OUTPERFORM

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Reason for report: **EARNINGS**

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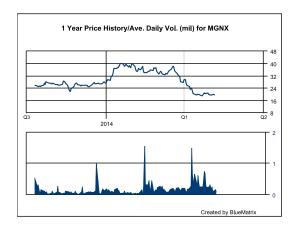
MACROGENICS, INC.

1Q14 Recap - Pipeline Progressing with MAGENTA Study to Initiate in 2H14

- · Bottom Line: MGNX reported 1Q14 financial results today and expects to be sufficiently capitalized to fund the company into 2017. The company's pipeline programs remain on track, and MGNX expects to have six clinical programs active by the end of 2015. MGNX also provided incremental details on MAGENTA, the Phase III margetuximab study to initiate in 2H14 in gastroesophageal cancer. We continue to view MGNX's diversified pipeline strategy positively since it generates several shots on goal, and it takes advantage of MGNX's innovative drug discovery and development capabilities. Reiterate Outperform rating and \$52 price target.
- Company is well capitalized to drive its product pipeline forward. MGNX reported total revenue of \$14.7M and net loss of \$3.1M for 1Q14 vs. our revenue estimate of \$20.3M and net loss of \$1.5M. Management expects that its current cash and equivalents should fund the company into 2017. MGNX ended 1Q14 with \$198.7M in cash and equivalents.
- Pipeline continues to progress with six clinical programs projected by the end of 2015. MGNX expects to initiate MAGENTA, a Phase III study of margetuximab in advanced gastroesophageal cancer, in 2H14. A Phase IIa study of margetuximab in metastatic breast cancer continues to enroll patients with data from Part A expected in late 2014. Timelines for MGA271 are on track with completion of the first three dose expansion cohorts expected by YE14, and the initiation of additional monotherapy expansion cohorts is expected in 2H14. MGD006 will be MGNX's first dual-affinity retargeting (DART)-based molecule to enter the clinic in 2Q14 in patients with acute myeloid leukemia. MGNX also plans to advance MGD007 into the clinic for colorectal cancer in 2H14 and two additional DARTs into the clinic for oncology in 2015. We believe first-inman DART data should further derisk MGNX's platform.
- · Additional details on MAGENTA provided. MAGENTA (Phase III margetuximab trial in gastroesophageal cancer) is 90% powered to detect an overall survival hazard ratio of 0.67. The trial will evaluate the 6mg/ kg weekly dose of margetuximab in combination with chemotherapy vs. chemotherapy alone in 425 Her2-positive patients refractory to two lines of prior therapy.



| ndex: 1,2 | 212.60 |
|-------------------------|-------------------------|
| | 19.64 |
| \$ | 52.00 |
| Sum-of-the-parts DCF ar | nalysis |
| 9 | \$41.00 |
| \$ | 18.35 |
| I): | 26.3 |
| nil): | 516.5 |
| | \$0.00 |
| | \$7.47 |
| | \$0.00 |
| | 0.0% |
| | Sum-of-the-parts DCF ar |



| Dec Yr | 1Q | 2Q | 3Q | 4Q | FY Rev | 1Q | 2Q | 3Q | 4Q | FY EPS | |
|-------------|---------|--------|--------|--------|--------|-----------|----------|----------|----------|----------|----|
| 2013A | \$10.6 | \$12.3 | \$20.2 | \$14.9 | \$58.0 | (\$2.80) | (\$0.29) | \$0.14 | (\$0.13) | (\$0.04) | NM |
| 2014E - New | \$14.7A | \$9.8 | \$9.8 | \$9.8 | \$44.0 | (\$0.12)A | (\$0.38) | (\$0.43) | (\$0.48) | (\$1.42) | NM |
| 2014E - Old | \$20.3 | \$7.9 | \$7.9 | \$7.9 | \$44.0 | (\$0.05) | (\$0.50) | (\$0.50) | (\$0.50) | (\$1.57) | NM |
| 2015E - New | | | | | \$48.0 | i | | | | (\$1.96) | NM |
| 2015E - Old | | | | | \$48.0 | | | | | (\$2.19) | NM |

Source: Company Information and Leerink Partners LLC Research

Revenues in \$MM.

