#### **US Equity Research**

18 June 2015

#### BUY

unchanged

PRICE TARGET US\$70.00 unchanged Price (18-Jun) US\$60.10

Ticker RDUS-NASDAQ

52-Week Range (US\$):
Avg Daily Vol (M):
Shares Out. (M):
Market Cap (US\$M):

FYE Dec	2014A	2015E	2016E
Revenue (US\$M)	0.0	0.0	82.9
EPS Adj&Dil (US\$)	(53.81)	(2.29)↑	(1.10)↑
Previous	(53.81)	(2.31)	(1.19)

7.46 - 51.22

277.6

29.7

1,072

Quarterly Revenue	Q1	Q2	Q3	Q4
2014A	0.0	0.0	0.0	0.0
2015E	0.0A	0.0	0.0	0.0
2016E	-	-	-	-

Quarterly EPS Adj&Dil	Q1	Q2	Q3	Q4
2014A	(50.45)	(2.22)	(0.59)	(0.55)
2015E	(0.47)A	(0.66)	(0.58)	(0.58)
2016E				



Radius is a biotechnology company focused on drugs for endocrine disorders, including osteoporosis.

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# **Estimates Revised**

# ACTIVExtend results positive for abaloparatide, RAD109 holds additional value

#### Abaloparatide extension study maintains strong Ph 3 results

Alendronate maintained the strong 86% reduction in new vertebral fractures vs. placebo (p<0.0001) in the 6-month alendronate extension period, which is very encouraging, in our view. Also, the arm receiving 18 months of abaloaparatide followed by 6 months of alendronate showed a 0.57 hazard ratio versus placebo for non-vertebral fractures vs. 0.72 for the teriparatide arm, which was not statistically significant vs. placebo.

## Wrist fracture data may open orthopedic opportunity

We are particularly interested in the 72% reduction in wrist fractures for abaloparatide followed by alendronate, as this may open an opportunity for the drug in the orthopedic market. Interestingly, there are ~440,000 wrist fractures per year in women, many of which are untreated, with only 20-25% of women tested or treated for osteoporosis following a fracture.

#### Expect YE15 FDA filing, launch in 2016

Radius expects to file the NDA for abaloparatide for the treatment of severe osteoporosis by YE15, and we expect approval in 2016. We maintain our \$560M US peak sales estimate, which assumes Radius captures ~50% of current market share from Forteo.

#### RAD-1901 holds upside, data expected YE15

We await initial data in breast cancer for RAD-1901 by YE15, likely at the San Antonio Breast cancer symposium. Based on activity for Seragon's drug with a similar mechanism of action, but better safety for RAD-1901, we could see exciting clinical activity. As of ASCO 2015, only a few patients had been enrolled in the RAD-1901 study, so no efficacy data were available. We currently model ~\$470M US peak sales and apply a 35% probability adjustment to arrive at \$10 per share US, and similar metrics to arrive at \$9 per share Ex-US. We have updated the estimated share count for 2015 and 2016, resulting in slightly higher EPS estimates of (\$2.29) for 2015 vs. (\$2.31) previously, and (\$1.10) for 2016 vs. (\$1.19) previously.

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Figure 1: RDUS valuation

Product	Peak Sales (\$MM)	Year	NPV at launch	Estimated launch	Time to launch	Probability Adjustment	Current Value (\$MM)	Scenario probability	Value / Share (EV/Sales)
abaloparatide									
US	\$358	2022	\$561	6/1/2016	1.0	85%	\$388	100%	\$21
Ex-US - co-promote	\$346	2021	\$390	1/1/2017	1.5	85%	\$254	50%	\$11
Ex-US - royalty	\$346	2021	\$189	1/1/2017	1.5	85%	\$135	50%	\$11
Total abaloparatide							\$641		\$44
RAD-1901									
US	\$467	2023	\$670			35%	\$235		\$10
Ex-US	\$427	2023	\$188			35%	\$66		\$9
Total RAD-1901							\$300		\$19
Total Product Value							\$641		\$64
Cash							243		\$7
Total Equity Value							884		\$70
Shares Outstanding (MM)							36		

