

SUMMARY

After clinical disappointments from Alzheimer's disease drugs that targeted the breakup of beta amyloid plaque, researchers turned their attention to tau, a brain-specific, axon-enriched protein. Beta amyloid plaque and tau appear to work together with distinct functions: Beta amyloid helps form the glue that holds plaque together after neurofibrillary tangles emerge from toxic tau. Beta amyloid can be described as the trigger, tau the bullet. The two proteins, however, seem to operate synergistically in a continuous feedback loop.

Several companies have advanced monoclonal antibodies into the clinic seeking to reduce the level of pTau as Alzheimer's treatments. Cassava Sciences is developing PTI-125 as a small molecule that restores the normal shape and function of filamin A (FLNA) in the brain. *SanaCurrents* assigns a **low pivotal sentiment** that Cassava will report positive topline results from its phase IIb trial of PTI-125 by mid-year 2020. Full results are expected by the fourth quarter.

Cassava's research on FLNA began 10 years ago with NIH funding. Cassava researchers assert when FLNA, a scaffolding protein, becomes altered, the normal function of neurons is disrupted. By correcting the FLNA protein, PTI-125 improves the function of certain brain receptors, consequently slowing neurodegeneration and exerting strong anti-neuroinflammatory effects.

In a 13-patient, open label phase IIa trial, Cassava reported significant reductions ($p < 0.01$) in both nitrated and phosphorylated forms of the tau protein; **each patient exhibited biomarker responses** to PTI-125. The results, presented in December 2019, sent Cassava's stock up more than 4-fold in just three weeks.

Cassava has enrolled 64 patients in the phase IIb trial. Unlike the previous trial, the phase IIb will be a randomized, double blind, placebo-controlled, trial in US patients with mild-to-moderate Alzheimer's disease. Patients will receive placebo, 50 mg or 100 mg of PTI-125 twice daily over 28 days. The trial objective is to investigate the safety and effects of PTI-125 on biomarkers of disease. The primary endpoint is improvement in patients' biomarkers of disease from baseline to Day 28.

THE EDGE

Cassava's stock shot up in December because PTI-125 demonstrated through biomarker measurement it potentially could pinpoint an Alzheimer's diagnosis through blood and serum tests.

While disabling the trigger function of beta amyloid plaque was regarded as the Holy Grail in treating Alzheimer's – as well as a mega-blockbuster – the drugs simply did not work. One reason often cited for the failures was the drugs had few *specific* targets. The Cassava approach is much more granular. In the phase IIa trial, PTI-125 produced declines in four types of tau: p-tau-T181, p-tau-T202, p-tau-T231, and a nitrated form of tau—n-tau-Y29. Cassava contends these cumulative declines can influence amyloid beta production, as well as both neuroinflammation and tau pathology.

PROBABILITY SENTIMENT

- SUPERIOR (highest)
- Pivotal
- Advantageous

Key Catalyst(s)

- Topline results from phase IIb trial of Alzheimer's drug PTI-125

Key Catalyst Date(s)

- mid-year 2020

Insider & Institutional Holdings

20.19% % of Shares Held by All Insider

39.17% % of Shares Held by Institutions

49.08% % of Float Held by Institutions

36 Number of Institutions Holding Shares

Key Executives

Mr. Remi Barbier, Founder, Chairman, CEO & Pres

Dr. Nadav Friedmann, COO, Chief Medical Officer & Director

Dr. George Thornton, Sr. VP of Technology

Mr. Michael Zamloot, Sr. VP of Technical Operations

Dr. Michael Marsman Pharm.D., Sr. VP of Regulatory Affairs

Location

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Austin, TX 78731
512-501-2444
<http://www.cassavasciences.com>

General Guidelines for SanaCurrents Strategy

Based on SanaCurrents' analytical model, Superior and Pivotal sentiments reflect a probability score of at least 60% that the company will announce a positive result to the specified catalyst. In our experience, a positive result typically increases the company's share price. An Advantageous sentiment reflects a score of less than 60%.

