Equity Research

Concert Pharmaceuticals, Inc.

CNCE: Q2 2014--Insight Into '354 Therapeutic Window

- Summary: On August 12, before the open, CNCE reported Q2 2014 earnings, featuring new data from the ongoing phase I work for CTP-354 that may support the potential for the drug to have a broader therapeutic window than traditional benzodiazepines, and warrant further exploration of the agent in the planned spasticity studies. Overall, we are encouraged by the continued progress with '354 and '499, as well as the partnered programs, and continue to believe Concert's valuation does not fully reflect the potential of its many proprietary and partnered programs. Adjusting 2014E/2015E EPS to -\$2.07/-\$1.09 from -\$1.88/-\$0.74. Adjusting valuation range to \$15-17 from \$21-23 based on updates to our model on potential out-year expenses and product launches.
- Financials: Concert reported a net loss of \$7.99MM, or EPS of -\$0.45, and ended Q2 with cash and equivalents of \$98.3MM. A slight uptick in quarterly R&D expense to \$6.2MM from \$5.8MM in Q2 2013 reflected increased spending on clinical development of '354, and the company expects this to continue to increase in H2 2014.
- New receptor occupancy data continues to support potential for '354 to demonstrate a high therapeutic index and long half-life supporting once-daily dosing, both of which could be important for commercial differentiation from marketed anti-spasticity therapies benzodiazepines. On the earnings call, Concert presented receptor of occupancy data for two patients receiving 6mg QD doses of '354, the first such data for the drug in repeat dosing, demonstrating that the drug provided GABA-A receptor occupancy of approximately 50% at five hours following dosing. There were indirect suggestions of a long half-life, which could speak to the potential for '354 to have differentiated 1x/day dosing, and no appreciable brain accumulation was observed. Both patients had very similar plasma concentrations and receptor occupancies, suggesting potential for good consistency patient to patient--a positive, though patient numbers are very small. The 50% occupancy approached the company's goal of 60% observed with single 20mg doses and which they had modeled might be achievable with the repeated 6mg dose, perhaps falling slightly below this bar due to the two patients having relatively high weights (220lbs on overage) which may have impacted the drug's volume of distribution. As expected, some neurologic side effects were seen, particularly transient dizziness and sleepiness, though it appears such side effects are likely more mild than the overt sedation that might be expected from traditional benzos dosed to reach comparable receptor occupancies. This suggests the potential for a broader therapeutic window, though we believe further escalation of the upper end of the dose range (CNCE is now assessing 12mg in the MAD study) and phase II studies assessing activity should better clarify '354's degree of differentiation.

Valuation Range: \$15.00 to \$17.00 from \$21.00 to \$23.00

Our valuation range is based on applying a 30x multiple to our 2022 estimated EPS and discounting at 15%, blended with 6x multiple of 2022 estimated sales, and discounting 12%. Key risks, in our view, are failure of '354 and/or '499 to show efficacy in subsequent studies and regulatory hurdles in spasticity or CKD.

Investment Thesis:

We believe Concert's proprietary and partnered candidates, and drug deuteration platform, will generate long-term value.

Please see page 5 for rating definitions, important disclosures and required analyst certifications
All estimates/forecasts are as of 08/12/14 unless otherwise stated.

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Outperform / V

Sector: Biotechnology Market Weight

Earnings Estimate Revised Down

	2013A	2014	2014E		2015E	
EPS		Curr.	Prior	Curr.	Prior	
Q1 (Mar.)	NE	(\$0.76) A	NC	NE		
Q2 (June)	NE	(0.45) A	(0.32)	NE		
Q3 (Sep.)	NE	(0.48)	(0.45)	NE		
Q4 (Dec.)	(4.66)	(0.50)	(0.49)	NE		
FY	(\$4.99)	(\$2.07)	(1.88)	(\$1.09)	(0.74)	
CY	(\$4.99)	(\$2.07)		(\$1.09)		
FY P/EPS	NM	NM		NM		
Rev.(MM)	\$25,408	\$4,608		\$22,520		

Source: Company Data, Wells Fargo Securities, LLC estimates, and Reuters NA = Not Available, NC = No Change, NE = No Estimate, NM = Not Meaningful V = Volatile, NO = Company is on the Priority Stock List

2014 quarterly EPS do not sum to full-year figure due to share count calculations.

