Zacks Small-Cap Research

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Corgenix Medical (CONX-OTC)

CONX: *Q3 Earnings - Revenue Hits Our Number But Soft GM Impacts Income*

Current Recommendation	**Outperform**
Prior Recommendation	N/A
Date of Last Change	10/18/11
Current Price (05/15/12)	$0.12
Target Price	**$0.50**

OUTLOOK

Corgenix is on the front-end of a transition away from reliance on certain legacy products to a more determined focus on growing higher potential areas of their business. Highlighted by the recent transfer of their international business and related joint product development agreements, the transition is expected to help jump-start revenue growth and form the basis for continued double-digit growth over the next several years.

Based on comparable valuations, Corgenix's shares trade significantly cheaper than warranted. We are maintaining our Outperform rating and $0.50 per share price target.

SUMMARY DATA

52-Week High	**$0.19**
52-Week Low	**$0.08**
One-Year Return (%)	**38.25**
Beta	**1.32**
Average Daily Volume (sh)	**24,912**
Shares Outstanding (mil)	**45**
Market Capitalization ($mil)	**$6**
Short Interest Ratio (days)	**N/A**
Institutional Ownership (%)	**0**
Insider Ownership (%)	**16**
Annual Cash Dividend	**$0.00**
Dividend Yield (%)	**0.00**
5-Yr. Historical Growth Rates	
Sales (%)	**1.0**
Earnings Per Share (%)	**N/A**
Dividend (%)	**N/A**
P/E using TTM EPS	**N/A**
P/E using 2012 Estimate	**N/A**
P/E using 2013 Estimate	**N/A**
Zacks Rank	**N/A**

Risk Level	**High,**
Type of Stock	**Small-Value**
Industry	**Med Products**

ZACKS ESTIMATES

Revenue
(in '000)

	Q1 (Sep)	Q2 (Dec)	Q3 (Mar)	Q4 (Jun)	Year (Jun)
2011	$1,978 A	$1,786 A	$1,829 A	$2,349 A	$7,942 A
2012	$2,178 A	$2,066 A	$2,717 A	$2,163 E	$9,124 E
2013					$10,975 E
2014					$12,491 E

Earnings per Share

	Q1 (Sep)	Q2 (Dec)	Q3 (Mar)	Q4 (Jun)	Year (Jun)
2011	-$0.01 A	$0.00 A	$0.00 A	$0.00 A	-$0.01 A
2012	-$0.00 A	-$0.00 A	-$0.00 A	-$0.00 E	-$0.01 E
2013					$0.00 E
2014					$0.01 E

Zacks Projected EPS Growth Rate - Next 5 Years % **N/A**

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Q3 2012 Financial Results

Corgenix reported financial results for the third fiscal quarter ending March 31, 2012 on May 14th. While revenue of $2.7 million was a quarterly record (+49% y-o-y) and just about dead-on with our $2.8 million estimate, net income (-$190k A vs. $188k E) came in lighter than we expected due to a relatively weak gross margin. Management again adjusted full-year revenue guidance downward, now looking for $9.1 million, revised down from $9.7 million three months ago, and $10.1 million issued with Q1 results. In contrast to earlier expectations, international sales have yet to return to growth which is the major cause of the downward revision to revenue guidance over the last few quarters. Management also noted that revenue from contract manufacturing, which can be highly irregular from q-to-q, is also expected to come in lower in Q4 than had been previously anticipated.

Despite the disappointing international revenue and downward revenue guidance for the rest of the current year, Corgenix has seen very strong growth in domestic sales and still expects to show double-digit revenue growth for fiscal 2012. The company also continues to make progress with development of new products for both the international and U.S. markets, some of which should reach commercialization in the near-term. Recent and near-term new product launches remain the major catalyst to driving our modeled revenue growth over the next several years. While we have made some downward revisions to our revenue estimates, the majority relates to the near-years (fiscal 2012, 2013) reflecting the drawn-out turn-around in international revenue and some delay with certain product launches. The impact to our out-years estimates is much more modest.

Following Q3 results we have adjusted our estimated current fiscal year (2012) revenue down from $9.7 million to $9.1 million. Our out-year (2015) revenue has moved from $14.4 million to $14.0 million. We are maintaining our Outperform rating on the stock and our $0.50/share price target.

