

Specialty Pharmaceuticals

Intercept Pharmaceuticals, Inc.

(ICPT) - BUY

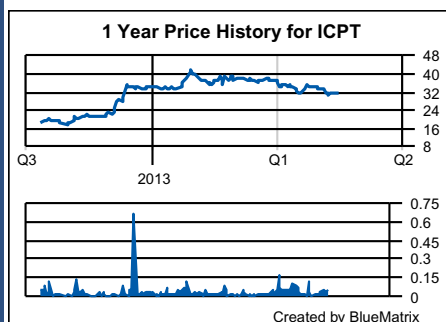
Price: **\$31.95**
Fair Value Estimate: \$48.00
52-Week Range: \$15.00-\$42.67
Market Cap (MM): \$537
Shr.O/S-Diluted (mm): 16.8
Average Daily Volume: 31,977
Dividend: NA
Book Value: \$0.52

FYE: Dec	2012A	2013E	2014E
Revenue (M):	\$2.4A	\$1.6E	\$1.6E

Quarterly Revenue (M):			
Q1	\$0.8A	\$0.4A	\$0.4E
Q2	\$0.8A	\$0.4E	\$0.4E
Q3	\$0.5A	\$0.4E	\$0.4E
Q4	\$0.4A	\$0.4E	\$0.4E

FYE: Dec	2012A	2013E	2014E
EPS:	\$(7.36)A	\$(2.09)E	\$(2.31)E
Prior EPS:		NC	NC

Quarterly EPS:			
Q1	\$(1.03)A	\$(0.62)A	\$(0.57)E
Q2	\$(1.75)A	\$(0.51)E	\$(0.56)E
Q3	\$(1.86)A	\$(0.52)E	\$(0.58)E
Q4	\$(2.02)A	\$(0.54)E	\$(0.59)E



Equity Research
Note

1Q13 Looks Fine, 2H13 Data Next Drivers

INVESTMENT CONCLUSION:

ICPT reported its 1Q13 figures yesterday which were roughly in line with JMS and consensus estimates. With the recent positive news from the interim look at the primary biliary cirrhosis (PBC) "supergroup" study in late April, we remain confident that the odds favor positive phase 3 data from the POISE trial on obeticholic acid (OCA) for PBC. This trial is expected to read out mid-14 and we continue to believe that the FDA will accept ICPT's surrogate endpoint as appropriate for predicting clinical outcomes, as the supergroup interim analysis demonstrates. 2H13 should be busy for ICPT as we anticipate seeing the full data set from the supergroup analysis as well as top-line data from the phase 2 trials of OCA in portal hypertension and bile acid diarrhea. We continue to see ICPT as one of our top small cap picks in specialty pharma, and we reiterate our BUY rating, \$48 fair value.

KEY POINTS:

- **Financials in line.** ICPT reported its 1Q13 with revenue of \$405M vs JMS \$400: (Cons \$500M) and EPS of (\$0.62) vs JMS (\$0.52) and Cons (\$0.39). R&D expenses were in-line with estimates coming in at \$4.8M vs JMS of \$5.2M and SG&A came in slightly below at \$2.4M vs JMS of \$3.2M. EPS differed on the non-cash item of revaluation of warrants which was (\$3.6M).
- **"Supergroup" interim data looked positive.** In late April ICPT announced the positive results of the "supergroup" retrospective analysis of over 2,100 patients with primary biliary cirrhosis (PBC) at the European Association for the Study of the Liver (EASL). The data presented further substantiated that the primary endpoint of lowering serum liver makers used in their phase 3 POISE trial were predictive of clinical outcomes. With the European Medicines Agency (EMA) already on board, we believe that this confirmatory study should tip the scales in the eyes of the FDA.
- **POISE data remains on-track for 2Q14.** We believe that the P3 POISE trail is setup for success as it closely mimics the successful phase 2 trial. Historically in clinical trial design, we've found that phase 3 trials which closely resemble successful phase 2 trails tend to have the best chance for success. Applying the phase 3 endpoints to the 10mg cohort of the phase 2 trial shows 40% of patients hitting the phase 3 trial primary endpoint (p=0.012).
- **PESTO & OBEDIAH top line 4Q13.** We continue to anticipate the phase 2 trials in portal hypertension (PESTO) and bile acid diarrhea (OBEDIAH) in 4Q13. While smaller markets than the anticipated 8M patient indication in non-alcoholic steatohepatitis, these could provide good validation of the efficacy of OCA in helping to repair liver function.
- **Maintain Buy rating, \$48 Fair Value:** Our \$48/share sum-of-the-parts is based on OCA for PBC at \$33/share, OCA for NASH at \$9/share, and \$6/share for cash (end 2014) and technology value.

