

Neothetics Inc.

NEOT - BUY - When Less Is More; Stock Looks Inexpensive Ahead of LIPO-202 Data in December

November 2, 2015

Conclusion: We remain positive on NEOT ahead of 3Q15 earnings results and what we believe will be positive data for LIPO-202 (for non-surgical abdominal fat reduction). Therefore, we think NEOT's LIPO-202 product is underappreciated for the following reasons: 1) We continue to think LIPO-202 could be the next market-leading botulinum toxin, with peak sales potential of over \$1B (expected launch in '18), but NEOT's market cap suggests that the Street is underestimating this opportunity; 2) We believe investing in NEOT could provide a good entry point into the body contouring market early in its growth and development; 3) We believe there is a free call option on LIPO-102, which has orphan designation for the treatment of exophthalmos. NEOT's valuation ascribes little value for this pipeline product, in our view. Similar companies have been valued at \$900MM to \$2B. NEOT has a market cap of only \$109MM, (EV \$46MM).

- **Expecting positive Phase III results for LIPO-202 in December.** Based on when the last patient was enrolled (nine-week trial, four to six weeks for analysis), we expect LIPO-202 data in December. LIPO-202 Phase II data were positive and the Phase III trial design is similar to the Phase II study.
- **Kybella approval presents positive read-through to LIPO-202.** Kybella is a product approved for the reduction of chin fat. Historically, the FDA's Dermatology Committee has been comfortable approving cosmetic drugs, but it had not reviewed such an application for many years until Kybella. Positive read-throughs from Kybella to NEOT include: 1) FDA was comfortable with Kybella's MRI measurements for fat (prior issues resolved); and 2) Kybella used co-endpoints (1 & 2 grade changes, NEOT also has co-endpoints).
- **Next potential catalyst is positive Phase III results for LIPO-202.** Other potential catalysts include: 1) LIPO-202 submission in 2H16; 2) LIPO-202 approval and launch in 2H17.

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NEOT BUY COMPANY UPDATE

Investment Thesis: Growth

SHARE PRICE	\$7.89
PRICE TARGET	\$16.00

Revenue (\$M) (FY Dec)	1Q	2Q	3Q	4Q	FY
2014	0.0	0.0	0.0	0.0	0.0
2015	0.0	0.0	0.0E	0.0E	0.0E
2016	—	—	—	—	0.0E

Adjusted EPS (\$)	1Q	2Q	3Q	4Q	FY
2014	(0.49)	(0.68)	(1.22)	(0.45)	(2.83)
P/E					NM
2015	(0.50)	(0.69)	(1.80)E	(1.75)E	(4.75)E
P/E					NM
2016	—	—	—	—	(4.79)E
P/E					NM

Market Data & Valuation Multiples

52-Week Range	\$5.92 - \$15.05
Shares Out (M)	13.7
Market Cap (M)	\$108
ADV (3 mo; 000)	83

SCENARIO	VALUATION	INVESTMENT THESIS
BULL CASE	2020E EPS: \$6.75 12-Month Bull-Case Scenario: \$28.00 Upside/Downside Potential: 255%	<ul style="list-style-type: none"> • LIPO-202 sales of \$515MM in 2020 • Additional business development • Additional pipeline advancements
BASE CASE	2020E EPS: \$5.63 12-Month Base-Case Scenario: \$16.00 Upside/Downside Potential: 103%	<ul style="list-style-type: none"> • LIPO-202 sales of \$472MM in 2020 • No additional business development • No additional pipeline advancements
BEAR CASE	2020E EPS: \$4.75 12-Month Bear-Case Scenario: \$7.00 Upside/Downside Potential: -11%	<ul style="list-style-type: none"> • LIPO-202 sales of \$410MM in 2020 • No additional business development • No additional pipeline advancements

VALUATION METHODOLOGY AND RISKS

We use a DCF analysis to get to our 12-month price target of \$16. We use a WACC of 28% and an exit multiple of 2x forward EBIT to arrive at our price target. This compares favorably to NEOT's peers, which trade, on average, at a forward EV/EBIT of 17x. This compares favorably to other dermatology companies that have been acquired in the industry for 13x EV/EBITDA, which means the EV/EBIT could be even higher. Risks to our valuation are as follows: 1) Neothetics faces competition in each of its markets from a number of large and small companies, some of which have greater financial, R&D, production, and other resources than Neothetics; 2) Pipeline failures would delay the company's time to achieve profitability; and 3) Neothetics' drugs may not be commercially successful. This could be a result of better competing products on the market and/or an economic downturn.