Revenue

Q3 revenue was $2.72 million, up 49% y-o-y and in-line with our $2.78 million estimate. North America sales were $2.46 million, up 59% y-o-y and dead-on with our $2.44 million estimate. International sales were $253k, down 8% y-o-y and 21% below our $320k estimate. We have adjusted our Q4 international sales estimate downward as traction in this business is clearly taking longer than management had previously anticipated. We continue to model international sales to return to growth in fiscal 2013, fueled by new product launches. We continue to believe that the automated version of AspirinWorks as well as other new product introductions, including those coming out of the ELITech joint product development agreement represent potentially significant growth opportunities for Corgenix's international business over the long-term.

Meanwhile, the domestic business has been very strong with most of the recent growth coming from AspirinWorks (ELISA format), contract manufacturing and contract R&D which has more than offset relative softness in some of the older product lines, namely ELISA phospholipid and coagulation. Contract manufacturing ($582k, +640%) in particular was a major catalyst to Q3 revenue growth as was AspirinWorks ($243k) which has seen sequential revenue growth every quarter since Q4 2011 and was up 167% y-o-y in Q3 2012. Key to stable, long-term revenue growth in the domestic business is new product introductions as revenue from contract manufacturing and contract R&D, which have accounted for approximately 80% of the y-o-y revenue growth through the first nine months of fiscal 2012, can be highly variable.

Net Income / EPS

Q3 net income and EPS came in at ($190k) and ($0.00), compared to our $188k and $0.00 estimates. Operating expenses were largely in-line with our numbers with almost all of the miss on net income coming from lower than modeled gross margin (37.3% A vs. 46.9% E). At least some of the relative weakness in GM (which had averaged ~ 47% in the first half of fiscal 2012) was a result of higher contract R&D and contract manufacturing revenue in the quarter, which are lower-profit segments.

Cash

Corgenix exited Q3 with $1.0 million in cash and equivalents, up from $812k at 12/31/2011. The sequential increase in cash came from 245k inflow from operations, $110k used in investing activities and net inflow of $66k in financing activities. Excluding changes in working capital, operating cash flow was slightly positive for the nine months ending 3/12/2012. Debt remained relatively insignificant at quarter end.

BUSINESS

A pioneer in the field of antiphospholipid antibody detection, Corgenix Medical Corporation is credited with development of the first FDA-cleared assay for measuring anticardiolipin antibodies. The association of antiphospholipid antibodies with thrombosis (antiphospholipid syndrome is a coagulation disorder) encouraged the company to develop assays for other bleeding and clotting disorders. Corgenix has quickly become a renowned expert in the diagnosis and measurement of blood coagulation, vascular occlusion and thrombosis. They offer an array of in vitro diagnostic (IVD) testing products in the areas of **liver diseases** (fibrosis and cirrhosis), **autoimmune disease** (bone and joint disorders) and **vascular disease**, including the only non-blood based test for determining the effectiveness of aspirin, and have sold over 12 million tests to-date.

Most of the company's diagnostic products are developed on an **ELISA** testing format, a method widely used in clinical testing laboratories. As a result of recent R&D collaborations, Corgenix has also implemented additional immunoassay platforms including **lateral flow** (a rapid test format similar to that commonly used for home pregnancy tests) and **immunoturbidimetry** (a liquid-based system for use on automated chemistry analyzers). Although revenue from these additional formats has been immaterial to-date, the immunoturbidimetry format in particular represents an integral portion of the company's growth strategy.

The company has remained competitive in an industry that has experienced substantial consolidation over the recent past by delivering high quality products and service in niche areas such as antiphospholipid syndrome and hemophilia. Over the course of history, sales and distribution have largely been handled in-house. Corgenix's considerable R&D, manufacturing and production capabilities have also afforded the award of several U.S. government contracts and grants as well as facilitated the consummation of partnerships with several leading IVD companies. And while certain products are experiencing rapid growth (i.e. - AspirinWorks), due to increased competition, some of Corgenix's initial areas of expertise now offer less upside opportunity. As a result, the company has begun a measured transition away from reliance on certain legacy tests to a greater focus on faster-growth products and segments. This includes leveraging key partnerships in the development of novel high-potential products and in the expansion of their sales and distribution reach. Corgenix will continue to leverage their R&D/manufacturing capabilities to facilitate product development as well as to attract valuable government grants and manufacturing contracts, all of which are expected to hold outsized opportunities for revenue growth.

Corgenix generates revenue from four main business segments; manufactured products, OEM (private label) products, contract manufacturing, and contract development. Approximately 18% of Corgenix's revenue comes from overseas (i.e. - ex-North America). International revenues fell 36% from 2010 to $1.46 million in fiscal 2011 while sales in North America increased by 8% over the same period to $6.48 million in 2011.