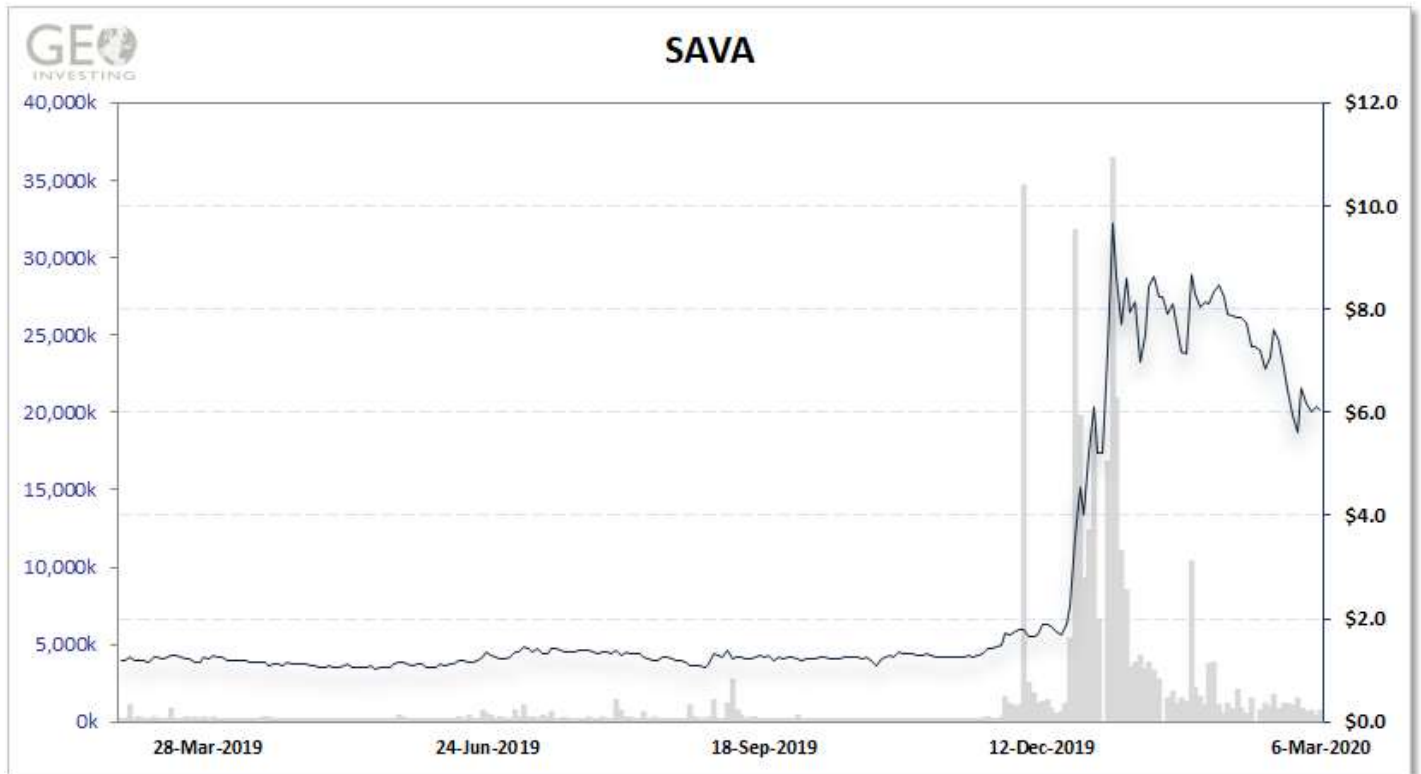
Cassava Sciences (NASDAQ:SAVA)

THE EDGE cont'd

Another edge for Cassava's phase IIb trial is the endpoint of improvement in patients' biomarkers. Patients' cognition and aptitude will not be assessed broadly. Rather, only the reduction in biomarkers at the two scheduled doses compared to placebo will be assessed. This endpoint, combined with the phase IIa results, point to Cassava repeating its earlier success.

Cassava has a long path ahead to demonstrate if PTI-125 can be the mega-blockbuster expected of the first approved Alzheimer's drug. The company, however, has a near-term plan to elevate its clinical standing and valuation.

Cassava is developing a diagnostic test called PTI-125Dx to detect Alzheimer's disease using only a blood sample. Such a comprehensive diagnostic does not exist now. Positive results from the phase IIb trial would validate, and likely accelerate, the PTI-125Dx test, pushing the company's market cap well past its current level of \$105 million.



Cassava Sciences (NASDAQ:SAVA)

Advanced Tearsheet

Cassava Sciences Inc		USD 6.04		-0.09		-1.31%	
Country	United States of America	Next PQFH End Date	Dec 31, 2019				
Exchange	NASDAQ Stock Market	Next FY End Date	Dec 31, 2019				
GICS Sector	Health Care	Next earnings Date	Nov 23, 2020				
GICS Industry	Pharmaceuticals	Latest earnings Date	Oct 29, 2019				
Currency	USD						

General Information		*in millions USD, except per share data	
Price (Previous Close)	6.04	Div. Yield, Last year (FY0)	
Price - 52 Week High	10.96	Shares Outstanding	17.2 MM
Price - Current vs 52wk High	-44.04%	Short Interest Shares	1.4 MM
Price - 52 Week Low	1.00	SI % of O.S	8.36%
Price - Current vs 52wk Low	504.00%	Market Capitalization	104
Beta - Sampling last 1y, Daily	0.95 vs S&P 500	Net Debt	20
		Enterprise Value	86