Risk-Free Rate	2.0%
Beta	1.8
Risk Premium	6%
Discount Rate	12%
EV/Sales	5.5

Source: Company Reports, Canaccord Genuity estimates



Figure 2: Radius income statement

			Rad	ius He	alth, In	C.						
(000's) [FY - DEC]												
Revenues	2013A	2014A	1Q15A	2Q15E	3Q15E	4Q15E	2015E	2016E	2017E	2018E	2019E	2020E
abaloparatide - US								82,902	122,229	165,478	212,938	264,914
abaloparatide - Ex-US								-	90,548	204,751	251,503	298,717
Total								82,902	212,777	370,229	464,441	563,631
Income Statement	2013A	2014A	1Q15A	2Q15E	3Q15E	4Q15E	2015E	2016E	2017E	2018E	2019E	2020E
Total Revenue	-	-	-	-	-	-	-	82,902	146,677	265,806	341,204	420,247
COGS	-	-	-	-	-	-	-	16,580	29,335	53,161	68,241	84,049
Gross Profit	-	-	•	•	•	-	-	66,322	117,341	212,645	272,963	336,197
Operating Expenses												
Research and development	60,536	45,719	11,559	16,270	15,173	13,525	56,527	47,591	47,070	57,626	77,561	111,588
General and administrative	6,829	13,674	4,756	4,956	5,156	5,356	20,224	58,031	47,670	59,806	76,771	94,556
Total Operating Expense	67,365	59,393	16,315	21,226	20,329	18,881	76,751	105,622	94,740	117,432	154,332	206,143
EBITDA												
Operating income	(67,365)	(59,393)	(16,315)	(21,226)	(20,329)	(18,881)	(76,751)	(39,301)	22,602	95,212	118,631	130,054
Other income (expense), net	9,085	(2,126)	(50)	(2,126)	(50)	(2,126)	(4,352)	(2,126)	(4,352)	(2,126)	(4,352)	(2,126)
Loss on retirement of note payable	0,000	(2,120)	(30)	(2,120)	(50)	(2,120)	(4,552)	(2,120)	(4,002)	(2,120)	(4,002)	(2,120)
Interest (expense) income, net	(2,410)	(768)	(692)	(768)	(692)	(768)	(2,920)	(768)	(2,920)	(768)	(2,920)	(768)
Accretion of preferred stock	( )	` ′	, ,	` ,	,	` ,	, ,	, ,	( , ,	,	( , ,	, ,
Pre-tax income (GAAP)	(60,690)	(71,501)	(17,057)	(24,120)	(21,071)	(21,775)	(84,023)	(42,195)	15,330	92,318	111,359	127,160
Pre-tax income (non-GAAP)												
Taylor (CAAD)									5,672	24.450	41,203	47.040
Taxes (GAAP)	37%		37%	37%	37%	270/	270/	37%		34,158		47,049
Tax rate (GAAP)	3176	37%	31%	31%	31%	37%	37%	3176	37%	37%	37%	37%
Net Income (GAAP)	(60,690)	(71,501)	(17,057)	(24,120)	(21,071)	(21,775)	(84,023)	(42,195)	9,658	58,161	70,156	80,111
GAAP EPS (diluted)	(\$3.97)	(\$53.81)	(\$0.47)	(\$0.66)	(\$0.58)	(\$0.58)	(\$2.29)	(\$1.10)	\$0.25	\$1.46	\$1.72	\$1.93
Basic shares outstanding	15,278	12,121	31,000	31,031	31,062	33,062	31,539	35,562	37,340	39,207	41,167	43,226
Diluted shares outstanding	15,278	17,578	36,269	36,305	36,342	37,541	36,614	38,346	39,113	39,896	40,694	41,507

Source: Company reports, Canaccord Genuity estimates



# Appendix: Important Disclosures

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#### **Target Price / Valuation Methodology:**

Radius Health - RDUS

Our \$70 price target is based on an adjusted EV/S methodologies.

#### Risks to achieving Target Price / Valuation:

Radius Health - RDUS

Risks to our outlook and price target include the following: the Phase 3 study for abaloparatide in osteoporosis may be negative, or fail to meet investor expectations, resulting in downside to shares and our price target. Also, Phase 3 data may be positive in terms of efficacy, but show an unexpected safety signal, also resulting in downside to our price target. Antibody formation was been seen in Phase 2 studies, with one patient showing potential evidence of neutralizing antibodies. Even assuming positive Phase 3 data for subcutaneous abaloparatide in osteoporosis, FDA approval may be delayed or may not occur at all, also resulting in downside to shares and our price target. FDA may also grant approval, but require large, lengthy and expensive post-approval studies, which could also result in downside to shares and our price target. Clinical data from other osteoporosis products including anti-sclerostin antibodies from Amgen. Merck. Eli Lilly and Novartis could be viewed as superior to abaloparatide, pressuring shares. Competition from existing and new osteoporosis products could also result in lower revenues that expected, leading to downside to our estimates and the share price. Although unlikely, a paragraph 4 challenge could be filed against Lilly's Forteo, a molecule closely related to abaloparatide, which investors may interpret as increasing risk for abaloparatide, and pressuring Radius shares. Forteo was approved as an NDA, where the ANDA pathway is well established. Even though Forteo is essentially a biologic, since it is a peptide, it is feasible although unlikely that a generic challenger could emerge. FDA has approved a generic version of Lovenox, a biologic approved via the NDA pathway, although the process took many years. If a generic version of Forteo were to reach the market, usage of abaloparatide could decline, resulting in downside to our estimates and price target. Also, if FDA were to approve a generic version of Copaxone, a peptide used to treat multiple sclerosis, investors may see increased risk of a generic challenge and approval for abaloparatide, as both products are classified as NDA filings for biologic peptides. A transdermal microneedle formulation for abaloparatide may not be feasible, which investors may view as negative for life cycle management and commercial competitive positioning for Radius, pressuring shares. Even if a microneedle formulation can be developed to show equal efficacy to the subcutaneous formulation. FDA may require a full clinical study versus a bridging study, which would require additional funding and time to approval.

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Rating	Coverag	Coverage Universe				
	#	%	%			
Buy	588	58.92%	32.99%			
Hold	322	32.26%	15.53%			
Sell	40	4.01%	5.00%			
Speculative Buy	48	4.81%	54.17%			
	998*	100.0%				

<sup>\*</sup>Total includes stocks that are Under Review

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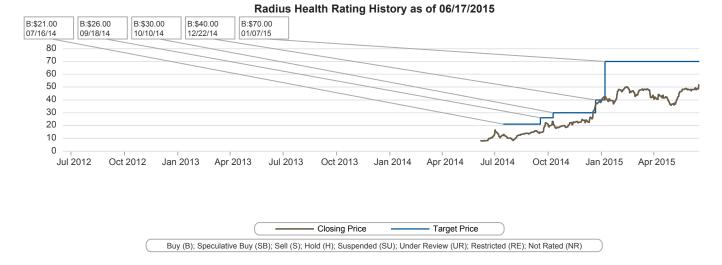
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