Research Analyst Certifications and Important Disclosures are on pages 5 - 6 of this report

EXHIBIT 1:

Sum-of-the-parts valuation: ICPT		
Segment	Valuation (000's)	Per share value
OCA for PBC in the US	\$678,791	\$33
OCA for NASH in the US	\$206,017	\$9
Cash (end of '14E) & tech value	\$144,676	\$6
	\$1,128,174	\$48
2014 fully diluted shares out		23,433

Source: Janney estimates

EXHIBIT 2:

Intercept Pharmaceuticals

Potential clinical trial timeline estimates

	2012A				2013E				2014E				2015E				2016E				2017E				2018E							
	1QA	2QA	3QA	4QA	1QA	2QE	3QE	4QE	1QE	2QE	3QE	4QE	1QE	2QE	3QE	4QE	1QE	2QE	3QE	4QE	1QE	2QE	3QE	4QE	1QE	2QE	3QE	4QE				
OCA (Obeticholic acid) for Primary Biliary Cirrhosis (PBC)																																
POISE US/EU phase 3 trial - started Jan 2012	phase 3 - US								data																							
PBC 4,000 patient Supergroup analysis, full data 4Q13	supergroup analysis				intrm				data																							
Phase 2 Long Term Safety trial (Monotherapy)	phase 2																															
NDA & EMA filing (likely partnered in EU)									NDA/EMA																							
FDA approval & launch (6-mo approval)													FDA								LAUNCH											
EU approval & launch (12-mo approval)																	EU								LAUNCH							
POISE US phase 3 outcomes trial (minimum 5 years to run)									phase 3 outcomes trial																							
OCA (Obeticholic acid) for Nonalcoholic steatoheptitis (NASH) liver disease																																
FLINT US phase 2b - enrollment completed	Phase 2b (march 2011 start)								data																							
FLINT US phase 3													phase 3								data											
FLINT US phase 3 outcomes trial (~5 years)																					phase 3 outcomes trial											
NDA filing																									NDA							
Japan OCA (NASH) - partner Dainippon Sumitomo Pharma (DSP)																																
Japanese phase 2 trial	phase 2 - Japan (two trials)																data															
Japanese phase 3 trial																									phase 3 - Japan							
OCA (Obeticholic acid) for Portal Hypertension in patients with cirrhosis																																
PESTO EU phase 2a 10mg trial	phase 2				data																											
PESTO EU phase 2a safety trial 25mg	phase 2								data																							
PESTO EU phase 2b trial													phase 2b				data															
PESTO EU phase 3 trial																					phase 3				data							
OCA - Bile Acid Diarrhea																																
OBEDIAH EU phase 2a trial - started July 2012	phase 2a								data																							
Additional clinical development									additional clinical development																							
INT-767 FXR agonist for diabetic nephropathy & chronic kidney diseases																																
Enabling studies, pre-IND									pre-IND studies																							
File IND with FDA																									IND							
Initiate clinical development																													timeline uncertain			

Source: Company reports and Janney estimates

Specialty Pharmaceuticals
Jim Molloy (617) 371-1528 jmolloy@janney.com

Intercept Pharmaceuticals

Quarterly income statement

(\$000's except per share)	2012A				2012A Year	2013E				2013E Year
	1QA	2QA	3QA	4QA		1QA	2QE	3QE	4QE	
Revenues										
License fees	\$759	\$759	\$523	\$405	\$2,446	\$405	\$400	\$400	\$400	\$1,605
Total Revenues	\$759	\$759	\$523	\$405	\$2,446	\$405	\$400	\$400	\$400	\$1,605
Expenses										
Cost of Goods Sold	0	0	0	0	0	0	0	0	0	0
Gross Margin	759	759	523	405	2,446	405	400	400	400	1,605
Research and Development	3,060	5,018	3,318	4,787	16,183	4,833	6,250	6,500	6,500	24,083
SG&A	1,059	944	991	2,183	5,177	2,397	3,500	3,500	4,000	13,397
Total Operating Expenses	4,119	5,962	4,309	6,970	21,360	7,229	9,750	10,000	10,500	37,479
Income (loss) from Ops	(3,360)	(5,203)	(3,786)	(6,565)	(18,914)	(6,824)	(9,350)	(9,600)	(10,100)	(35,874)
Warrant Revaluation income	678	301	(1,418)	(24,187)	(24,626)	(3,683)				(3,683)
FOREX loss on liquidation		(198)								
Interest & dividend income	10	7	13	61	(100)	296	10	10	10	326
Interest expense	(7)	0	3	0	(4)					0
QTP grant					0					0
Pretax Income (Loss)	(2,680)	(5,092)	(5,187)	(30,691)	(43,644)	(10,210)	(9,340)	(9,590)	(10,090)	(39,230)
Div. pref stock, not declared	(750)	(750)	(1,000)	(130)	(2,630)	0				0
Net income/(loss)	(3,430)	(5,842)	(6,187)	(30,821)	(46,274)	(10,210)	(9,340)	(9,590)	(10,090)	(39,230)
EPS as reported	(\$1.03)	(\$1.75)	(\$1.86)	(\$2.02)	(\$7.36)	(\$0.62)	(\$0.48)	(\$0.49)	(\$0.51)	(\$2.10)
Shares out (000)	3,330	3,330	3,330	15,223	6,283	16,558	19,308	19,558	19,808	18,808
Fully diluted shares (000)	11,725	14,326	14,326	19,238	19,238	19,423	22,058	22,308	22,558	21,587