- **Manufactured Products**: CONX currently manufactures and sells 52 diagnostic products worldwide to hospitals, clinical testing laboratories, universities, biotechnology and pharmaceutical companies and research institutions. In North America, Corgenix's labeled products are sold through a small internal sales force as well as through a number of independent sales representatives. Corgenix, through a wholly-owned subsidiary located in the U.K., had also been handling sales of these products overseas but in October 2010 transferred that responsibility to the ELITech Group, who now serves as the company's international master distributor and is Corgenix's largest shareholder. Revenue from Corgenix's manufactured products fell about 7% from 2010 to $6.03 million in fiscal 2011 and accounted for approximately 76% of Corgenix's total revenue.

- **OEM Products:** CONX sells some of their manufactured IVD products (mainly antiphospholipid and coagulation products) to other diagnostic companies for resale under their own brand names.

- **Contract Manufacturing:** CONX provides contract manufacturing services to other medical diagnostic companies. Contract manufacturing revenue fell almost 21% from 2010 to $356k in fiscal 2011 and accounted for approximately 4% of Corgenix's total revenue. This has substantially rebounded through the first nine months of 2012 and been a major catalyst to Corgenix's recent top-line growth.

- **Contract Development:** CONX provides contract development services to strategic partners including ELITech, Tulane University and the National Institutes of Health (NIH). Contract development revenue grew 87% from 2010 to $1.15 million in fiscal 2011 and accounted for approximately 14% of Corgenix's total revenue.

Corgenix's total revenue fell about 4% from 2010 to $7.94 million in fiscal 2011 with EPS decreasing from ($0.00) to ($0.01) over the same period. This trend is not expected to repeat itself in 2012, however. Corgenix's recent restructurings, highlighted by the transfer of their international sales operation and consummation of a related amended co-development agreement with ELITech, along with key additions to the U.S. sales force, new product launches and additional product development contracts are expected to help jump-start revenue growth in 2012. Management's most recent guidance (provided on the Q3 earnings call on May 14, 2012) is for revenue of approximately $9.1 million - this guidance is trimmed from the $9.7 million provided in February 2012, which was revised down from ~$10 million provided in December 2011. The most recent guidance still represents implied growth of ~15% from 2011.

FACILITIES

Corgenix is headquartered near Denver, CO where, along with its corporate offices, it maintains its research and development, manufacturing and distribution facilities in a 32k square foot building. Corgenix is ISO certified and all of the products are CE marked for sale in Europe.

Headquarters



Product Development



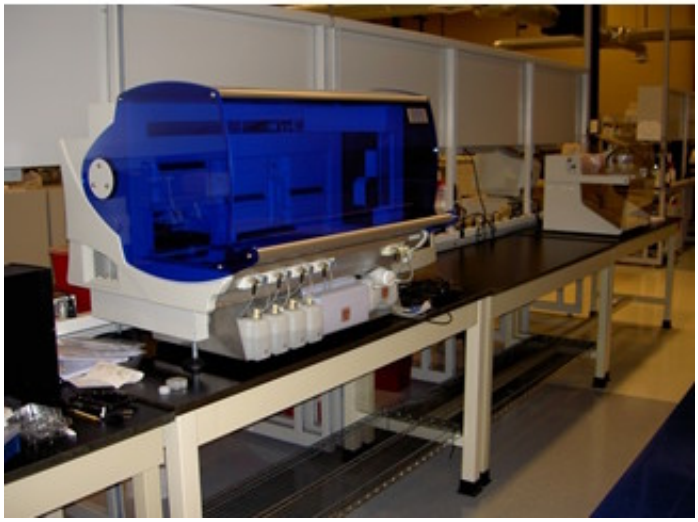
R&D and Production



Automated Filling Line



ELISA Instrumentation



Bulk Reagent Production



SOURCE: Corgenix Medical Corporation

PRODUCTS

The majority of Corgenix's product lines are developed on an ELISA testing format which are used by trained technicians in clinical labs for the diagnosis of autoimmune disorders, vascular disease and liver diseases. Testing is done with a standard 96-sample well microplate, allowing for automated high-throughput processing with quantitative (i.e. - accurate, non-subjective) results. (The following product and contract descriptions are per Corgenix's most recent 10-K).

