



Charles C. Duncan, PhD

(212) 906-3510

(212) 906-3505

Gena H. Wang, PhD
gwang@jmpsecurities.com

(212) 906-3528

Roy Buchanan, PhD
rbuchanan@jmpsecurities.com

(212) 906-3514

Jason N. Butler, PhD jbutler@jmpsecurities.com

cduncan@jmpsecurities.com

Biotechnology - Company Report

Horizon Pharma, Inc. (1,3)

Diligence Highlights Commercial Progress, Potential Duexis Competitive Advantage

MARKET OUTPERFORM

HZNP \$7.36

Price	\$7.36	FY Dec		2010A	2011E	2012E
Target Price	\$16.00	Revenue (M)	1Q		\$1.8A	\$6.5
52-Wk Range	\$4.53 - \$9.34	rtovonao (m)	2Q		\$1.3A	\$10.5
Shares Out. (M)	19.1		3Q		\$0.5	\$14.8
Market Cap. (M)	\$140.8		4Q		\$1.1	\$18.3
Average Daily Vol. (000)	41		FY	\$4.7	\$4.7	\$50.1
Float (M)	\$6.7			*	*	*****
. 1001 ()	Ψ0			2010A	2011E	2012E
LT Debt (M)	\$10.4	EPS	1Q		(\$5.13)A	(\$0.62)
Cash (M)	\$51.8		2Q		(\$7.78)A	(\$0.40)
Enterprise Value (M)	\$99.4		3Q		(\$0.63)	(\$0.23)
Cash/Share	\$2.71		4Q		(\$0.73)	(\$0.06)
			FY	(\$8.91)	(\$4.40)	(\$1.25)
			P/E	NM	NM	NM
			Previous FY		NC	NC
			CY	(\$8.91)	(\$4.40)	(\$1.25)
			PE	NM	NM	NM

 ${\it NC\ indicates\ no\ change\ to\ previous\ estimate.}\ {\it NE\ indicates\ no\ previous\ estimate.}$

Source: Company reports and JMP Securities

INVESTMENT HIGHLIGHTS

- KOL and management diligence provides enhanced conviction on our outlook for Duexis' imminent launch; reiterate Market Outperform rating and a \$16 price target for Horizon. We recently had an opportunity to meet with Horizon's management and are conducting ongoing diligence with physician thought leaders. As a result, we have incrementally increased visibility on the near-term launch of Duexis in the U.S., expected this quarter. The company has hired the Duexis sales force of 80 reps, and approximately 60 of those reps have been fully trained. We maintain our expectation for a solid launch from Duexis given management's marketing experience and feedback from physicians suggesting the product is likely to be well received based on its clinical value. Our \$16 price target is derived from 4x our estimated U.S. revenues and 7x estimated E.U. royalties for Duexis and Lodotra in 2017 with a discount rate of 30%.
- Duexis' value proposition is multifaceted and backed by strong execution to date. In our view, Duexis is a product offering bringing unique value to the pain management space. The product has much needed safety advantages lacking from the majority of the highly-prescribed NSAID class. While Duexis' clinical advantages would be theoretically achieved by multiple individual pills (up to 16 daily), this prescribing behavior hasn't occurred in actual clinical practice due to a combination of prescribing and patient compliance habits. Duexis provides immediate NSAID relief (unlike competing NSAID/acid-blocker combination offerings) with three pills a day, lowering the burden on the patient's tolerance, memory, and patience.
- Duexis' patent position increasingly robust. In addition, our continued diligence with legal experts supports our view on the increasingly robust Duexis patent estate, and we fully expect any ANDA challenges that may arise to be settled or fail in court. Earlier this month, Horizon announced the allowance of a key U.S. formulation patent for Duexis (Application #12/324808, "Stable Compositions of Famotidine and Ibuprofen", Pub. App. #20090142393), which our diligence suggests should be the more robust Duexis patent and provides protection until 2028. The company also received an allowance for a method-of-use patent in September (Application #11/779,204, "Methods and Medicaments for Administration of Ibuprofen", Pub. App. #20080063706), providing protection until 2026. Tables provided by Horizon that describe the allowed and pending IP for Duexis are shown in Figures 1 and 2.