CATALYST CALENDAR

Date	Driver	Upcoming Event
2015		
End of '15	LIPO-202	Topline Phase 3 results
2016+		
2H16	LIPO-202	NDA filing
3Q16	LIPO-202	Safety Update
1H17	LIPO-202	FDA AdCom
2H17	LIPO-202	FDA approval
4Q17 or End of 2017	LIPO-202	Launch
2026 to 2031+	LIPO-202	Patents expire

Source: Guggenheim Securities, LLC and Company reports

LIPO-202 MARKET POTENTIAL

We believe LIPO-202 addresses a large, unmet need in aesthetics (\$2B market opportunity today, by our estimate). LIPO-202 appears to offer a better option to existing options for fat reduction. Liposuction involves significant risk, has a long recovery time, and takes four to six months to see an effect. Current non-invasive treatments are lower risk than liposuction; however, these still require equipment space and take two to four months to see an effect. Patients see results with LIPO-202 in one to two months.

There is a lot of opportunity for a faster-acting, non-invasive fat reduction procedure because non-invasive options have only ~1% market share. In the United States, lipoplasty is one of the most common surgical procedures (ASAPS 2013, Neothetics Market Research 2014). Annually:

- Aggregate lipoplasty revenue is over \$1B
- Over 364K procedures are performed (+16% Y/Y)
- Females account for 86% of these procedures
- Lipoplasty is the most common cosmetic surgery for men

According to Neothetics, ~50% of patients who consider a body contouring procedure decide against a procedure

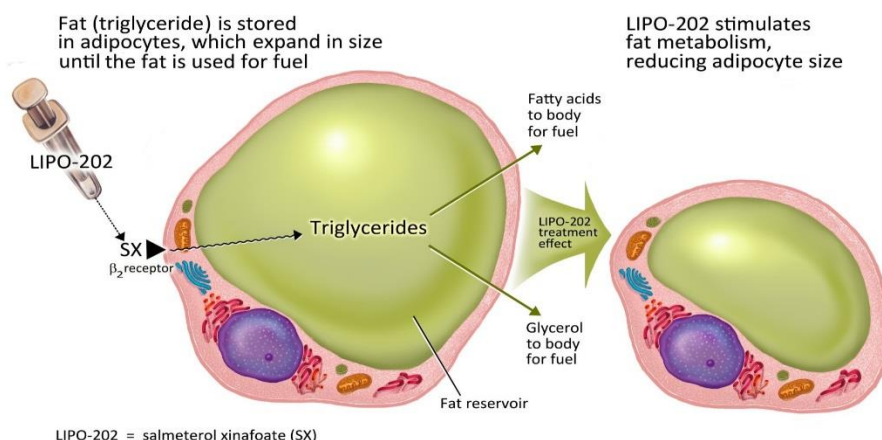
- Demonstrates a need for alternative options
- We estimate the cosmetic injectable patient market is currently over \$2B annually, and these patients would be an easy target for LIPO-202 because they are already familiar with the process associated with other aesthetic injections.

WHAT IS LIPO-202?

LIPO-202 Salmeterol Xinafoate for Injection is a first-in-class, injectable drug treatment for targeted subcutaneous abdominal fat reduction (non-surgical liposuction).

LIPO-202 has a well understood targeted mechanism of action to shrink fat cells.

- It enhances the body's natural lipolysis pathway, resulting in a reduction in size and volume of fat cells in the treatment area.
 - 20 anterior abdomen injections
 - Spaced 4 cm apart
 - Drug diffuses ~ 2 cm in diameter
- Salmeterol xinafoate is a highly selective, long-acting β_2 -adrenergic receptor agonist. Adrenergic receptors play a major role in the regulation of several processes in the body, including fat cell metabolism.
- As shown in the Figure below, salmeterol xinafoate activates β_2 -adrenergic receptors located on human fat cells and triggers the metabolism of triglycerides in these cells to free fatty acids and glycerol by means of the natural process of lipolysis.
- Administering LIPO-202 evenly across the abdomen shrinks fat cells and reduces central abdominal bulging due to subcutaneous fat.
- LIPO-202 reduces local fat stores and the bulges they create with no inflammatory reaction.



Source: Neothetics September 12, 2014 S-1 pages 93

LIPO-202 PHASE II DATA - ENDPOINTS

There are no FDA subjective or objective tools for measuring subcutaneous fat in torso; subjective and objective tools assessed by Neothetics include:

Composite Ratings of Patient Assessment and Clinician Rating Scales: Patient and Clinician Photonumeric rating scale (PnS), Abdominal Contour Scale (P-ACS), and Patient-Reported Patient-Global Abdominal Perception Scale (PGAPS).

- **CPnS** is a clinician rating of the amount of bulging in the central abdomen on a six-point photonumeric scale pursuant to which the clinician performs a match-to-sample from two gender-specific scales of lateral profile torso pictures with progressively larger abdominal bulges.
- **P-GAPS** is a patient self assessment of the amount of bulging in the central abdomen on a five-point ordinal scale, as follows:
 - 0 = Flat
 - 1 = Almost Flat
 - 2 = Slight Bulge, Not Flat
 - 3 = Bulge
 - 4 = Big Bulge

Measurements of Patient Satisfaction: Abdominal Contour Questionnaire (ACQ)

- **ACQ** is a 10-item patient questionnaire on the impact of bulging in the central abdomen, each on an ordinal scale.

