

Karyopharm Therapeutics Inc. (KPTI)

Reports FY2Q14; Reviews Robust Parallel Registration Paths with Selinexor

MARKET DATA	
Price	\$35.48
52-Week Range:	\$15.50 - \$47.98
Shares Out. (M):	32.6
Market Cap (\$M):	\$1,156.6
Average Daily Vol. (000):	197.0
Cash (M):	\$226
Cash/Share:	\$6.93
Enterprise Value (M):	\$1,041
Float (M):	14.6
LT Debt (M):	\$0
Source: Thomson Reuters and JMP Securities LLC	

FY DEC		2013A	2014E	2015E					
Revenue (\$M)) 1Q		\$0.2A	\$0.0					
	2Q	\$0.4	\$0.0A	\$0.0					
	3Q	\$0.0	\$0.0	\$0.0					
	4Q	\$0.0	\$0.0	\$0.0					
	FY	\$0.0	\$0.0	\$0.0					
EPS	1Q		(\$0.46)A						
	2Q	(\$5.39)	(\$0.55)A						
	3Q	(\$3.66)	(\$0.58)						
	4Q	(\$0.47)	(\$0.63)						
	FY	(\$5.59)	(\$2.20)	(\$4.75)					
Previou	s FY	NC	(\$1.89)	(\$5.18)					
Source: Company	Source: Company reports and JMP Securities LLC								



MARKET OUTPERFORM | Price: \$35.48 | Target Price: \$50.00

INVESTMENT HIGHLIGHTS

Heightened visibility provided on path forward in DLBCL amid a flurry of trial development activity; reiterate our Market Outperform rating and \$50 price target based on our DCF, CAGR, and SOTP methodologies. KPTI recorded net operating loss and EPS of \$16.4MM and (\$0.55), respectively, higher than JMP estimates of \$13MM and (\$0.44), primarily on higher R&D spend. We remind investors that as a development stage biotech company, KPTI continues to be a story of clinical execution with its lead asset selinexor, rather than an earnings story. Key highlights from the conference call included protocol updates related to the planned registration-directed Phase II trial of selinexor in R/R DLBCL, increasing the likelihood of RRMM serving as a fourth registration-directed indication in the near term (in addition to AML, Richter's syndrome, and DLBCL). On the heels of a recently completed secondary financing that raised net proceeds of ~\$113MM, KPTI is adequately resourced to advance selinexor to inflection points on multiple fronts, in our view, both in heme malignancy and solid tumors, wherein we maintain a high level of confidence in its clinical success. Changes to our model reflect the impact of 2Q14 actual results and moderately increased operating expense estimates, with minimal impact to our valuation.

Feedback from FDA supports accelerated path forward in R/R DLBCL on updated Phase II protocol design. As noted on the call and in this morning's press release, the SADAL protocol has been modified to a two-arm, 1:1 randomized Phase IIb trial, evaluating high-dose (flat dose of 100mg = 60mg/2) and low-dose (50mg = 35mg/m2) selinexor plus dexamethasone in 200 patients with ≥3L disease, maintaining ORR and duration of response as the basis for potential accelerated approval. Based on the relatively greater unmet need, the trial will target at least fifty percent enrollment of patients with GCB subtype disease. Based on positive feedback from EMA indicating the potential for registration using the modified protocol, SADAL will recruit both in the U.S. and EU beginning in 4Q14. Confirmatory studies in DLBCL could take the shape of randomized trials in earlier line patients, evaluating standard of care (Rituximab chemoimmunotherapy), with and without selinexor. A Phase I trial of selinexor plus Rituximab in aggressive NHL is currently underway.

