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# **Endocyte, Inc. (ECYT)**

Reports Q1:11 EPS, Conditional EU Approval Process To Begin Q3:11

- Endocyte reported Q1:11 earnings; importantly, the company announced that it began enrolling the Phase III PROCEED trial of EC145 (with EC20, the companion prognostic). Recall that the PROCEED trial will treat platinum-resistant ovarian cancer patients whose tumors over-express the folate receptor (identified through use of EC20). The company plans to enroll more than 500 patients, with a primary endpoint of progression-free survival and also measure overall survival benefits.
- The conditional marketing application for EC145 and EC20 for the EU could be filed in August. Recall that the company announced it would seek conditional approval for EC145 and EC20 in the EU based on its positive Phase II PRECEDENT trial in platinum-resistant ovarian cancer patients with tumors that over-express the folate receptor.
- Two recent conditional approvals—Arzerra and Votrient—took 14 and 16 months, respectively, from application submission to conditional approval.
- We estimate EU revenues beginning in 2013, based on accelerated approval
  in the region. Recall that conditional approval in the EU allows access to
  therapies in areas of high unmet medical need. Recall also that a conditional
  marketing application is valid for 1 year, renewable and can be converted to full
  marketing authorization upon completion of obligatory studies determined by the
  EMA.
- Reiterate OUTPERFORM rating and \$20 price target. We arrive at our \$20 price target by taking the sum of the per share values of US sales and EU royalties in 2015. We value the US opportunity for EC145 in the platinum-resistant ovarian cancer setting at \$10 which is derived by taking 6x our 2015 sales estimate, discounted at 25% annually. We value the EU opportunity for EC145 in the platinum-resistant ovarian cancer setting at \$10 a share, based upon a 6x multiple of EU royalties, discounted 25% annually.
- Risks to our price target include 1) failure of EC145 for PROC in the clinic; and 2) failure to receive marketing approval in the US or the EU for EC145 for PROC.

FYE Dec	2010A		2011E			2012E	
REV	ACTUAL	CURR.	PREV.	CONS.	CURR.	PREV.	CONS.
Q1 Mar	\$0.0A	\$0.0A			\$0.0E		\$0.0E
Q2 Jun	0.0A	0.0E		0.0E	0.0E		0.0E
Q3 Sep	0.0A	0.0E		0.0E	0.0E		0.0E
Q4 Dec	0.0A	0.0E		0.0E	0.0E		0.0E
Year*	\$0.0A	\$0.0E		\$0.0E	\$0.0E		\$6.0E
Change							
	2010A		2011E			2012E	
EPS	ACTUAL	CURR.	PREV.	CONS.	CURR.	PREV.	CONS.
Q1 Mar	\$0.00A	(\$0.43)A	(\$0.55)A		(\$0.31)E	(\$0.38)E	(\$0.38)E
Q2 Jun	0.00A	(0.23)E	(0.35)E	(0.30)E	(0.33)E	(0.43)E	(0.43)E
Q3 Sep	(1.10)A	(0.25)E	(0.36)E	(0.34)E	(0.31)E	(0.38)E	(0.38)E
Q4 Dec	(0.22)A	(0.28)E	(0.36)E	(0.37)E	(0.31)E	(0.37)E	(0.37)E
Year*	(\$1.32)A	(\$1.12)E	(\$1.57)E	(\$1.39)E	(\$1.25)E	(\$1.56)E	(\$1.25)E
P/E							
Change	93%	15%			-11%		

Consensus estimates are from Thomson First Call.

May 5, 2011

**Price** 

\$11.29

Rating

## **OUTPERFORM**

12-Month Price Target **\$20** 

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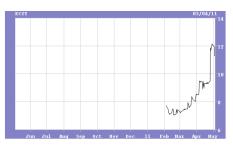
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Company Information	
Shares Outst (M)	29.7
Market Cap	\$335.1
52-Wk Range	\$6.15 - \$12.91
Book Value/sh	\$2.62
Cash/sh	\$3.19
Enterprise Value (M)	\$240.3
LT Debt/Cap %	0
Cash Burn (M)	\$-29.8

#### **Company Description**

Endocyte is developing novel small molecule drug conjugates and companion imaging prognostics. Their lead candidate, EC145, is in PIII trials in the PROC setting, targets cancers expressing the folate receptor that is also expressed on many solid tumors.



Source: Thomson Reuters

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<sup>\*</sup> Numbers may not add up due to rounding.



### Endocyte, Inc. (ECYT: NASDAQ; OUTPERFORM)

H1:11 End of Phase II meeting with EU EMEA for EC145 H1:11 Initiation of a Phase III trial for EC145 in women with platinum-resistant ovarian cancer H1:11 Initiation of a Phase III trial for EC20 folate imaging agent (companion prognostic for EC145) Jun. 4-8, 2011 Final Phase II PRECEDENT trial data for EC145 at ASCO (Chicago, IL) 6-month IPO lockup period ends Aug. 3, 2011 Potential submission of application for conditional approval in the EU based on Phase II Aug., 2011 PSMA targeting SMDC trial expected to be complete YE:11 Complete Phase I Trial for EC0489 DAVLBH payload SMDC in solid tumors YE:11 Complete Phase I Trial for EC0225 DAVLBH/mitomycin-C payload SMDC in solid tumors YE:11 Q1:12 Overall survival data from Phase IIb PRECEDENT trial of EC145 in platinum-resistant ovarian cancer

2012 Potential EU launch if accelerated approval granted by the EMA

2013 Final analysis of Phase III PROCEED trial of EC145 2014 Potential US launch of EC145 in PROC setting

# **Financial Model**



Gregory R. Wade, Ph.D.