Ticker	CNCE
Price (08/12/2014)	\$8.47
52-Week Range:	\$7-17
Shares Outstanding: (MM)	17.9
Market Cap.: (MM)	\$151.6
S&P 500:	1,932.52
Avg. Daily Vol.:	149,418
Dividend/Yield:	\$0.00/0.0%
LT Debt: (MM)	\$0.0
LT Debt/Total Cap.:	0.0%
ROE:	NM
3-5 Yr. Est. Growth Rate:	NM
CY 2014 Est. P/EPS-to-Growth:	NM
Last Reporting Date:	08/12/2014
	Before Open

Source: Company Data, Wells Fargo Securities, LLC estimates, and Reuters

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Together we'll go far



Company Description:

Concert Pharmaceuticals (CNCE), Inc., headquartered in Lexington, Massachusetts, is a biotechnology company focused on improving therapies in a variety of areas such as neurology and kidney disease, using its platform technology around deuterium substitution. The company's lead development candidate is CTP-354, a GABA-A partial agonist in development for muscle spasticity. Another development candidate is CTP-499, a phosphodiesterase enzyme inhibitor in development for diabetic kidney disease. Alongside '354 and '499, its partnered pipeline includes CTP-730, a deuterated anti-inflammatory drug partnered with Celgene, AVP-786 for major depressive disorder partnered with Avanir, and JZP-386, a deuterated version of Xyrem for narcolepsy, partnered with Jazz.

• FDA feedback on '499 path forward should help set stage for partnership discussions. The company was optimistic after receiving FDA feedback suggesting a standard phase III design, likely 96 weeks in duration, which could potentially be comprised of a single study, use an enriched population, and contain a higher dose, could be possible. FDA also signaled their comfort with safety and view that the drug has activity. CNCE continues to pursue a partnership for the asset, and we believe such regulatory clarity, and aspects of the pivotal design which could potentially reduce the costs a partner might bear, could be attractive to a prospective partner, especially given the potential considerable investment likely required prior to commercialization.

Upcoming Milestones

Product	Event	Timeline	
CTP-354	Report full data from ph.I MAD study	Oct 2014 (ANA)	
	Initiate ph.II trial in SCI spasticity patients	year-end 2014	
	Initiate ph.II trial in MS spasticity patients	1H15	_
	Report ph.II spasticity results	1H16	•
CPT-499	SPA discussions with FDA	2H14	
	Sign partnership agreement	2H14+	
AVP-786	Ph.II for treatment-resistant MDD first patient enrollment	Sep-14	
	Ph.II for treatment-resistant MDD completion	year-end 2014	
CTP-730	Ph.I study initiation	2H14	
	Milestone from CELG for phase I study	2015	
JZP-386	Ph.I study initiation	2014	

Source: Company reports and Wells Fargo Securities, LLC estimates

Product Pipeline

Product (partner)	Indication/mechanism	Status
CTP-354	Spasticity, anxiety, pain; subtype selective GABAa receptor modulator	Phase I
CTP-499	Diabetic nephropathy; multi-subtype selective inhibitor of phosphodiesterases	Phase II
AVP-786 (AVNR)	Neurologic and psychiatric disorders, depresion; deuterium- substituted dextromethorphan analog plus low-dose quinidine	Entering phase II
JZP-386 (JZP)	Narcolepsy; deuterium-substituted Xyrem analog	Phase I
CTP-730 (CELG)	Inflammatory diseases	Entering phase I
JZP-386 (JZP)	Narcolepsy; deuterium-substituted Xyrem analog	Phase I
C-10068	Pain and seizures; deuterium-substituted dextromethorphan/analog	Preclinical
d-ivacaftor	CF, COPD	Preclinical
d-praziquantel (NIH)	Parasitic diseases	Preclinical

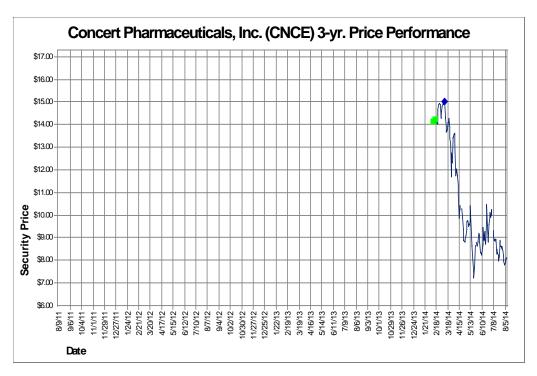
Source: Company reports and Wells Fargo Securities, LLC

Concert Pharma (CNCE) Statement of Operations (Income Statement)