Encouraging myeloma activity in combination low-dose dex is the key study to watch in 2H14. Recall that in Phase I data presented at EHA, selinexor in combination with low-dose dexamethasone showed a surprising 50% response rate in eight heavily pretreated myeloma patients (clinical benefit rate of 75%), vastly improving over single-digit response rates observed as a single agent. A majority of responding patients reportedly remain on study, while two target patients have been enrolled onto the combination expansion cohort. Abstracts submitted to EMSO could offer the opportunity for increased visibility on duration of response, although we suspect that it won't be until

Michael G. King, Jr. mking@jmpsecurities.com (212) 906-3520 Eric Joseph, PhD ejoseph@jmpsecurities.com (212) 906-3514



ASH that we learn whether the combination is more broadly active. Assuming the data support an accelerated path forward for selinexor/dex in later-line MM, forthcoming ISTs evaluating selinexor plus standard myeloma therapies (e.g., Kyprolis) should confer solid positioning with respect to confirmatory trial design.

Greater clarity on solid tumor development by year end. Phase I evaluation of selinexor continues in the encouraging indications including prostate cancer, ovarian cancer, and sarcoma. KPTI reiterated plans to select one or two of these indications for further clinical development by year end.

We remain encouraged by the signs of selinexor activity across a wide range of tumor types, both solid and liquid, exemplified by the data presented at ASCO and EHA. We believe Karyopharm is on the verge of bringing an entirely new class of chemotherapy agent to the market with broad activity and acceptable tolerability. We remind the reader that Karyopharm holds the worldwide rights to selinexor.

FIGURE 1. Selinexor Clinical Trials

Trial No.	Sponsor	Phase	Indication	Combo Partner	Pt Size	FPI
NCT01607892	KPTI	1	Various Heme Malignancies (MAD)		250	May-12
NCT01607905	KPTI	I	Various advance solid tumors		90	May-12
NCT02146833	KPTI	II	Metastatic prostate cancer		50	May-14
TBD	KPTI	II	SADAL - ≥3L R/R DLBLC, low and hi dose Selines	cor	200	4Q14
NCT02088541	KPTI	II	SOPRA - R/R Elderly AML vs physician's choice		150	Apr-14
NCT02138786	KPTI	II	SIRRT - R/R Richter's Transformation		50	4Q14
NCT02025985	KPTI	II	SIGN - Gynaecologic malignancies (ovarian, endometrial, cervical)		63	Apr-14
NCT01986348	KPTI	II	KING - Glioblastoma		30	Mar-14
NCT02178436	KPTI	I/II	Pancreatic cancer and PDAC	Gem/Abraxane	43	Not yet recruiting
NCT01896505	KPTI	1	Food effect study		20	Sep-13
NCT02186834	Moffit	I/II	Multiple myeloma	Dexamethasone, Doxil	47	Not yet recruiting
NCT02199665	U. Chicago, NCI	I	Refractory Multiple Myeloma	Kyprolis, Dexamethasone	48	Not yet recruiting
NCT02093403	Ohio State	1	R/R and Elderly Untreated AML	Dacogen	42	Mar-14
NCT02120222	Ohio State	1	Recurrent melanoma		20	Not yet recruiting
NCT02137356	Sheba Med Ctr	1	Neoadjuvant rectal neoplasms	Chemoradiation	28	Not yet recruiting
NCT02069730	U of T		Salivary gland cancers		30	Not yet recruiting
NCT02091245	Dana Farber	1	Childhood relapsed ALL/AML		28	Apr-14
NCT02078349	Ntl Univ. Hosp, Singapore	I	Asian solid tumor study		30	Mar-14

Source: Company filings

August 8, 2014 2



FIGURE 2. Upcoming Catalysts

Timing	Drug	Catalyst
ESMO	Selinexor	Updated Phase I presentations in solid tumor
4Q14	Selinexor	Initiation of second pivotal Phase II/III study in (3L+ DLBCL; SADAL)
4Q14	Selinexor	Initiation of Phase II in Richter's syndrome (SIRRT)
ASH	Selinexor	Updated RRMM Phase I data in combination with dexamethasone
2H14	KPT-350	IND completion for use in inflammation, auto-immune, and anti-viral indications
2H14	PAK Inhibitor	IND completion for use in oncology indications
Source: Com	pany presentations	

REVIEW OF 2Q14 FINANCIALS AND CHANGES TO OUR MODEL

As noted above, KPTI recorded a 2Q14 net operating loss of \$16.4MM, greater than our estimate of \$13MM. Specifically, R&D spend of \$13.2MM was higher than our \$10.2MM estimate, as was G&A spend of \$3.3MM compared to our \$2.8MM estimate. Changes to our FY14 estimates reflect the impact of 2Q14 results. Forward-looking EPS estimates are primarily impacted by increased outstanding share count estimates, resulting from the recent secondary offering of 2.6MM shares. A comparison of 2Q14 results versus JMP and consensus estimates is detailed in Figure 3. Changes to our model are detailed in Figure 4.