GAAP EPS. Quarterly figures may not sum to annual total due to change in shares out.



INVESTMENT THESIS

We rate MGNX Outperform. MGNX is a leader in the area of immune-modulation and is a fully integrated R&D-driven biotechnology company. The company is focused on developing new antibody-based therapeutics for cancer and autoimmune diseases and is based on a suite of platform technologies that allow rapid generation of novel therapeutic antibodies with superior properties. MGNX has applied its antibody discovery and engineering platform to generate a proprietary product pipeline and to enter into strategic collaborations that provide the company with funding and leverage the additional expertise of partners. We believe MGNX shares will appreciate as the company advances its proprietary and partnered pipeline products. We also believe MGNX could close additional product development partnerships and existing partners could license further products.

VALUATION

Our \$52 price target for MGNX shares in 12 months is based on a discounted cash flow (DCF) sum-of-parts analysis. We use a 12% discount rate for probability of success-weighted margetuximab (25%), MGA271 (15%), and MGD006 (20%) sales. Based on our DCF analysis, we attribute \$8/share to margetuximab, \$11/share to MGA271, \$9/share to MGD006, and \$18/share to the preclinical pipeline and platform and the remainder to expected cash in one year.

RISKS TO VALUATION

Developmental pipeline agents face clinical and regulatory development risk, as well as commercial risks. MGNX also faces execution risk and financial risk. We estimate that MGNX's current cash will be sufficient to fund operations into 2017, and the company may have additional financing needs before turning cash flow positive.

| | | | | Difference |
|--------------------------------|------------|------|--------|------------|
| 1Q14 (\$M, except EPS) | Leerink 1Q | L4E | 1Q14A | (Leerink) |
| Collaborative research revenue | 20 | 0.0 | 14.4 | (5.6) |
| Grant revenue | | 0.3 | 0.3 | 0.1 |
| Royalties | | - | - | - |
| Product sales | | - | - | - |
| Total Revenue | 20 | 0.3 | 14.7 | (5.5) |
| COGS | | - | - | - |
| R&D | 10 | 5.8 | 14.6 | (2.2) |
| SG&A | ! | 5.0 | 3.3 | (1.7) |
| Operating expenses | 2: | 1.8 | 17.8 | (3.9) |
| Operating income (expense) | (: | 1.5) | (3.1) | (1.6) |
| Interest income (expense) | | - | - | - |
| Other income (expense) | | - | 0.0 | 0.0 |
| Total Other income (expense) | | - | 0.0 | 0.0 |
| EBT | (: | 1.5) | (3.1) | (1.6) |
| Tax expense (income) | | - | - | - |
| Net income | (: | L.5) | (3.1) | (1.6) |
| GAAP EPS | (0. | 05) | (0.12) | (0.06) |
| Common shares outstanding | 2. | 7.5 | 26.3 | (1.2) |

| BS & CFS | Leerink 1Q14E | 1Q14A | Difference (Leerink) |
|--------------------|---------------|-------|-------------------------|
| Cash & equivalents | 193.2 | 198.7 | 5.6 |
| Debt | - | - | - |
| Deferred revenue | - | - | - |