Objective Measure: Tape Measure, Canfield 3-D, MRI, Calipers, Laser-Guided Tape Measure, and Ultrasound

- **Laser-Guided Manual Tape Measure Procedure.** Neothetics' procedure involves a precise and reproducible measure of circumference at three levels on the abdomen using patient standardization instructions, such as positioning, posture, breathing, a self-tensioning tape measure, the company's treatment area grid, consisting of a temporary tattoo applied to the central abdomen, and a tripod-mounted laser level to assure horizontal placement of the tape measure.

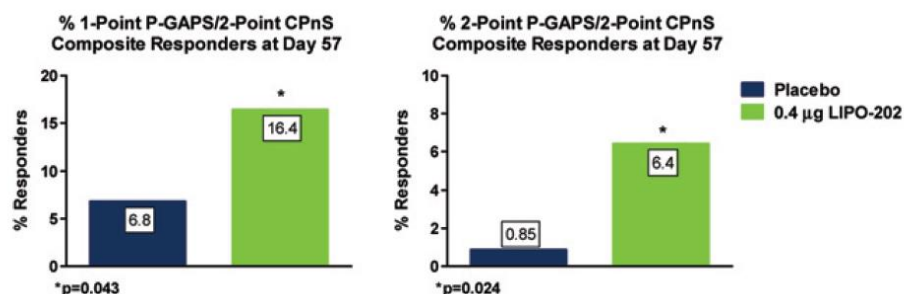
LIPO-202 PHASE II DATA - RESULTS

NEOT completed its largest (N=513) Phase II trial, RESET, in July 2013. Statistically significant reductions in central abdominal bulging due to subcutaneous fat in non-obese patients from baseline at day one and from placebo were demonstrated with a 0.4 µg weekly dose on the key clinical endpoint measures. The 0.4 µg dose is the most effective dose and will be taken forward.

#1. Significant Increase in Responders to LIPO-202 Treatment Compared to Placebo

	Neothetics' Clinical Endpoint Definition	FDA Division of Dermatologic and Dental Product
Responder Endpoint Definition	1-point/grade improvement on P-GAPS that is corroborated by treating clinician as at least a 2-point/grade improvement on the CPnS	2-point/grade improvement on a patient scale that is corroborated by the treating clinician as at least a 2-point/grade improvement on a clinician scale
RESET Results in Patients Treated with 4µg dose weekly for 8 weeks	LIPO-202 treated patients showed 16.4%, versus 6.8% of placebo patients, which translates to statistically significant improvement of p=0.043	LIPO-202 treated patients showed 6.4%, versus less than 1% of placebo patients, which translates to statistically significant improvement of p=0.024

Source: Guggenheim Securities LLC and Neothetics S-1 dated 9/12/14 pages 96 to 97



Source: Neothetics September 12, 2014 S-1 pages 96 to 97

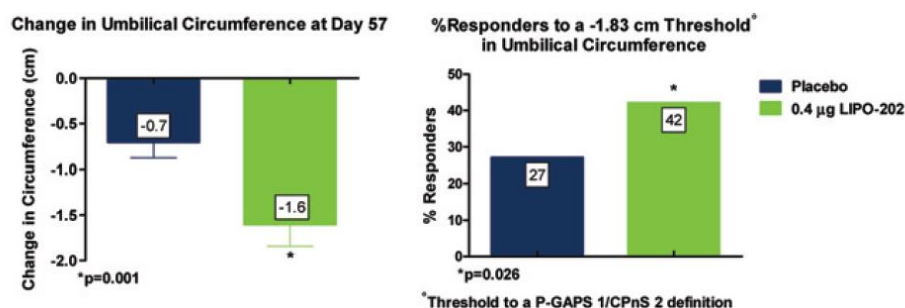
- According to Neothetics, the inverse dose response was a result of Tachyphylaxis. This is a decrease in the response to a drug after its administration. This can sometimes be caused by depletion or marked reduction of the amount of neurotransmitter responsible for creating the drug's effect, or by the depletion of receptors available to which the drug or neurotransmitter can bind. This depletion is caused by the cells reducing the number of receptors in response to their saturation.
- Safety is same percent injection site reaction as placebo; most adverse events were from the needle sticks themselves.

#2. Significant Reduction of Umbilical Circumference by LIPO-202

Laser-guided manual tape measure showed that the 0.4 µg weekly dose produced significant reduction in abdominal circumference at the umbilical circumference compared to placebo, whether expressed as:

- Mean change from baseline.** LIPO-202 0.4 µg dose reduced on average by 1.6 cm, versus 0.7 cm for placebo, which is statistically significant (p=0.001).
- Percentage of responders to a clinically meaningful threshold of a reduction of at least 1.83 cm as defined by 1-point P-GAPS and 2-point CPnS responder definition.** Approximately 42% of patients treated with 0.4 µg dose had a reduction of

at least 1.83 cm, versus 27% of placebo treated patients, which is statistically significant ($p=0.026$).