Source: Company reports and Janney estimates

Specialty Pharmaceuticals
Jim Molloy (617) 371-1528 jmolloy@janney.com

Intercept Pharmaceuticals

Annual income statement

(\$000's except per share)	2012A	2013E	2014E	2015E	2016E	2017E	2018E
Revenues							
OCA sales	-	-	-	\$27,799	\$150,913	\$396,849	\$609,266
OCA royalties (EU & Japan)	-	-	-	-	6,849	35,454	60,236
License fees	\$2,446	\$1,605	\$1,600	1,600	1,800	2,000	2,000
Total Revenues	\$2,446	\$1,605	\$1,600	\$29,399	\$159,562	\$434,303	\$671,502
Expenses							
Cost of Goods Sold	-	-	-	6,745	22,637	59,527	73,112
Gross Margin	2,446	1,605	1,600	22,654	136,925	374,776	598,390
Research and Development	16,183	24,083	28,000	30,000	34,000	45,000	65,000
SG&A	5,177	13,397	18,250	25,000	41,500	93,750	132,500
Total Operating Expenses	21,360	37,479	46,250	55,000	75,500	138,750	197,500
Income (loss) from Ops	(18,914)	(35,874)	(44,650)	(32,346)	61,425	236,026	400,890
Warrant Revaluation income	(24,626)	(3,683)	-	-	-	-	-
FOREX loss on liquidation	-	-	-	-	-	-	-
Interest & dividend income	(100)	326	40	60	75	100	100
Interest expense	(4)	-	-	-	-	-	-
Pretax Income (Loss)	(43,644)	(39,230)	(44,610)	(32,286)	61,500	236,126	400,990
Taxes	-	-	-	-	-	47,225	136,337
Div. pref stock, not declared	(2,630)	-	-	-	-	-	-
Net income/(loss)	(46,274)	(39,230)	(44,610)	(32,286)	61,500	188,901	264,654
EPS as reported	(\$7.36)	(\$2.10)	(\$2.18)	(\$1.51)	\$2.52	\$6.54	\$8.80
Shares out (000)	6,283	18,808	20,433	21,558	22,708	23,883	25,083
Fully diluted shares (000)	19,238	21,587	23,433	25,058	24,408	28,883	30,083
Margin & expense analysis							
COGS as % prod sales	NA	NA	NA	24%	14%	14%	11%
R&D	NA	NA	NA	102%	21%	10%	10%
SG&A	NA	NA	NA	85%	26%	22%	20%
Op. margin cont. ops	NA	NA	NA	-110%	38%	54%	60%
Taxes	NA	NA	NA	0%	0%	20%	34%
Net margin	NA	NA	NA	-110%	39%	43%	39%
Year-over-year change							
Net revenue	NA	NA	NA		443%	172%	55%
R&D	42%	49%	16%	7%	13%	32%	44%
SG&A	23%	159%	36%	37%	66%	126%	41%
Operating income	NA	NA	NA	NA	NA	284%	70%
Net income	NA	NA	NA	NA	NA	207%	40%

Specialty Pharmaceuticals

Source: Company reports and Janney estimates

Jim Molloy (617) 371-1528 jmolloy@janney.com

Company Description

Intercept Pharmaceuticals is a development stage biopharmaceutical company that is developing therapeutics to treat chronic liver diseases utilizing proprietary bile acid chemistry. The lead candidate is obeticholic acid (OCA), a bile acid analog and farnesoid X receptor (FXR) agonist that is in phase 3 trials for the treatment of primary biliary cirrhosis. ICPT is also conducting phase 2a clinical trials with OCA to treat portal hypertension; phase 2b clinical trials for the nonalcoholic steatohepatitis treatment (NASH) in sponsored by the National Institute of Diabetes and Digestive and Kidney Diseases; and phase 2a trials for the treatment of bile acid diarrhea.