Estimates revision momentum		*in millions USD, except EPS				
Last reported FY	Current Year FY1			Next Year FY2		
	6mth ago	3mth ago	Today	6mth ago	3mth ago	Today
Consensus date	9/7/2019	12/7/2019	3/7/2020	9/7/2019	12/7/2019	3/7/2020
Revenue	0	0	0	25	25	25
% diff to today		100%	100%	-100.0%	-100.0%	
Gross Profit	0			19	19	
% diff to today		100%	100%	100%	100%	
EBIT	-4	-3	-6	12	15	15
% diff to today		-21.3%	0.0%	20.9%	110%	
EBITDA	-4	-8	-8	39	39	39
% diff to today		0.0%	0.0%	0.0%	0.0%	
EPS	-0.43	-0.31	-0.31	0.00	0.14	0.14
% diff to today		-17.4%	0.0%	70.0%	110%	

Consolidated Income Statement		*in millions USD, except diluted EPS (adj.) and common shares					
Period end date	FY.5	FY.4	FY.3	FY.2	FY.1	FY0	5yr-CAGR
	12/31/2013	12/31/2014	12/31/2015	12/31/2016	12/31/2017	12/31/2018	
Sales	41	0	0	0	0	0	-100.0%
Gross Income	41	0	0	0	0	0	-100.0%
EBITDA (adj.)	31	-12	-14	-16	-12	-4	
EBIT (adj.)	21	-8	-11	-15	-8	-4	
Earnings Before Tax (adj.)	31	-8	-11	-15	-8	-4	
Net Income (adj.)	32	-11	-15	-16	-8	-4	
Diluted EPS (adj.)	4.30	-1.88	-2.17	-2.31	-1.82	-1.17	
Common Shares	7	7	7	7	7	17	21.0%

Margins & Profitability Ratios		vs. last 5y					
Period end date	FY.5	FY.4	FY.3	FY.2	FY.1	FY0	average (right)
	12/31/2013	12/31/2014	12/31/2015	12/31/2016	12/31/2017	12/31/2018	
Gross Margin	100.0%						
R&D % of Sales	12.0%						
SG&A % of Sales	11.0%						
EBITDA Margin (adj.)	76.3%		0.0%				
EBIT Margin (adj.)	76.2%						
Net Margin	76.7%						
Tax Rate	-0.2%						
ROA	39.0%		-29.4%	-50.0%			
ROE	102.7%		-30.8%	-61.3%			

Per Share Metrics		*in USD		
Book Value	FY.2	FY.1	FY0	FY1
	2.80	1.47	1.14	0.90
Free Cash Flow	FY.2	FY.1	FY0	FY1
	-1.97	-1.2E	-0.28	
Dividend	FY.2	FY.1	FY0	FY1
Earnings (EPS)	FY.2	FY.1	FY0	FY1
	0.31	-1.82		-0.31

Valuation ratios		*in USD		
EV/Sales	LTM	NTM	FY1	FY2
				3.5 x
EV/EBITDA	LTM	NTM	FY1	FY2
	100x		100x	2.2 x
PE	LTM	NTM	FY1	FY2
	100x		100x	43.1 x
PE to Growth (PEG)	LTM	NTM	FY1	FY2
	100x		100x	
EV/FCF	LTM	NTM	FY1	FY2
	100x		100x	
P/Sales	LTM	NTM	FY1	FY2
	6.0 x		6.0 x	4.2 x
P/BV	LTM	NTM	FY1	FY2
	6.0 x		6.0 x	
P/Tangible BV	LTM	NTM	FY1	FY2
	6.0 x		6.0 x	

Consolidated Balance Sheet		*in millions USD	
Period end date	FY.2	FY.1	FY0
	12/31/2016	12/31/2017	12/31/2018
Cash & ST Investments	19	10	
Inventory			
Total Assets	19	11	20
Current Liabilities	1	1	1
Total Debt	0	0	0
Total Liabilities	1	1	1
Total Shareholders Equity	19	10	20

Consolidated Cash Flow Statement		*in millions USD	
Period end date	FY.2	FY.1	FY0
	12/31/2016	12/31/2017	12/31/2018
Net Income	15		
D&A	6	6	0
Change in WC	-2	1	-1
Cash From Operations	12	8	5
Capital Expenditures	0	0	
Cash From Investments	-2	2	8
Shares Repurchased			
Cash From Financing	0	0	14
Cash & Cash Eq - Period end	17	10	20

Disclosure of General Guidelines for SanaCurrents Strategy

In most situations, SanaCurrents partners take a position in a stock (or buy shares) shortly after a report on a catalyst is published. This is particularly true if the catalyst is within 90 days of the published report. For catalysts with horizons of 6-9 months, or longer, SanaCurrents anticipates the biopharma and device stocks from which it selects catalysts can rise or fall 20-30% in the months following the date of the published report. While buying at a dip, naturally, is preferred, SanaCurrents does not forecast or model in swings in share price prior to the catalyst. SanaCurrents' analytics score models the probability a compelling catalyst will be positive. SanaCurrents expects a positive announcement regarding a catalyst will drive a company's stock to higher level than the price on the date of the report, based on the company's value and capital structure when the report was published. That said, subscribers should exercise their preferred, individual discipline when timing trades, buy or sell, regarding the catalyst forecast.

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