FOR DISCLOSURE AND FOOTNOTE INFORMATION, REFER TO THE JMP FACTS AND DISCLOSURES SECTION



FIGURE 1: Duexis Patent Estate

Subject Matter	Publication No.	Territory	Key Claims	Patent Term
Admixture	WO07012019	All Major Markets	Dosage Form with 800 mg Ibuprofen / 26.6 mg Famotidine TID Administration Prevention of Ibuprofeninduced Ulcers with Dosage Form	2026
Separate Compartments	WO08011426	All Major Markets	Dosage Form Architecture Barrier Layer Prevents Famotidine Degradation Prevention of Ibuprofeninduced Ulcers with Dosage Form	2026
Unit Dose I	US080021078	US	Dosage Form Architecture Barrier Layer and Ibuprofen Surrounds Famotidine Prevention of Ibuprofeninduced Ulcers with Dosage Form	2026

Source: Company reports

FIGURE 2: Additional Duexis Patents

Subject Matter	Publication No.	Territory	Key Claims	Patent Term
Unit Dose II	US080020040	US	Dosage Form Architecture pH Independent Barrier Layer Prevention of Ibuprofeninduced Ulcers with Dosage Form	2026
Unit Dose III	US080063706 US090264484	US	Dosage Form Architecture Optimal Barrier Layer Prevention of Ibuprofeninduced Ulcers with Dosage Form	2026
Unit Dose IV	US090142393	US	- Dosage Form Architecture - Preventing "Work-arounds"	2028
Commercial Formulation	Not Yet Published	Worldwide	- Commercial Formulation - Additional "Work-arounds"	2031
Phase 3 Study Results	Not Yet Published	Worldwide	Treatment of RA Combination Therapies Patient Subgroups	2031

Source: Company reports



INVESTMENT RISKS

Regulatory risk. The FDA, and/or other ex-U.S. regulatory agencies, could reject any of the firms', or its partners', future regulatory filings or require additional studies prior to granting approval.

Commercial risk. If successfully developed and approved, Horizon's products may face competition both from approved products and also potentially from new product candidates in development by biotechnology and pharmaceutical companies. The company may also face IP risk from competing brand or generic products or product candidates.

Balance sheet risk. The expenses associated with drug development and commercialization are high. Horizon may return to the capital markets to secure additional financing to fund current or future development programs or marketing efforts. Horizon had approximately \$5.7MM in cash and equivalents at the end of 2Q11 and netted approximately \$46MM from its July IPO that we believe will be sufficient to fund operations through 2011. We have projected a raise of ~56MM in 2Q12 at \$15/share. However, the company may also complete one or multiple ex-U.S. partnerships for Duexis which would reduce the need for equity financing.

COMPANY DESCRIPTION

Horizon Pharma is a specialty pharmaceutical company focused on the development and commercialization of novel drug formulations for the treatment of pain and inflammatory indications, particularly arthritis. The company has two approved products, Duexis in the U.S. and Lodotra in Europe. The primary near/mid-term drivers for Horizon are successful execution on the launch of Duexis in the U.S. (planned for 4Q11) as well as U.S. approval of Lodotra (NDA filing expected in 3Q11). Horizon has partnered Lodotra in Europe and Asia, with Mundipharma and Merck/Serono, and intends to secure a partner for Duexis for ex-U.S. geographies.