- **Autoimmune Disease Products:** sales of autoimmune products fell about 25% from 2010 to $127k in fiscal 2011, representing approximately 2% of the company's total revenue. The products are used for the diagnosis and monitoring of autoimmune diseases, including rheumatoid arthritis, systemic lupus erythematosus, mixed connective tissue disease, Sjogren's Syndrome, dermatopolymyositis and scleroderma**.** The company's autoimmune product line includes tests for antinuclear antibodies (ANA) screening, dsDNA, Sm, SM/RNP, SSA, SSB, Jo-1, Scl-70, Histones, Centromere, Mitochondria, MPO, PR3, Thyroglobulin, LKM-1, anti Ribosomal P, BP-180, DSG-1, DSG-3, anti-polymer antibodies and thyroid peroxidase. Of Corgenix's 21 autoimmune products, two are manufactured in-house, with the remainder produced by other companies and sold through Corgenix's distribution network.

- **Vascular Disease Products:** Corgenix's vascular disease business, which includes antiphospholipid and coagulation products, is Corgenix's single largest source of revenue, accounting for almost 60% of the company's total sales in fiscal 2011. The vascular disease products are sold both as labeled (i.e. - "manufactured products") as well as OEM. OEM-related revenue accounts for approximately 18% of total coagulation and antiphospholipid sales. Corgenix expects OEM-related revenue to show a consistent decline as a percentage of total sales as the company shifts its focus towards higher growth products and segments.

  - **Antiphospholipid antibody testing:** sales of antiphospholipid products fell about 7% from 2010 to $3.39 million in fiscal 2011, representing approximately 43% of Corgenix's total revenue.

Antiphospholipid antibodies are associated with the presence of both venous and arterial thrombosis (clotting), thrombocytopenia (low platelet count that can result in bleeding), and recurrent miscarriage. These antibodies are frequently found in patients with systemic lupus erythematosis (SLE), and other autoimmune diseases, as well as in some individuals with no apparent previous underlying disease. They are also found in patients with antiphospholipid syndrome, an important medical condition with serious clinical manifestations such as chronic and recurrent venous (deep vein) thrombosis, as well as arterial thromboembolic disease, including heart attacks, strokes and pulmonary embolism. Thrombocytopenia has been attributed to the temporary removal of platelets from circulation during a thrombotic episode (clot formation). Corgenix manufactures and sells 11 antiphospholipid antibody testing products including aCL IgG, IgA, and IgM; anti-phosphatidylserine ("aPS") IgG, aPS IgA, aPS IgM; anti-ß2-Glycoprotein I ("aß2GPI") IgG, aß2GPI IgA, and aß2GPI IgM; and anti-Prothrombin ("aPT") IgG and IgM.

A potential significant bright spot for Corgenix's vascular disease business is with its "AtherOx" technology. AtherOx is being developed as a novel technology utilizing oxidized low density lipoprotein (oxLDL) in the assessment of cardiovascular disease, the nations number one killer. AtherOx development has been ongoing and the candidate recently entered clinical studies in the U.S. Corgenix now hopes to have study data available by the end of calendar 2012. The prior timeline was to hopefully to have an FDA submission prior to the end of the current year, although that may now not happen management. Nonetheless, management believes AtherOx could be a significant product if and when it reaches the market. As there is only limited information related to development and potential for eventual FDA approval we do not currently include any contribution from AtherOx in our financial model.

- **Bleeding / Clotting risk tests (coagulation):** sales of bleeding / clotting tests fell almost 15% from 2010 to $1.33 million in fiscal 2011, representing approximately 17% of Corgenix's total revenue.

  In people that suffer from excessive clotting or bleeding, Corgenix's tests are used to help find the source of the disorder such as von Willebrand's Disease (Hemophilia B). Corgenix's portfolio of bleeding / clotting risks tests include Protein C Antigen ELISA, Protein S Antigen ELISA, Monoclonal Free Protein S ELISA, von Willebrand Factor Antigen ELISA, von Willebrand Factor Activity Test; abp Ristocetin, and Collagen Binding Assay. Corgenix manufactures 5 of the 20 bleeding / clotting tests and markets the other 15 which are manufactured by other companies.

  We expect coagulation may be another area that holds future promise for Corgenix, especially as it relates to expansion into tests for other biological markers on ELITech's immunoturbidimetry testing platform.

- **AspirinWorks:** sales of the AspirinWorks test were $406k in fiscal 2011, representing approximately 5% of Corgenix's total revenue. The test is sold almost exclusively in North America. Sales of the test doubled from the $202k posted in fiscal 2010 and are on-track to double again in fiscal 2012 (to about $800k) as additional customers adopt the test and existing customers place more orders. AspirinWorks revenue was $564k in the first nine months of 2012, up 108% from the same period 2011. A 28% increase in reimbursement under existing CPT codes, which became effective in 2011, has also beeen a catalyst in driving demand for the test.