FIGURE 3. 2Q14 Actuals versus JMP and Consensus Estimates

Karyopharm Therapeutics (KPTI)	2Q14 Results									
Abridged Income Statement (\$ MM)	JMP Estimate	Street Consensus	Actual	Variance (JMP vs. Actual)						
Total Revenues	-	0.15	0.02	0.02						
Operating Expenses	12.95	14.80	16.47	3.5						
Research and development	10.20		13.16	3.0						
General and administrative	2.75		3.31	0.6						
Operating income (loss)	(12.95)	(14.65)	(16.45)	3.5						
Other income (expense)	0.00	1.65	0.01	0.01						
Interest income	0.00		0.01	0.01						
Pretax income (loss)	(12.95)	(13.00)	(16.44)	(3.49)						
Provision for Income Tax	0.00	(3.30)	0.00	_						
Net income (loss)	(12.95)	(16.30)	(16.44)	(3.49)						
EPS Calculations										
Basic EPS	\$ (0.44)									
Diluted EPS	\$ (0.44)	\$ (0.51)	\$ (0.55)	\$ (0.12)						
Desir shares systematics	00.040		00.050	0.047						
Basic shares outstanding	29.612		29.659	0.047						
Diluted shares outstanding	29.612		29.659	0.047						

Source: JMP Securities LLC, Thomson Reuters, Company filings



FIGURE 4. Changes to Our Income Statement

Karyopharm Therapeutics (KF	3Q ⁻	14E	4Q ²	14E	FY 2	014E	FY 2	015E	FY 2	016E	FY 2	017E
(\$ MM)	Old	New	Old	New	Old	New	Old	New	Old	New	Old	New
Sales ROW Royalties Other	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	44.1	44.1	103.6 6.8	103.6 6.8
Total Revenue	-	-	-	-	-	-	-	-	44.1	44.1	110.4	110.4
cogs	0.0	0.0	0.0	0.0	0.2	0.2	0.0	0.0	4.4	4.4	9.3	9.3
Gross Profit	-	-	-	-	0.17	0.19	-	-	39.7	39.7	101.1	101.1
Operating Expenses Research and development General and administrative Operating income (loss)	13.5 10.6 2.9 (13.5)	18.1 14.5 3.6 (18.1)	15.7 11.7 4.0 (15.7)	20.5 16.5 4.0 (20.5)	56.0 43.5 12.6 (55.9)	69.0 55.1 13.8 (68.8)	153.6 78.3 75.3 (153.6)	153.6 96.5 57.1 (153.6)	234.4 125.2 109.2 (194.7)	232.4 149.6 82.8 (192.7)	302.8 169.0 133.8 (201.8)	303.4 201.9 101.5 (202.3)
Other income (expense)	-	-	-	-	-	-	-	-	-	-	-	-
Interest income Interest expense	-	-	-	-	-	-	-	-	-	-	-	-
Pretax income Provision for Income Tax	(13.5) -	(18.1) -	(15.7) -	(20.5)	(55.9)	(68.8)	(153.6)	(153.6) -	(194.7) -	(192.7) -	(201.8)	(202.3)
Net income	(13.5)	(18.1)	(15.7)	(20.5)	(55.9)	(68.8)	(153.6)	(153.6)	(194.7)	(192.7)	(201.8)	(202.3)
Basic EPS Diluted EPS	\$ (0.46) \$ (0.46)	, ()	\$ (0.53) \$ (0.53)	\$ (0.63) \$ (0.63)	,						,	\$ (5.61) \$ (5.61)
Basic shares outstanding Diluted shares outstanding	29.62 29.62	30.99 30.99	29.63 29.63	32.32 32.32	29.52 29.52	31.20 31.20	29.67 29.67	32.36 32.36	31.50 31.50	34.19 34.19	33.35 33.35	36.04 36.04