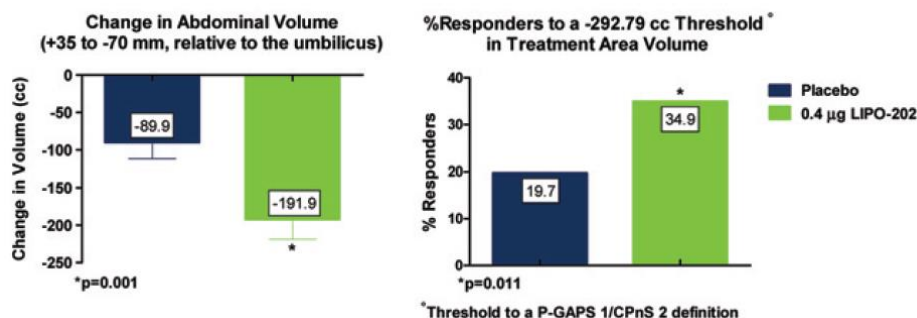


Source: Neothetics September 12, 2014 S-1 pages 97 to 98

#3. Significant Reduction in Treatment Area Volume Produced by LIPO-202.

- Average change from baseline.** 0.4 µg dose reduced treatment area volume on average by 161.6 cubic centimeters (cc), compared to 89.9 cc for placebo, which is statistically significant ($p=0.001$)
- Percentage of responders to a clinically meaningful threshold for reduction in volume of 292.79 cc as defined by the 1-point/grade P-GAPS and 2-point/grade CPnS responder definition.** Approximately 34.9% of patients treated with 0.4 µg dose had a reduction of at least 292.79 cc, compared to 19.7% for placebo, which is statistically significant ($p=0.011$).

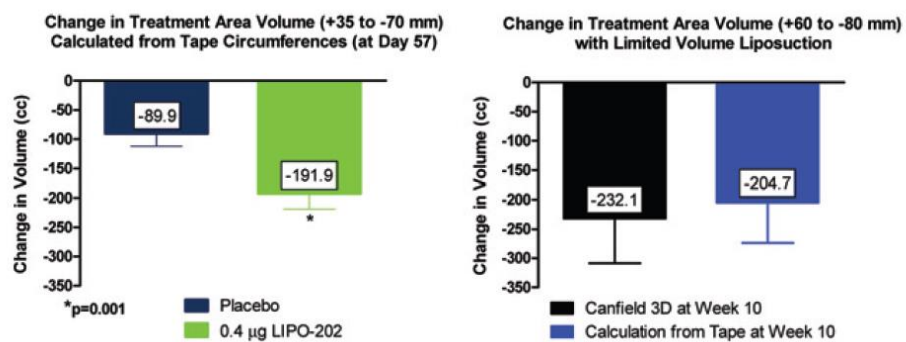
Change from baseline and change from placebo treatment effects with 0.4 µg dose of LIPO-202 were enhanced on all outcome measures in subjects who remained weight neutral or lost weight.



Source: Neothetics September 12, 2014 S-1 pages 98

#4. Similar Reduction in Treatment Area Volume Produced by LIPO-202 and Limited Volume Liposuction in Separate Studies.

In a non-drug, limited volume (VAL-CL-10) liposuction study conducted in a similar population over a similar area, mean reduction in treatment area volume of 200 cc was seen in both eight weeks of treatment with 0.4 µg dose in RESET and limited volume liposuction as assessed 10 weeks after surgery.



Source: Neothetics September 12, 2014 S-1 pages 98

LIPO-202 PHASE III ENDPOINTS

Phase III endpoints will be similar to the RESET Phase II trial.

The Phase III study will be a randomized, double-blind, placebo-controlled study using P-GAPS (Patient-Global Abdominal Perception Scale), CPnS (Clinician Photnumeric Scale), laser-guided tape endpoints, and ultrasound.

- **Primary Endpoints:** % responders with > 1-point improvement P-GAPS and > 2-point CPnS, then > 2-point P-GAPS and > 2-point CPnS.
- **Second Endpoint:** % responders with > 1.83cm decrease in umbilical circumference by laser-guided tape measure.
- **Other Studies:** Bridging PK (to Advair), retreatment study, durability of effect and long-term safety study, special population (obese), and supportive studies.

Based on a meeting with FDA, Neothetics will initiate and complete an additional exploratory evaluation of two-dimensional ultrasound as a secondary measure in the Phase III trials. During Neothetics' end of Phase II meeting with the FDA, the Agency expressed concerns regarding Neothetics' proposed endpoint tools used to assess the efficacy of LIPO-202. The FDA wanted to know if a more accurate tool than laser-guided tape measure could be used to assess the reduction of subcutaneous fat. Therefore, Neothetics is using both two-dimensional ultrasound and laser-guided tape measure in the Phase III trials.