	2012A	2013A	1QA	20A	3QE	4QE	2014E	2015E	2016E	2017E	2018E	2019E	2020E	2021E	2022E
Revenues (1)															
Revenues from CTP-354													\$48,920	\$102,009	\$166,608
Royalties from sales of CTP-499												\$16,192	\$35,289	\$58,517	\$84,650
Royalties from sales of A VP-786														\$10,032	\$23,732
Royalties from sales of JZP-386															\$95,568
Milestones	\$1,500	\$2,000	\$0	\$0	\$0	S	\$	\$14,000	\$30,000	\$15,000	\$34,000	\$65,000	\$0	\$63,000	\$52,000
License and research and development revenue	\$11,349	\$23,408	\$1,613	\$1,235	\$880	\$880	\$4,608	\$8,520	\$12,520	\$12,520	\$12,520	\$9,000	\$9,000	\$9,000	\$9,000
Total revenues, net	\$12,849	\$25,408	\$1,613	\$1,235	\$880	\$880	\$4,608	\$22,520	\$42,520	\$27,520	\$46,520	\$90,192	\$93,210	\$242,559	\$431,558
Expenses															
Cost of goods sold													\$4,892	\$9,669	\$15,536
Research and development	\$24,193	\$21,790	\$5,594	\$6,243	\$6,450	\$6,950	\$25,237	\$30,284	\$31,799	\$33,389	\$34,724	\$35,766	\$36,839	\$37,944	\$39,082
Selling, general and administrative	\$7,266	\$8,028	\$2,538	\$2,718	\$2,800	\$2,900	\$10,956	\$12,380	\$12,875	\$13,391	\$16,069	\$48,206	\$81,950	\$147,510	\$184,387
Total operating expenses	\$31,459	29,818	\$8,132	\$8,961	9,250	\$9,850	\$36,193	\$42,665	\$44,674	\$46,779	\$50,793	\$83,972	\$123,681	\$195,122	\$239,006
Operating Income	(\$18,610)	(\$4,410)	(\$6,519)	(\$7,726)	(\$8,370)	(\$8,970)	(\$31,585)	(\$20,145)	(\$2,154)	(\$19,259)	(\$4,273)	\$6,220	(\$30,471)	\$47,436	\$192,552
Investment income	\$22	\$21	\$0	\$16	\$23	\$21	\$60	\$89	\$113	\$110	\$84	\$73	\$77	\$100	\$202
Interest and other expense	(\$1,856)	(\$1,667)	(\$431)	(\$280)	(\$207)	(\$148)	(\$1,066)	(\$294)	\$0	\$	\$0	\$0	\$0	%	\$0
CTP-499 commercialization payment to GSK											(\$2,750)				
(Loss) income before benefit from income taxes	(\$20,444)	(\$6,056)	(\$6,950)	(066'2\$)	(\$8,554)	(260,6\$)	(\$32,591)	(\$20,349)	(\$2,041)	(\$19,149)	(\$6,939)	\$6,293	(\$30,394)	\$47,536	\$192,754
Benefit (expense) from income taxes	0\$	\$0	\$0	\$0	\$0	O\$	\$0	\$0	0\$	0\$	\$0	(\$126)	\$0	(\$2,852)	(\$15,420)
Net (loss) income	(\$20,444)	(\$6,056)	(\$6,950)	(066'2\$)	(\$8,554)	(260'6\$)	(\$32,591)	(\$20,349)	(\$2,041)	(\$19,149)	(\$6,939)	\$6,167	(\$30,394)	\$44,684	\$177,334
Accretion on redeemable convertible preferred stock	(\$388)	(968\$)	(\$22)		(\$100)	(\$100)	(\$252)	(\$300)	0\$	\$0	\$0	\$0	\$0	0\$	\$0
Net loss applicable to common shareholders	(\$20,832)	(\$6,452)	(\$2,005)	(\$7,990)	(\$8,654)	(\$9,197)	(\$32,846)	(\$20,649)	(\$2,041)	(\$19,149)	(\$6,939)	\$6,167	(\$30,394)	\$44,684	\$177,334
Earnings per share (EPS)	(\$16.15)	(\$4.99)	(\$0.76)	(\$0.45)	(\$0.48)	(\$0.50)	(\$2.07)	(\$1.09)	(\$0.10)	(\$0.95)	(\$0.33)	\$0.26	(\$1.29)	\$1.70	\$6.59
Shares Outstanding (Basic)	1,290	1,292	9,188	17,937	18,137	18,337	15,900	18,937	19,537	20,137	20,737	21,337	23,603	24,203	24,803
Shares Outstanding (Diluted)	1,290	13,237	11,296	20,045	20,245	20,445	18,008	21,045	21,645	22,245	22,845	23,445	25,711	26,311	26,911

Source: Company reports and Wells Fargo Securities, LLC Note: We do not provide estimates of 2015 quarterly EPS at this point, 2014 quarterly EPS does not sum to full year EPS due to share count calculations Note: in 000's & except per share amounts; FY ends 12/31 (1) Not probability weighted

Required Disclosures



	Date	Publication Price (\$)	Rating Code	Val. Rng. Low	Val. Rng. High	Close Price (\$)
	2/13/2014		IPO at \$14.00			•
	3/10/2014		Abrahams, M.D.			
•	3/10/2014	14.88	1	21.00	23.00	15.03

Source: Wells Fargo Securities, LLC estimates and Reuters data

 Symbol Key
 Rating Code Key

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 1 Outperform/Buy
 SR Suspended

 ▲ Rating Upgrade
 ■ Analyst Change
 2 Market Perform/Hold
 NR Not Rated

 ● Valuation Range Change
 □ Split Adjustment
 3 Underperform/Sell
 NE No Estimate

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CNCE: Key risks, in our view, are failure of '354 and/or '499 to show efficacy in subsequent studies and regulatory hurdles in spasticity or CKD.

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As of: August 12, 2014

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