Source: JMP Securities LLC, Company filings

FIGURE 5. Updated Income Statement

Income Statement (\$MM)	1Q	14A :	2Q14A	3Q14E	4Q14E	2014E	2015E	2016E	2017E	2018E	2019E	2020E
Product Sales and Royalties:												
Selinexor												
US Sales							-	44.1	103.6	308.1	842.2	1,463.8
ROW Royalties							-	-	6.8	17.1	43.2	111.9
Total Product Sales and Royalties		0.0	0.0	0.0	0.0	0.0	0.0	44.1	110.4	325.2	885.3	1,575.8
Collaboration Revenue		0.0	0.0									
Total Revenue		0.2	0.0	0.0	0.0	0.0	0.0	44.4	440.4	205.0	005.0	4 575 0
Total Revenue		0.2	0.0	0.0	0.0	0.0	0.0	44.1	110.4	325.2	885.3	1,575.8
Cost of Goods Sold								4.4	9.3	24.6	67.4	117.1
Gross Profit		0.2	0.0	0.0	0.0	0.2	0.0	39.7	101.1	300.5	818.0	1,458.7
Operating Expenses:												
Research and Development		11.0	13.2	14.5	16.5	55.1	96.5	149.6	201.9	226.1	244.2	256.4
General and administrative		2.9	3.3	3.6	4.0	13.8	57.1	82.8	101.5	116.7	128.3	141.2
Total operating expenses		13.9	16.5	18.1	20.5	69.0	153.6	232.4	303.4	342.8	372.6	397.6
Operating income (loss)		(13.7)	(16.4)	(18.1)	(20.5)	(68.8)	(153.6)	(192.7)	(202.3)	(42.3)	445.4	1,061.0
operating mooms (1965)		(10.1)	(10.1)	(10.1)	(20.0)	(00.0)	(100.0)	(102.1)	(202.0)	(12.0)	110.1	1,001.0
Other income (expense):												
Interest income		0.0	0.0									
Interest expense												
Total other income, net		0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Change in fair value of preferred stock warrant liability												
Foreign currency transaction gain (loss)												
Pretax income (loss)		(13.7)	(16.4)	(18.1)	(20.5)	(68.7)	(153.6)	(192.7)	(202.3)	(42.3)	445.4	1,061.0
Income tax benefit (provision)						0.0	0.0	0.0	0.0	0.0	0.0	(159.2)
Tax Rate						0%	0%	0%	0%	0%	0%	15%
Comprehensive income (loss)		(13.7)	(16.4)	(18.1)	(20.5)	(68.7)	(153.6)	(192.7)	(202.3)	(42.3)	445.4	901.9
Accretion of redeemable convertible preferred stock		(12.7)	(16.4)	(40.4)	(20.5)	(60.7)	(452.0)	(100.7)	(202.2)	(40.0)	11E 1	001.0
Net income (loss) attributable to common stockholders		(13.7)	(16.4)	(18.1)	(20.5)	(68.7)	(153.6)	(192.7)	(202.3)	(42.3)	445.4	901.9
Basic EPS to common shareholders	\$	(0.46) \$	/	\$ (0.58)								\$ 24.81
Diluted EPS to common shareholders	\$	(0.46) \$	(0.55)	\$ (0.58)	\$ (0.63)	\$ (2.20)	\$ (4.75)	\$ (5.64)	\$ (5.61)	\$ (1.17)	\$ 11.97	\$ 24.18
Basic shares outstanding		29.6	29.7	31.0	32.3	31.2	32.4	34.2	36.0	36.1	36.3	36.4
Diluted shares outstanding		29.6	29.7	31.0	32.3	31.2	32.4	34.2	36.0	36.1	37.2	37.3
% change in diluted shares outstanding		20.0	20.1	01.0	02.0	414.3%	3.7%	5.7%	5.4%	0.3%	2.9%	0.3%
70 Grange III dilated shares batstanding						T17.3/0	3.1 /0	J.1 /0	J.4/0	0.5/0	2.3/0	0.370