	Clinical Trial	# of Patients	Trial Purpose	Expected Trial Initiation	Data Expected
Pivotal	LIPO-202-CL-18	800	Pivotal Phase III clinical trial of safety & efficacy	1H15	End of 2015
	LIPO-202-CL-19	800	Pivotal Phase III clinical trial of safety and efficacy (identical design to LIPO-202-CL18)	1H15	End of 2015
Supporting	LIPO-202-CL-12	24	Comparative bioavailability of LIPO-202 and Advair Diskus 500/50 Clinical bridge for 505(b)(2) NDA	1H15	1H15
	LIPO-202-CL-21	120	Safety in a special population of obese patients	1H15	2H15
	LIPO-202-CL-22	120	Long-term safety of repeated cycles of treatment	1H15	1H16
	LIPO-202-CL-23	200	Long-term safety and durability of efficacy in responders to treatment	2H15	2H16
Supplement	LIPO-202-CL-25	10 to 12	Exploratory study in submental fat	1H15	2H15
	LIPO-202-CL-26	10 to 12	Exploratory study in lipomas	1H15	2H15

Source: Guggenheim Securities LLC and Neothetics S-1 dated 9/12/14 pages 99 to 100.

FINANCIAL MODEL

Neothetics - Quarterly Income Statement Analysis 2012-2025E

(\$ in Millions, except per share amounts)																			CAGR 20E-25E
(Year Ended December 31)	2012	2013	2014	1Q15A	2Q15A	3Q15E	4Q15E	2015E	2016E	2017E	2018E	2019E	2020E	2021E	2022E	2023E	2024E	2025E	
Total Revenue	0.1	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	109.1	231.4	471.5	657.7	850.1	949.8	1,059.1	1,178.8	20.1
COGS	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	27.3	25.5	42.4	52.6	68.0	76.0	84.7	94.3	
Gross Profit	0.1	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	81.8	206.0	429.0	605.1	782.1	873.8	974.3	1084.5	
R&D	3.2	11.4	5.2	4.7	7.5	18.3	17.3	47.8	35.1	37.0	37.7	38.5	30.0	30.6	31.2	31.8	32.5	33.1	
SG&A	2.6	3.0	4.4	1.9	1.7	6.5	7.3	17.5	55.0	95.0	110.0	168.0	171.4	174.8	178.3	181.8	185.5	189.2	
Total Operating Expense	5.8	14.4	9.6	6.6	9.2	24.8	24.6	65.2	90.1	132.0	147.7	206.5	201.4	205.4	209.5	213.7	218.0	222.3	
Operating Income	(5.7)	(14.4)	(9.6)	(6.6)	(9.2)	(24.8)	(24.6)	(65.2)	(90.1)	(132.0)	(65.9)	(0.5)	227.7	399.7	572.6	660.1	756.4	862.1	30.5
Interest Income	0.0	0.0	0.0	0.0	0.0	0.2	0.6	0.8	0.1	0.7	1.3	0.3	0.1	1.1	3.1	6.3	10.1	14.5	
Interest Expense	(0.9)	(0.1)	(0.4)	(0.3)	(0.3)	(0.2)	(0.2)	(1.0)	(0.9)	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	
Other Income/(Expense)	(1.2)	(0.5)	(0.9)	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	
Non-Operating Items	(2.1)	(0.6)	(1.2)	(0.3)	(0.3)	(0.0)	0.3	(0.2)	(0.8)	0.7	1.3	0.3	0.1	1.1	3.1	6.3	10.1	14.5	
Pre-tax Income	(7.8)	(15.0)	(10.8)	(6.9)	(9.5)	(24.8)	(24.3)	(65.5)	(90.9)	(131.3)	(64.7)	(0.2)	227.8	400.8	575.7	666.4	766.5	876.7	
Tax Rate	0.0%	0.0%	0.0%	0.0%	0.0%	0.0%	0.0%	0.0%	0.0%	0.0%	0.0%	0.0%	40.0%	40.0%	40.0%	40.0%	40.0%	40.0%	
Taxes	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	91.1	160.3	230.3	266.6	306.6	350.7	
Net Income	(7.8)	(15.0)	(10.8)	(6.9)	(9.5)	(24.8)	(24.3)	(65.5)	(90.9)	(131.3)	(64.7)	(0.2)	136.7	240.5	345.4	399.8	459.9	526.0	30.9
Diluted Shares Outstanding	3.1	3.1	3.8	13.7	13.7	13.8	13.9	13.8	19.0	24.0	24.1	24.2	24.3	24.4	24.5	24.6	24.7	24.8	
Adjusted Diluted EPS	(\$2.57)	(\$4.81)	(\$2.83)	(\$0.50)	(\$0.69)	(\$1.80)	(\$1.75)	(\$4.75)	(\$4.79)	(\$5.48)	(\$2.69)	(\$0.01)	\$5.63	\$9.87	\$14.12	\$16.28	\$18.65	\$21.24	30.4
% Change																			
Total Revenue	-100.0%	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	112.2%	103.7%	39.5%	29.3%	11.7%	11.5%	11.3%	
COGS	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	-6.6%	66.7%	24.0%	29.3%	11.7%	11.5%	11.3%	
Gross Profit	-100.0%	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	151.8%	108.3%	41.0%	29.3%	11.7%	11.5%	11.3%	
R&D	252.3%	-54.8%	NM	NM	NM	NM	NM	823.2%	-26.5%	5.4%	2.0%	2.0%	-22.1%	2.0%	2.0%	2.0%	2.0%	2.0%	
SG&A	14.8%	48.4%	NM	NM	NM	NM	NM	295.5%	214.9%	72.7%	15.8%	52.7%	2.0%	2.0%	2.0%	2.0%	2.0%	2.0%	
Operating Income	NM	NM	NM	NM	NM	NM	NM	NM	38.1%	46.5%	-50.0%	-99.2%	-42984.7%	75.6%	43.3%	15.3%	14.6%	14.0%	
Non-Operating Items	NM	NM	NM	NM	NM	NM	NM	NM	229.7%	-188.1%	84.1%	-74.6%	-56.7%	648.4%	197.1%	99.1%	61.4%	43.9%	
Pre-tax Income	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	75.9%	43.7%	15.7%	15.0%	14.4%	
Tax Rate	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	0.0%	0.0%	0.0%	0.0%	0.0%	
Taxes	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	75.9%	43.7%	15.7%	15.0%	14.4%	
Net Income	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	75.9%	43.7%	15.7%	15.0%	14.4%	
Adjusted Diluted EPS	87.4%	-41.1%	2.5%	2.6%	48.2%	290.0%	67.6%	1.0%	14.3%	-51.0%	-99.7%	NM	75.2%	43.1%	15.3%	14.6%	13.9%		

