating income	(6.1)	3.6	7.4	(5.1)	7.2	(7.4)	1.6	(3.7)	(33.0)	(53.5)	23.1	51.1	45.1	124.0
Net Interest/Investment income	0.0	0.0												0.0
(interest expense)	(0.0)	(0.0)	0.0	0.0	0.0	0.0	0.0	0.0						0.1
Other non-operating income (expense)	`- ′	(0.0)	-	0.0				-	-	-	-	-	-	-
Interest and other, Net	0.0	(0.0)	-	-	-	-	-	-	-	-	-	-	-	-
Pre-tax income	(6.1)	3.6	7.5	(5.1)	7.2	(7.3)	1.6	(3.6)	(33.0)	(53.5)	23.1	51.1	45.1	124.1
Income tax expense (benefit)	_	_	_					_	_	_	_	_	-	_
Net income (loss)	(6.1)	3.6	7.5	(5.1)	7.2	(7.3)	1.6	(3.6)	(33.0)	(53.5)	23.1	51.1	45.1	124.1
Basic EPS	(6.58)	3.93	0.61	(0.30)	0.42	(0.42)	0.09	(0.21)	(1.89)	(3.05)	1.31	2.89	2.53	6.94
Diluted EPS	(6.58)	0.33	0.61	(0.30)	0.42	(0.42)	0.09	(0.21)	(1.89)	(3.05)	1.31	2.89	2.53	6.94
Basic shares outstanding	0.9	0.9	12.3	17.2	17.3	17.4	17.5	17.4	17.4	17.5	17.6	17.7	17.8	17.9
Diluted shares outstanding	0.9	11.0	12.3	17.2	17.3	17.4	17.5	17.4	17.4	17.5	17.6	17.7	17.8	17.9
			-				-				-		-	

Source: Canaccord Genuity and company reports



Figure 2: GLYC pNPV analysis

Product Development

Drug name	Indication	Status	Launch	Years to Laui Launch	Years to nch plus 6	Success S	ales (US\$m)	Probability weighted Peak Sales (US\$m)	Royalty	Profitability	Probability weighted Peak Profit (US\$m)	Discount Factor	NPV (US\$)
GMI-1070	Sickle cell disease	Phase 3	2017	3	9	60%	1212.4	727.4	18%	90% Total	117.84	5.90	17.40 17.40

Source: Canaccord Genuity and company reports

Investment risks

Clinical risk -- GlycoMimetics' current Phase 3 trial may not be successful. We note that there has been relatively little drug development in SCD and no successful clinical trials that have led to the approval of drugs for the disease based on the endpoint GlycoMimetics is pursuing. Further, the Phase 2 trial of GMI-1070 did not meet its primary endpoint (the same endpoint in the Phase 3 trial design) due to data variability. However, we feel the expansion of the patient numbers in the upcoming Phase 3 trial will compensate for the data variability inherent to SCD trials.

Clinical risk -- Additional clinical investigation may show GMI-1070 to have an unacceptable safety and tolerability signal. GMI-1070's selectin-based mechanism could potentially interfere with immune responses to infection, thereby increasing risk of infections, opportunistic and otherwise. The only serious adverse event in the Phase 2 trial was one case that was controlled and resolved without discontinuation of treatment.

Regulatory risk -- GMI-1070 may not be approved by the FDA and/or EMA despite Phase 3 success. We note the only approved drug for treatment of SCD is indicated for the prevention of the number of crises, rather than reduction of the duration of crisis (the trial design). There is no precedent for the approval of SCD drugs based on reduction of length of hospitalization.

Competitive risk -- GlycoMimetics faces potential competition from other agents seeking to decrease the time to resolution of SCD crises, as well as indirect competition from agents being developed to prevent the onset of crises.

Commercialization risk -- There is little to no precedent for the successful promotion to the sickle cell disease market. Hydroxyurea, the only currently-approved drug for the treatment of SCD, has been available in multiple generic forms for SCD for a number of years, and is no longer promoted by Bristol-Myers, its original SCD sponsor. Commercial uptake has historically been extremely limited. As such, we see no precedent for the successful launch and promotion of a drug for SCD.

Reimbursement risk -- There is no guarantee that GlycoMimetics, or its partners, will garner reimbursement for GMI-1070. There has historically been significant skepticism regarding the market opportunity in SCD due to concerns about insurance coverage rates of the affected population. We also note GMI-1070's initially pursued indication would dictate its use in the hospital setting, which will require prior approval from hospital P&T committees.



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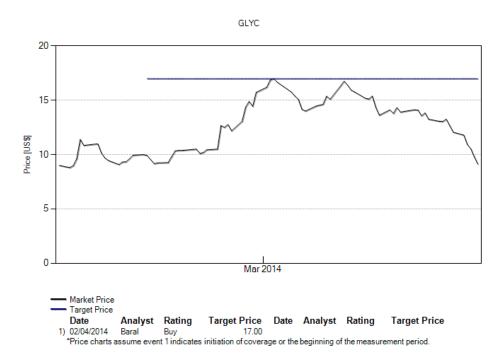
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Site Visit:

An analyst has visited GlycoMimetics' material operations in Gaithersburg, Maryland. No payment or reimbursement was received from the issuer for the related travel costs.

Price Chart:*



Distribution of Ratings: Global Stock Ratings (as of 31 March 2014)

Coverage Universe					
Rating	#	%	IB Clients %		
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Speculative Buy	43	4.4%	55.8%		
Hold	317	32.1%	13.2%		
Sell	45	4.6%	4.4%		
	988*	100.0%			

^{*}Total includes stocks that are Under Review



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Company	Disclosure
GlycoMimetics	1A, 2, 3, 5, 7

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