IMPORTANT DISCLOSURES

Research Analyst Certification

I, Jim Molloy, the Primarily Responsible Analyst for this research report, hereby certify that all of the views expressed in this research report accurately reflect my personal views about any and all of the subject securities or issuers. No part of my compensation was, is, or will be, directly or indirectly, related to the specific recommendations or views I expressed in this research report.

Janney Montgomery Scott LLC ("Janney") Equity Research Disclosure Legend

Janney Montgomery Scott LLC intends to seek or expects to receive compensation for investment banking services from Intercept Pharmaceuticals, Inc. in the next three months.

The research analyst is compensated based on, in part, Janney Montgomery Scott's profitability, which includes its investment banking revenues.

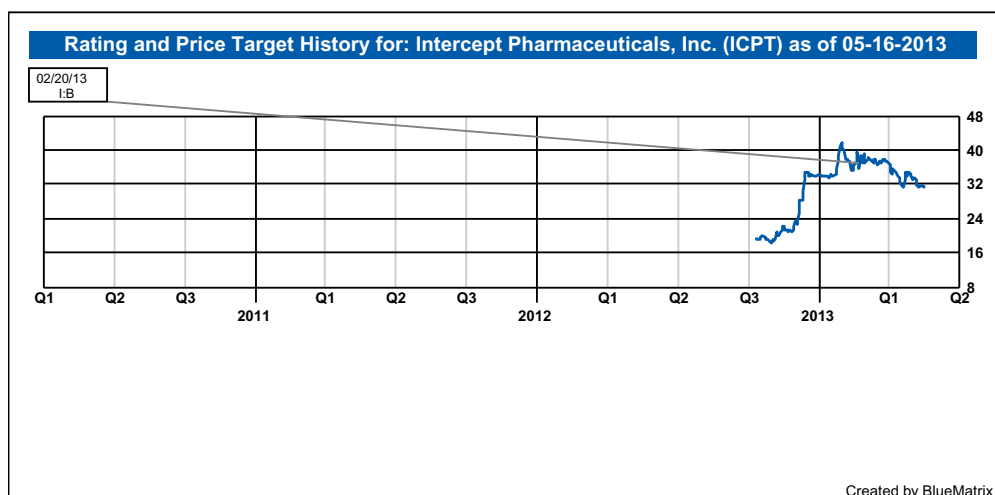
Definition of Ratings

BUY: Janney expects that the subject company will appreciate in value. Additionally, we expect that the subject company will outperform comparable companies within its sector.

NEUTRAL: Janney believes that the subject company is fairly valued and will perform in line with comparable companies within its sector. Investors may add to current positions on short-term weakness and sell on strength as the valuations or fundamentals become more or less attractive.

SELL: Janney expects that the subject company will likely decline in value and will underperform comparable companies within its sector.

Price Charts



Janney Montgomery Scott Ratings Distribution as of 3/31/13

Rating	Count	Percent	IB Serv./Past 12 Mos.	
			Count	Percent
BUY [B]	214	52.58	38	17.76
NEUTRAL [N]	189	46.44	20	10.58
SELL [S]	4	0.98	0	0.00

***Percentages of each rating category where Janney has performed Investment Banking services over the past 12 months.**

Other Disclosures

Janney Montgomery Scott LLC, is a U.S. broker-dealer registered with the U.S. Securities and Exchange Commission and a member of the New York Stock Exchange, the Financial Industry Regulatory Authority and the Securities Investor Protection Corp.

This report is for your information only and is not an offer to sell or a solicitation of an offer to buy the securities or instruments named or described in this report. Interested parties are advised to contact the entity with which they deal or the entity that provided this report to them, should they desire further information. The information in this report has been obtained or derived from sources believed by Janney Montgomery Scott LLC, to be reliable. Janney Montgomery Scott LLC, however, does not represent that this information is accurate or complete. Any opinions or estimates contained in this report represent the judgment of Janney Montgomery Scott LLC at this time and are subject to change without notice.

Investment opinions are based on each stock's 6-12 month return potential. Our ratings are not based on formal price targets, however, our analysts will discuss fair value and/or target price ranges in research reports. Decisions to buy or sell a stock should be based on the investor's investment objectives and risk tolerance and should not rely solely on the rating. Investors should read carefully the entire research report, which provides a more complete discussion of the analyst's views. Supporting information related to the recommendation, if any, made in the research report is available upon request.