Source: Guggenheim Securities, LLC and Company Reports

(1) Assume 40% tax rate when profitable, to be conservative.

Neothetics - Quarterly Margin Analysis 2012-2025E

	2012	2013	2014	1Q15A	2Q15A	3Q15E	4Q15E	2015E	2016E	2017E	2018E	2019E	2020E	2021E	2022E	2023E	2024E	2025E
Total Revenue	100.0%	100.0%	100.0%	100.0%	100.0%	100.0%	100.0%	100.0%	100.0%	100.0%	100.0%	100.0%	100.0%	100.0%	100.0%	100.0%	100.0%	100.0%
COGS	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	25.0%	11.0%	9.0%	8.0%	8.0%	8.0%	8.0%	8.0%
Gross Margin	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	75.0%	89.0%	91.0%	92.0%	92.0%	92.0%	92.0%	92.0%
R&D	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	34.6%	16.6%	6.4%	4.7%	3.7%	3.4%	3.1%	2.8%
SG&A	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	100.9%	72.6%	36.3%	26.6%	21.0%	19.1%	17.5%	16.1%
Oper. Inc.	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	-60.5%	-0.2%	48.3%	60.8%	67.4%	69.5%	71.4%	73.1%
Operating Expenses	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	135.5%	89.2%	42.7%	31.2%	24.6%	22.5%	20.6%	18.9%
Non-Oper. Items	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	1.2%	0.1%	0.0%	0.2%	0.4%	0.7%	1.0%	1.2%
Pretax Income	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	-59.3%	-0.1%	48.3%	60.9%	67.7%	70.2%	72.4%	74.4%
Net Income	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	-59.3%	-0.1%	29.0%	36.6%	40.6%	42.1%	43.4%	44.6%

Source: Guggenheim Securities, LLC and Company Reports

Neothetics - Quarterly Revenue Model 2012-2025E

(\$ in Millions)																		
(Year Ended December 31)	2012	2013	2014	1Q15A	2Q15A	3Q15E	4Q15E	2015E	2016E	2017E	2018E	2019E	2020E	2021E	2022E	2023E	2024E	2025E
Base Business																		
LIPO-102	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
LIPO-202	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	109.1	231.4	471.5	657.7	850.1	949.8	1,059.1	1,178.8
Total Base Business Revenues	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	109.1	231.4	471.5	657.7	850.1	949.8	1,059.1	1,178.8
New Products																		
Product 1	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Product 2	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Product 3	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Total New Product Revenues	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Other	0.1	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Total Revenues	0.1	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	109.1	231.4	471.5	657.7	850.1	949.8	1,059.1	1,178.8
%Change																		
Base Business																		
LIPO-102		NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM
LIPO-202		NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	112.2%	103.7%	39.5%	29.3%	11.7%	11.5%	11.3%
Total Base Business Revenues		NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	112.2%	103.7%	39.5%	29.3%	11.7%	11.5%	11.3%
New Products																		
Product 1		NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM
Product 2		NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM
Product 3		NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM
Total New Product Revenues		NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM
Other		-100.0%	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM
Total Revenues		-100.0%	NM	NM	NM	NM	NM	NM	NM	NM	NM	112.2%	103.7%	39.5%	29.3%	11.7%	11.5%	11.3%