5/5/2011

# Endocyte Inc.

Annual Financial Results & Projections (\$ in thousands except per share data)

Ticker: ECYT (Nasdag)

	FY:10A	FY:11E	FY:12E	FY:13E	FY:14E	FY:15E
Revenue:						
Sales	0	0	0	0	12,862	137,433
License fee and other revenues	0	0	0	9,772	38,395	83,402
Contracts	0	0	0	0	0	0
Total Revenues	\$0	\$0	\$0	\$9,772	\$51,257	\$220,836
Cost and Expenses:						
Costs of goods sold	0	0	0	0	1,929	20,615
Research and Development	14,561	20,601	27,797	27,611	28,423	40,604
Sales, General and Administrative	6,039	9,282	12,831	16,242	19,084	19,859
Other	0	0	0	0	0	0
Total Costs and Expenses	\$20,600	\$29,884	\$40,628	\$43,852	\$49,436	\$81,078
Operating Income (loss)	(20,600)	(29,884)	(40,628)	(34,081)	1,821	139,758
Net Interest Income (Expense)	(1,059)	51	797	988	702	1,246
Other income / (Expense)	1,564	3	0	0	0	0
Income Before Income Taxes	(20,095)	(29,830)	(39,831)	(33,093)	2,523	141,004
Net Income	(\$20,095)	(\$29,830)	(\$39,831)	(\$33,093)	\$7,265	\$129,337
GAAP Basic EPS with sFAS123	(1.32)	(1.12)	(1.25)	(1.01)	0.22	3.96
GAAP Diluted EPS with sFAS123	(1.32)	(1.12)	(1.25)	(1.01)	0.22	3.95
Weighted shares outstanding	15,220	26,521	31,813	32,869	32,403	32,698
Fully diluted shares outstanding	15,220	26,521	31,813	32,884	32,411	32,710



#### **Analyst Certification**

I, Gregory R. Wade, Ph.D., David M. Nierengarten, Ph.D., Christopher N. Marai, Ph.D., certify that the views expressed in this report accurately reflect my personal opinion and that I have not and will not, directly or indirectly, receive compensation or other payments in connection with my specific recommendations or views contained in this report.

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Outperform: Expect the total return of the stock to outperform relative to the median total return of the analyst's (or the analyst's team) coverage universe over the next 6-12 months.

Neutral: Expect the total return of the stock to perform in-line with the median total return of the analyst's (or the analyst's team) coverage universe over the next 6-12 months.

Underperform: Expect the total return of the stock to underperform relative to the median total return of the analyst's (or the analyst's team) coverage universe over the next 6-12 months.

The Investment Ratings are based on the expected performance of a stock (based on anticipated total return to price target) relative to the other stocks in the analyst's coverage universe (or the analyst's team coverage).\*

Rating Distribution (as of March 31, 2011)	Investment Banking Relationships (as of March 31, 2011)
Outperform: 55%	Outperform: 11%
Neutral: 37%	Neutral: 4%
Underperform: 8%	Underperform: 0%

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#### Wedbush Equity Research Disclosures as of May 5, 2011

Company	Disclosure
Endocyte, Inc.	1,3,4,5,7

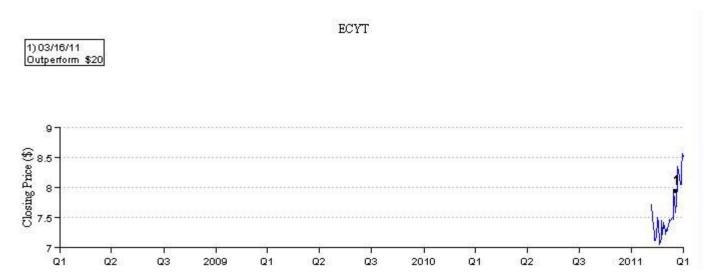
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\* WS changed its rating system from (Strong Buy/Buy/Hold/Sell) to (Outperform/ Neutral/Underperform) on July 14, 2009. Please access the attached hyperlink for WS' Coverage Universe: <a href="http://www.wedbush.com/services/cmg/equities-division/research/equity-research">http://www.wedbush.com/services/cmg/equities-division/research/equity-research</a> Applicable disclosure information is also available upon request by contacting Ellen Kang in the Research Department at (213) 688-4529, by email to <a href="mailto:ellen.kang@wedbush.com">ellen.kang@wedbush.com</a>, or the Business Conduct Department at (213) 688-8090. You may also submit a written request to the following: Business Conduct Department, 1000 Wilshire Blvd., Los Angeles, CA 90017.

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