AspirinWorks kit



SOURCE: www.aspirinworks.com

Approximately 25% of people are resistant to the benefits of aspirin, a daily dose of which is often recommended to reduce the risk of heart attack or stroke. AspirinWorks is a urine test and the only FDA-cleared test to help doctors accurately determine aspirin's effectiveness by measuring the levels of thromboxane in urine. The test can be administered at the point of care (i.e. - doctor's office) with results available within minutes, a significant advantage over blood sample tests that must be evaluated in a clinical lab. The test allows physicians to adjust aspirin dosage or prescribe different therapy for those individuals that show aspirin resistance. The AspirinWorks test was developed in conjunction with biotechnology company Creative Clinical Concepts and biochemical research products manufacturer Cayman Chemical Company. Certain technology patents covering the test are owned by McMaster University which granted Corgenix exclusive rights to them.

Corgenix, in collaboration with Randox Laboratories, is also working on an immunoturbidimetry version (i.e. - automated version) of the test which would run on clinical lab analyzers, affording high volume throughput. Management has been shooting for regulatory approval and launch of test in Europe before the end of the current fiscal year (6/30/2012) but that may now be delayed. Following eventual launch, we expect this automated test to provide a strong complement in growing the AspirinWorks franchise.

- **Liver Disease test:** sales of the company's hyaluronic acid (HA), liver disease tests fell almost 19% from 2010 to $783k in fiscal 2011, representing approximately 10% of Corgenix's total revenue. The test is used to measure the levels of hyaluronic acid in the blood to help assess the degree of liver fibrosis and cirrhosis in chronic liver disease. We expect revenue from this business to turn around and show positive growth in fiscal 2012. Liver disease is another area where we think there is significant upside potential from new products via the ELITech development agreement and a segment which we model to grow at a 20%+ annual rate over the next several years.

R&D CONTRACTS / GRANTS

Corgenix's contract development business currently covers two main areas; government contracts and grants which are mainly involved in infectious diseases, and a joint-development agreement with ELITech. Revenue from this segment increased 87% in fiscal 2011 to $1.15 million. As R&D and grant revenue started to become more meaningful, during fiscal 2011 Corgenix began reclassifying related expenses from R&D expense to cost of goods sold (2011 10-K reflects this reclassification for 2011 as well as for 2010). R&D contracts / grants gross margin was 32% in 2011, compared to approximately 52% for the remainder of Corgenix's combined businesses.

- **Government Contracts / Grants:** These are typically one to three year contracts. Corgenix serves as a sub-contractor to Tulane University for several NIH funded grants and contracts related to development of diagnostics, vaccines and therapeutics for hemorrhagic fever viruses. Corgenix serves as the principal investigator for a $600k NIH funded two-year contract to develop recombinant diagnostic tests for the filoviruses (Ebola and Marburg), and has engaged three subcontractors (Tulane University, Autoimmune Technologies and the Scripps Research Institute) to assist in the development. Ebola and Marburg viruses are indigenous to Africa, cause hemorrhagic fever and can be fatal. Development of these tests are still in the early stages.

  The company, along with Tulane and other collaboration partners, are also working on development of Lassa virus test kits as part of an NIH grant. Lassa virus, spread by rodents and potentially fatal, is endemic in parts of Africa. The Lassa kits are currently being used in clinical trials in Africa, which could wrap up by calendar year-end 2012, followed by regulatory submissions. The regulatory process is expected to be relatively simple and commercial sales in Africa could potentially commence by early 2013. Corgenix has rights to manufacture and market these infectious disease tests upon gaining all the requisite approvals - upon commercialization, sales will flow through ELITech (international distributor). While sales of tests for infectious diseases related to developing countries may not offer much in the way of significant revenue for Corgenix, the company's capabilities in developing these tests should bode well for procuring additional government grants.

  In an effort to continue to grow this business, Corgenix regularly submits for additional government grants (related to both infectious diseases and bio-terrorism) and recently indicated that they are in discussions with a number of other potential R&D contract partners that could materialize in the future (no timeline was offered). We model this segment to grow at a low double-digit to mid-teens annual rate over the next several years as a result of new government contracts and the ELITech deal (described below).