Source: Guggenheim Securities, LLC and Company Reports

Neothetics - Annual Revenue Model 2012-2025E

(\$ in Millions)	Year Ended December 31st,																										CAGR 18E-25E		
	2012	2013	2014	2015E	2016E	2017E	2018E	2019E	2020E	2021E	2022E	2023E	2024E	2025E	2013E/ 2012	2014/ 2013	2015E/ 2014	2016E/ 2015E	2017E/ 2016E	2018E/ 2017E	2019E/ 2018E	2020E/ 2019E	2021E/ 2020E	2022E/ 2021E	2023E/ 2022E	2024E/ 2023E		2025E/ 2024E	
Base Business																													
LPO-102	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	
LPO-202	0.0	0.0	0.0	0.0	0.0	0.0	109.1	231.4	471.5	657.7	850.1	949.8	1,059.1	1,178.8	NM	NM	NM	NM	NM	NM	112.2%	103.7%	39.5%	29.3%	11.7%	11.5%	11.3%	40.5%	
Total Base Business Revenues	0.0	0.0	0.0	0.0	0.0	0.0	109.1	231.4	471.5	657.7	850.1	949.8	1,059.1	1,178.8	NM	NM	NM	NM	NM	NM	112.2%	103.7%	39.5%	29.3%	11.7%	11.5%	11.3%	40.5%	
New Products																													
Product 1	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	
Product 2	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	
Product 3	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	
Total New Product Revenues	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	
Other	0.1	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	-100.0%	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	
Total Revenues	0.1	0.0	0.0	0.0	0.0	0.0	109.1	231.4	471.5	657.7	850.1	949.8	1,059.1	1,178.8	-100.0%	NM	NM	NM	NM	NM	NM	112.2%	103.7%	39.5%	29.3%	11.7%	11.5%	11.3%	40.5%

Source: Guggenheim Securities, LLC and Company Reports

Neothetics - Cash Flow Analysis 2012-2025E

(\$ in Millions)														
(Year ended December 31)	2012	2013	2014	2015E	2016E	2017E	2018E	2019E	2020E	2021E	2022E	2023E	2024E	2025E
Cash flows provided by Operating Activities:														
Net Income/(loss)	(7.8)	(15.0)	(10.8)	(65.5)	(90.9)	(131.3)	(64.7)	(0.2)	136.7	240.5	345.4	399.8	459.9	526.0
Depreciation and Amortization	0.1	0.1	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Other	2.1	0.6	1.6	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Net Changes in Working Capital	(1.8)	1.4	(0.3)	0.0	0.0	0.0	(31.2)	(18.4)	(45.0)	(31.8)	(33.7)	(15.3)	(16.5)	(17.8)
Net cash provided by Operating Activities	(7.4)	(12.9)	(9.6)	(65.5)	(90.9)	(131.3)	(95.8)	(18.5)	91.7	208.6	311.7	384.5	443.4	508.3
Cash flows from Investing Activities														
Purchase/Proceeds of property and equipment	0.0	0.0	(0.0)	(0.0)	(0.0)	(0.0)	(0.0)	(0.0)	(0.0)	(0.0)	(0.0)	(0.0)	(0.0)	(0.0)
Other	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Net cash used in Investing Activities	0.0	0.1	(0.0)	(0.0)	(0.0)	(0.0)	(0.0)	(0.0)	(0.0)	(0.0)	(0.0)	(0.0)	(0.0)	(0.0)
Cash flows from Financing Activities														
Proceeds/Payment on Notes Payable and Bank Loan	2.4	(0.4)	9.8	0.1	0.2	(10.0)	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Issuance of Common Stocks and Other	0.0	0.0	57.8	0.0	150.0	200.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Issuance of Preferred Stock for Cash, net of offering	10.2	6.5	13.6	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Net cash (used in) provided by Financing Activities	12.7	6.1	81.2	0.1	150.2	190.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Net (decrease) increase in cash and equivalents	5.3	(6.7)	71.6	(65.4)	59.2	58.7	(95.8)	(18.6)	91.7	208.6	311.7	384.5	443.4	508.2
Cash and equivalents at beginning of year	5.8	11.1	4.4	75.9	10.6	69.8	128.5	32.7	14.1	105.9	314.5	626.2	1,010.7	1,454.1
Cash and equivalents at end of year	11.1	4.4	75.9	10.6	69.8	128.5	32.7	14.1	105.9	314.5	626.2	1,010.7	1,454.1	1,962.4