JMP FACTS AND DISCLOSURES

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Company		Disclosures	
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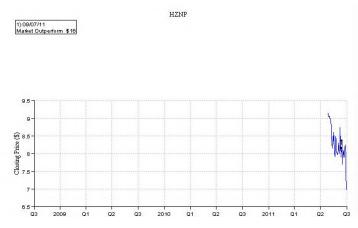
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Market Outperform	Buy	207	66%	Buv	207	66%	58	28%
Market Perform	Hold	105	33%	Hold	105	33%	7	20% 7%
Market Underperform	Sell	3	1%	Sell	3	1%	0	0%
TOTAL:		315	100%		315	100%	65	21%

Stock Price Chart of Rating and Target Price Changes:

Note: First annotation denotes initiation of coverage or 3 years, whichever is shorter. If no target price is listed, then the target price is N/A. In accordance with NASD Rule 2711, the chart(s) below reflect(s) price range and any changes to the rating or price target as of the end of the most recent calendar quarter. The action reflected in this note is not annotated in the stock price chart. Source: Jovus and JMP Securities.



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JMP SECURITIES LLC

600 Montgomery Street, Suite 1100, San Francisco, CA 94111-2713, www.jmpsecurities.com

Peter V. Coleman Director of Equity Research (415) 869-4455

Financial Services Capital Markets David Trone	(212) 906-3525	Real Estate Hotels & Resorts William C. Marks	(415) 835 8044
Steven Fu, CFA	(212) 906-3525	William C. Warks	(415) 835-8944
Chris Ross, CFA	(212) 906-3532	Housing & Housing Supply Chain Michael G. Smith	(415) 835-8965
Consumer & Specialty Finance, Comm John Hecht	<u>ercial Banks</u> (415) 835-3912	Land Development	
Kyle M. Joseph	(415) 835-3940	Michael G. Smith	(415) 835-8965
Financial Processing & Outsourcing		Real Estate & Property Services	
David M. Scharf	(415) 835-8942	William C. Marks	(415) 835-8944
Kevane A. Wong	(415) 835-8976	Real Estate Technology Michael G. Smith	(415) 835-8965
Insurance Matthew J. Carletti	(312) 768-1784	REITs: Healthcare	
Christine Worley	(312) 768-1786	Peter L. Martin, CFA	(415) 835-8904
	,	Aaron Hecht	(415) 835-3963
Market Structure David M. Scharf	(415) 835-8942	REITs: Office & Industrial	
David IVI. Scrian	(413) 633-6942	Mitch Germain	(212) 906-3546
Kevane A. Wong	(415) 835-8976		(= :=) 555 55 :5
Residential & Commercial Real Estate	Finance	Technology	
Steven C. DeLaney	(404) 848-7773	Clean Technology	
Trevor Cranston, CFA	(415) 869-4431	Alex Gauna	(415) 835-8998
Trevor Cranston, CFA	(415) 869-4431	Semiconductors Alex Gauna	(415) 835-8998
Healthcare		Matt Danziger	(415) 869-4454
Biotechnology		Software	
Charles C. Duncan, PhD	(212) 906-3510	Patrick Walravens	(415) 835-8943
Roy Buchanan, PhD	(212) 906-3514	Greg McDowell	(415) 835-3934
Jason N. Butler, PhD	(212) 906-3505	Peter Lowry	(415) 869-4418
Gena H. Wang, PhD	(212) 906-3528	One of Ma Decoral	(445) 005 0004
Liiga A. Dayka	(242) 760 4705	Greg McDowell	(415) 835-3934
Liisa A. Bayko Heather Behanna, PhD	(312) 768-1785 (312) 768-1795		
rication Benatina, i no	(312) 700-1733		
Jason N. Butler, PhD	(212) 906-3505		
Healthcare Facilities & Services			
Peter L. Martin, CFA	(415) 835-8904		
Aaron Hecht	(415) 835-3963		
Healthcare Services			
Constantine Davides, CFA	(617) 235-8502		
Tim McDonough	(617) 235-8504		
Medical Devices J. T. Haresco, III, PhD	(415) 869-4477		

For Additional Information

Mark Lehmann President, JMP Securities (415) 835-3908 Erin Seidemann Vice President, Publishing (415) 835-3970