Source: JMP Securities LLC, Company filings



Company Description

Karyopharm Therapeutics (KPTI) is a Natick, MA-based, clinical-stage biopharmaceutical company focused on the discovery and development of novel first-in-class drugs directed against nuclear transport targets for the treatment of cancer and other major diseases. Karyopharm's Selective Inhibitors of Nuclear Export (SINE) compounds function by preventing the export of tumor suppressor proteins from the nucleus, driving accumulation and restoration of function. The company's lead pipeline candidate, selinexor (KPT-330), is a Phase I orally available small molecule inhibitor of XPO1, set to initiate pivotal Phase II/III evaluation in various hematologic malignancies in 2014. Karyopharm is also developing selinexor and SINE as potential therapies for autoimmune and inflammatory disease, viral infections, and wound healing.

Investment Risks

Clinical. Drug development is an inherently risky business. Clinical trials always carry a risk of failure and Karyopharm's assets (Selinexor (KPT330), KPT-350, PAK4 inhibitor, verdinexor or future drug candidates) may fail to demonstrate meaningful enough levels of efficacy in current or future clinical trials.

Regulatory and commercial. The ability of Karyopharm to market its drugs depends upon the drugs obtaining approval from the FDA and foreign regulatory agencies. Failure to achieve approval or delays in the timelines to approval could negatively impact the company's share price.

Competitive. Hematologic malignancies, including multiple myeloma, indolent non-Hodgkin lymphoma, and acute myeloid leukemia represent increasingly competitive fields and Karyopharm faces competition from both commercial and development-stage companies with product(s) or product candidates addressing similar clinical indications. Some of these companies may possess substantially greater R&D and commercial resources than Karyopharm. As such, there is no assurance Karyopharm will be competitive or differentiated from other drug products.

Financial. Following its IPO, Karyopharm ended 1Q14 with approximately \$156MM in cash and cash equivalents. We anticipate the company is likely to seek additional equity financing in the form of a secondary offering in order to complete the development of its drug candidates, creating dilution risk for existing shareholders.



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JMP Securities was manager or co-manager of a public offering of securities for Karyopharm Therapeutics Inc. (KPTI) in the past 12 months, and received compensation for doing so.

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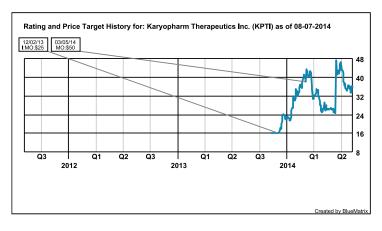
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							# Co's	
							Receiving	
							IB	
		# Co's	%		# Co's	%	Services in	% of Co's
	Regulatory	Under	of	Regulatory	Under	of	Past 12	With This
JMP Rating	Equivalent	Coverage	Total	Equivalent	Coverage	Total	Months	Rating
MARKET OUTPERFORM	Buy	267	60.14%	Buy	267	60.14%	97	36.33%
MARKET PERFORM	Hold	137	30.86%	Hold	137	30.86%	18	13.14%
MARKET UNDERPERFORM	Sell	4	0.90%	Sell	4	0.90%	0	0%
COVERAGE IN TRANSITION		36	8.11%		36	8.11%	0	0%
TOTAL:		444	100%		444	100%	115	25.90%

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 guarter. The action reflected in this note is not annotated in the stock price chart. Source: JMP Securities.