Source: Guggenheim Securities, LLC and Company Reports

Neothetics - Balance Sheet Analysis 2012-2025E

(\$ in Millions)														
(Year ended December 31)	2012	2013	2014	2015E	2016E	2017E	2018E	2019E	2020E	2021E	2022E	2023E	2024E	2025E
Assets														
Cash and Cash Equivalents	11.1	4.4	75.9	10.6	69.8	128.5	32.7	14.1	105.9	314.5	626.2	1,010.7	1,454.1	1,962.4
Accounts receivables	0.0	0.0	0.0	0.0	0.0	0.0	18.2	37.9	76.0	104.4	132.8	146.1	160.5	175.9
Inventory	0.0	0.0	0.0	0.0	0.0	0.0	13.0	11.6	18.4	21.9	27.2	29.2	31.4	33.7
Prepaid expenses and other current assets	1.5	0.1	0.9	0.9	0.9	0.9	0.9	0.9	0.9	0.9	0.9	0.9	0.9	0.9
Total Current Assets	12.6	4.5	76.9	11.5	70.7	129.4	64.8	64.6	201.3	441.7	787.2	1,187.0	1,646.9	2,172.9
Property and equipment, net	0.2	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Restricted cash	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Other assets	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Total Other Assets	0.2	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
TOTAL ASSETS	12.8	4.5	76.9	11.5	70.8	129.5	64.8	64.6	201.3	441.8	787.2	1,187.0	1,646.9	2,172.9
Liabilities & Shareholder's Equity														
Accounts Payable	0.7	0.6	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0
Accrued Expenses	0.6	0.7	0.9	0.9	0.9	0.9	0.9	0.9	0.9	0.9	0.9	0.9	0.9	0.9
Current Portion of Term Loan Payable	0.6	0.2	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Other Liabilities	0.0	0.0	9.7	9.8	10.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Total Liabilities	1.9	1.5	11.7	11.8	11.9	1.9	1.9	1.9	1.9	1.9	1.9	1.9	1.9	1.9
Total Stockholder's Equity	10.9	3.0	65.2	-0.2	58.9	127.6	62.9	62.7	199.4	439.8	785.3	1,185.1	1,645.0	2,171.0
TOTAL LIABILITIES & STOCKHOLDERS' EQUITY	12.8	4.5	76.9	11.5	70.8	129.5	64.8	64.6	201.3	441.8	787.2	1,187.0	1,646.9	2,172.9

Source: Guggenheim Securities, LLC and Company Reports

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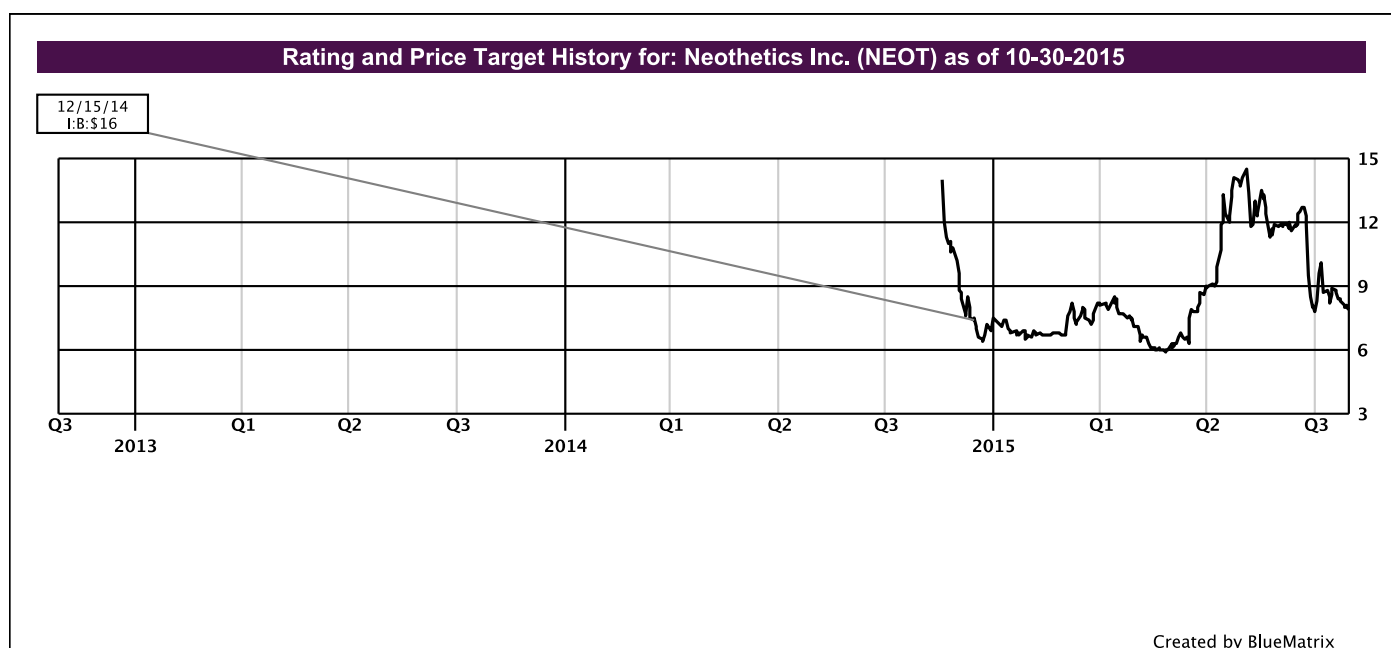
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