Source: Leerink Partners Estimates and Company Filings

| Program | Target | Platform | Partner | Indication | Current Status | Next milestone | Timing |
|--------------|---------------|------------|--------------------------|----------------------------|----------------|--|-----------|
| Margetuximab | Her2 | Fc | proprietary | Breast cancer (Her2 IHC2+) | Phase IIa | Phase IIa data | late 2014 |
| | | | | | | Initiate Phase IIb/III | 2015 |
| | | | | 3rd line Gastric cancer | Phase I | Initiate Phase III (MAGENTA) | 2H14 |
| | | | | | | Phase III data | 2018 |
| | | | | Other cancers (bladder) | Phase I | Initiate Phase II | 2015 |
| | | | | HER2-expressing tumors | Phase I | Additional dosing data | 2014/2015 |
| MGA271 | B7-H3 | Fc, CSLC | Servier (EU rights) | Solid tumors | Phase Ib | Phase Ib initiation of additional cohorts | 2H14 |
| | | | | | | Phase Ib dose-expansion data monotherapy | 4Q14/1H15 |
| | | | | | | Phase Ib dose-expansion data monotherapy (addl. cohorts) | 2015 |
| | | | | | | Phase Ib dose-expansion data monotherapy (Servier cohorts) | 2015 |
| | | | | | | Phase Ib initiation of combination studies | 1Q15 |
| | | | | | | Phase Ib dose-expansion data combinations | 2015 |
| | | | | | | Servier opt-in | 2015 |
| MGD006 | CD123 x CD3 | DART | Servier (EU rights) | AML | IND | Phase I DE data | 2H15 |
| MGD007 | gpA33 x CD3 | DART, CSLC | Servier (EU rights) | Colorectal cancer | Preclinical | IND accepted | mid-14 |
| | | | | | | Initiate Phase I | 2H14 |
| | | | | | | Phase I data | 2H15 |
| | | | | | | Servier opt-in | 2014/15 |
| MGD010 | CD32B x CD79B | DART | proprietary | Autoimmune (SLE, RA) | Preclinical | Partnership | 2014/15 |
| | | | | | | Initiate Phase I | 2015 |
| MGD011 | undisclosed | DART | proprietary | Oncology | Preclinical | Initiate Phase I | 2015 |
| MGD012 | undisclosed | DART | proprietary/Servier (EU) | Oncology | Preclinical | Initiate Phase I | 2015 |
| Teplizumab | CD3 | Fc | proprietary | Type 1 Diabetes | IS Study | Partnership | n/a |

Source: SEC Filings, Leerink Partners Estimates

| MGNX P&L | 2011A | 2012A | 2013A | 1Q14A | 2Q14E | 3Q14E | 4Q14E | 2014E | 2015E |
|--------------------------------|-------|--------|--------|--------|--------|--------|--------|--------|--------|
| Collaborative research revenue | 47.1 | 59.6 | 56.8 | 14.4 | 9.5 | 9.5 | 9.5 | 43.0 | 48.0 |
| Grant revenue | 10.2 | 4.2 | 1.3 | 0.3 | 0.2 | 0.2 | 0.2 | 1.0 | - |
| Royalties | - | - | - | - | - | - | - | - | - |
| Product sales | - | - | - | - | - | - | - | - | - |
| Total Revenue | 57.2 | 63.8 | 58.0 | 14.7 | 9.8 | 9.8 | 9.8 | 44.0 | 48.0 |
| cogs | - | - | - | - | - | - | - | - | - |
| R&D | 41.1 | 45.4 | 46.6 | 14.6 | 16.0 | 17.5 | 18.9 | 67.0 | 83.0 |
| SG&A | 10.9 | 10.2 | 11.1 | 3.3 | 4.0 | 4.0 | 4.0 | 15.3 | 20.0 |
| Operating expenses | 52.0 | 55.6 | 57.7 | 17.8 | 20.0 | 21.5 | 22.9 | 82.3 | 103.0 |
| Operating income (expense) | 5.2 | 8.2 | 0.37 | (3.1) | (10.2) | (11.7) | (13.2) | (38.3) | (55.0) |
| Total Other income (expense) | 1.5 | 0.2 | (0.6) | 0.0 | - | - | - | 0.0 | - |
| ЕВТ | 6.7 | 8.4 | (0.3) | (3.1) | (10.2) | (11.7) | (13.2) | (38.3) | (55.0) |
| Tax expense (income) | - | - | - | - | - | - | - | - | - |
| Net income | 6.7 | 8.4 | (0.3) | (3.1) | (10.2) | (11.7) | (13.2) | (38.3) | (55.0) |
| GAAP EPS | 6.55 | 7.72 | (0.04) | (0.12) | (0.38) | (0.43) | (0.48) | (1.42) | (1.96) |
| Common shares outstanding | 1.0 | 1.1 | 6.8 | 26.3 | 26.7 | 27.1 | 27.5 | 26.9 | 28.0 |
| BS & CFS | 2011A | 2012A | 2013A | 1Q14A | 2Q14E | 3Q14E | 4Q14E | 2014E | 2015E |
| Cash & equivalents | 55.2 | 47.7 | 116.5 | 198.7 | 189.0 | 177.9 | 165.5 | 165.5 | 117.8 |
| Debt | - | - | - | - | - | - | - | - | - |
| Change in Cash | 18.3 | (7.5) | 68.7 | 73.2 | (9.7) | (11.1) | (12.4) | 40.9 | (47.8) |
| Cash from operations | 6.8 | (6.6) | (14.2) | (3.1) | (9.3) | (10.6) | (11.9) | (34.1) | (45.8) |
| Net income (loss) | 6.7 | 8.4 | (0.3) | (3.1) | (10.2) | (11.7) | (13.2) | (38.3) | (55.0) |
| Share based comp | 2.3 | 0.8 | 0.9 | 0.6 | 1.6 | 1.7 | 1.8 | 6.6 | 8.2 |
| D&A | 1.1 | 1.0 | 1.2 | 0.4 | 0.4 | 0.4 | 0.4 | 1.6 | 1.0 |
| Other (Change in WC) | (3.5) | (16.7) | (16.0) | (1.0) | (1.0) | (1.0) | (1.0) | (4.0) | - |
| Cash from investing | (0.5) | (0.9) | (3.0) | (0.4) | (0.4) | (0.4) | (0.4) | (1.8) | (2.0) |
| CapEx | (0.5) | (0.9) | (3.0) | (0.4) | (0.4) | (0.4) | (0.4) | (1.8) | (2.0) |
| Acquisitions | - | - | - | - | - | - | - | - | - |
| Other | - | - | - | - | - | - | - | - | - |
| Cash from financing | 12.1 | 0.0 | 85.9 | 76.8 | - | _ | - | 76.8 | - |
| Equity issue (buyback) | 12.1 | 0.0 | 85.9 | 76.8 | - | - | - | 76.8 | - |
| Debt issue (principal payment) | - | - | - | - | - | - | - | - | - |
| Other | - | - | - | - | - | - | - | - | - |