Karyopharm Therapeutics Inc. (KPTI)



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Jeffrey H. Spurr Director of Research (415) 835-3903

RESEARCH PROFESSIONALS

FINANCIAL SERVICES

Alternative Asset Managers		Medical Devices & Supplies	
Devin Ryan	(212) 906-3578	David Turkaly	(212) 906-3563
Brian McKenna	(212) 906-3545	John Gillings	(212) 906-3564
		Ou a state a Discours a south a sta	
Commercial & Specialty Finance		Specialty Pharmaceuticals	(0.40) 000 0=00
Christopher York	(415) 835-8965	Oren G. Livnat, CFA	(212) 906-3566
Hannah Kim, CFA	(415) 835-8962	Nazibur Rahman	(212) 906-3519
Consumer Finance			
David M. Scharf	(415) 835-8942	REAL ESTATE	
Jeremy Frazer	(312) 768-1796		
Jeremy i razer	(312) 700-1790	Housing & Land Development	
Financial Processing & Outsourcing		Peter L. Martin, CFA	(415) 835-8904
David M. Scharf	(445) 925 9042	Aaron Hecht	(415) 835-3963
	(415) 835-8942	Bharathwajan Iyengar	(415) 835-3902
Jeremy Frazer	(312) 768-1796		()
Insurance		Lodging & Leisure	
Matthew J. Carletti	(312) 768-1784	Robert A. LaFleur	(212) 906-3510
		Whitney Stevenson	(212) 906-3538
Christine Worley	(312) 768-1786		(= :=) ::::
Investment Banks & Brokers		Property Services	
	(242) 000 2570	Mitch Germain	(212) 906-3546
Devin Ryan	(212) 906-3578	Peter Lunenburg	(212) 906-3537
Brian McKenna	(212) 906-3545	. 5.5	(= :=) 000 000:
Mortgage Operating Companies		REITs: Healthcare, Residential, & Specia	lty
		Peter L. Martin, CFA	(415) 835-8904
REITs: Agency, Hybrid, & Commercial M		Aaron Hecht	(415) 835-3963
Steven C. DeLaney	(404) 848-7773	Arthur Kwok	(415) 835-8908
Trevor Cranston, CFA	(415) 869-4431	Addid Note	(110) 000 0000
Charter Robinson	(757) 613-8955	REITs: Office, Industrial, & Diversified	
Benjamin Zucker	(212) 906-3529	Mitch Germain	(212) 906-3546
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HEALTHCARE		reter Luneriburg	(212) 906-3537
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Biotechnology	(040) 700 4705		(415) 835-3963
Liisa A. Bayko	(312) 768-1785	Aaron Hecht	
Andrew Prigodich, PhD	(312) 768-1788	Bharathwajan Iyengar	(415) 835-3902
Bhumika Sharma, PhD	(312) 768-1795		
Jason N. Butler, PhD	(212) 906-3505	TECHNOLOGY	
Caroline Palomeque	(212) 906-3509	TEOTINOEOGT	
Michael G. King, Jr.	(212) 906-3520	Communications Equipment & Internet S	Cocurity
Bryan Czyzewski, PhD	(212) 906-3577	Erik Suppiger	
Eric Joseph, PhD	(212) 906-3514	11 0	(415) 835-3918
• •	,	John Lucia	(415) 835-3920
Healthcare Services & Facilities		Internet & Digital Media	
Peter L. Martin, CFA	(415) 835-8904	Ronald V. Josey III	(242) 006 2520
Aaron Hecht	(415) 835-3963		(212) 906-3528
Arthur Kwok	(415) 835-8908	Andrew Boone, CFA	(415) 835-3957
	()-/	Michael Wu	(415) 835-8996
Life Science Tools & Diagnostics		Software	
J. T. Haresco, III, PhD	(415) 869-4477		(445) 025 0042
Marie T. Casey, PhD	(415) 835-3955	Patrick Walravens	(415) 835-8943
	(-,	Peter Lowry	(415) 869-4418
Medical Devices		Greg McDowell	(415) 835-3934
J. T. Haresco, III, PhD	(415) 869-4477	Minelana 9 Claud Committee Tool	
Marie T. Casey, PhD	(415) 835-3955	Wireless & Cloud Computing Technolog	
	(-,	Alex Gauna	(415) 835-8998

ADDITIONAL CONTACTS

Thomas R. Wright Director of Equities (212) 906-3599 Dan Wychulis Director of Institutional Sales (617) 235-8530 **600 Montgomery Street, Suite 1100** San Francisco, CA 94111 www.jmpsecurities.com