Source: Leerink Partners Estimates and Company Filings



Disclosures Appendix Analyst Certification

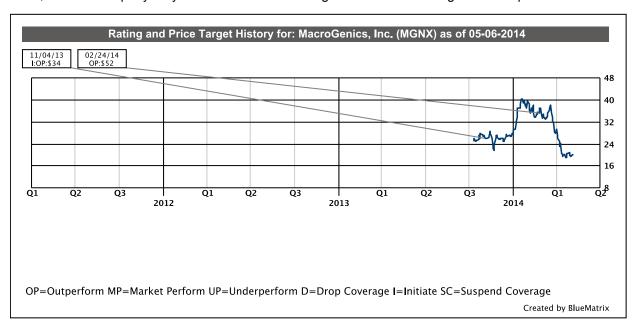
I, Michael Schmidt, Ph.D., certify that the views expressed in this report accurately reflect my views and that no part of my compensation was, is, or will be directly related to the specific recommendation or views contained in this report.

Valuation

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| | Distribution of Ratings/Investment Bank | ing Services (IB | , | erv./Past 12 Mos. |
|-----------|---|------------------|-------|----------------------|
| Rating | Count | Percent | Count | Percent |
| BUY [OP] | 131 | 68.23 | 46 | 35.11 |
| HOLD [MP] | 61 | 31.77 | 3 | 4.92 |
| SELL [UP] | 0 | 0.00 | 0 | 0.00 |

Explanation of Ratings

Outperform (Buy): We expect this stock to outperform its benchmark over the next 12 months.

<u>Market Perform (Hold/Neutral):</u> We expect this stock to perform in line with its benchmark over the next 12 months.

<u>Underperform (Sell):</u> We expect this stock to underperform its benchmark over the next 12 months. The degree of outperformance or underperformance required to warrant an Outperform or an Underperform rating should be commensurate with the risk profile of the company.

For the purposes of these definitions the relevant benchmark will be the S&P 600® Health Care Index for issuers with a market capitalization of less than \$2 billion and the S&P 500® Health Care Index for issuers with a market capitalization over \$2 billion.

Important Disclosures

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Leerink Partners LLC makes a market in MacroGenics, Inc.

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