- **ELITech Joint Product Development Agreement:** The ELITech Group (ELITech) is a worldwide manufacturer and distributor of IVD equipment, chemical analyzers and reagents. ELITech distributes diagnostic products for clinical chemistry, microbiology, immunology, and molecular biology through its own sales force as well as a distribution network to over 100 countries.

  ELITech also serves as Corgenix's international master distributor and is the company's largest shareholder. In 2010 Corgenix entered into a product co-development agreement with ELITech called the Joint Product Development Agreement. In July 2011 the agreement was amended and extended until July 2014 (with options

to extend it further). The current agreement has two phases; Phase I entails modification of Corgenix's existing assay technology for use in ELITech's analyzers (testing instruments) while Phase II is focused on the development of new Corgenix Immunoturbidimetry (IT) assays for use in ELITech analyzers.

Corgenix recently began beefing up their manufacturing capabilities to support production of these new assays. Corgenix will manufacture and sell the assays to ELITech (for sales overseas) and will also have the option of selling derivatives of the assays in North America. In mid-February 2012 Corgenix announced the first of the two products, an IT-format (automated) hyaluronic acid test, received CE Mark - the test will now be launched through ELITech. The other product has completed development and as of the Q1 2012 earnings call was expected to be CE Marked in the near term as well. Upon launch, CONX will announce exactly what this products is. Although we have no particular insight, we assume this product also launches before the current year end.

In conjunction with the original (2010) joint development agreement, Corgenix and ELITech (along with Wescor, a wholly owned subsidiary of ELITech) entered into a Common Stock Purchase agreement. Among other terms (which we detail later) Wescor is required to purchase 3.33 million shares of CONX's common stock for an aggregate $500k ($0.15 / share) upon the successful completion of two new Corgenix assays (related to the joint development agreement). Wescor will also receive a warrant to purchase 1.67 million CONX shares at $0.15 / share. Wescor bought the 3.33 million shares on September 15, 2011.

Corgenix expects to continue pumping out new products on a regular basis through their partnership with ELITech and has indicated several tests are under development and could reach commercialization in the not-too-distant future. We expect new products through this deal, coupled with ELITech's significant international distribution reach, to be a significant driver of Corgenix's revenue (including reviving sales of the vascular disease segment) over the next several years.

CONTRACT MANUFACTURING

CONX provides contract manufacturing services to other medical diagnostic companies. Their most significant contract manufacturing customer is BG Medicine, a Boston, MA based biotechnology company. While revenue from this segment can be somewhat irregular from period-to-period (i.e. - quarter-to-quarter as well as year-to-year) as a result of contracts churn, as this business has historically accounted for only a small portion of total revenue (~5% - 7%), this volatility has only a relatively minor impact on Corgenix's financial performance.

Contract manufacturing revenue significantly ramp through the first nine months of 2012, increasing by more than $700k from the same period in 2011 as result of an increase in business under existing accounts and commencement of activity under recently signed contracts. Corgenix is also in the midst of expanding their testing platforms (from almost exclusively ELISA to also focusing on immunoturbidimetry and lateral flow) which not only broadens their potential customer base, it also allows them to hold onto existing clients that switch between testing platforms - all of this is expected to benefit CONX's manufacturing business.

ELITech AGREEMENTS / INTERNATIONAL TRANSITION

In July 2010 Corgenix and ELITech / Wescor entered into an agreement calling for Wescor to purchase a stated value of CONX common stock on three separate occasions. Two of the three stock purchases were contingent on Corgenix meeting pre-specified milestones.

The first purchase came in July 2010 in the amount of $1.25 million for 8.33 million shares of common ($0.15 / share). Wescor also received warrants to purchase 4.17 million shares at a $0.15 strike. In conjunction with closing of this first tranche, the companies entered into a Master Distribution Agreement and the aforementioned Joint Product Development Agreement (the latter which we explained previously) - terms of the Master Distribution Agreement stated that closing of the second tranche was contingent on ELITech becoming Corgenix's exclusive international distributor.

The second tranche, an investment of $250k for 1.67 million shares (also included warrants for 833k shares @ $0.15 strike), closed in October 2010, commensurate with the transfer of Corgenix's international distribution activity (handled by Corgenix UK) over to ELITech. Corgenix took a $486k charge in fiscal 2011 related to the shuttering of Corgenix UK.

The third and final tranche, as discussed earlier, was for an investment of $500k for 3.33 million shares (along with warrants to purchase 1.67 million shares @ $0.15) and contingent upon the successful completion of two new Corgenix assays (per terms of the First Amended Joint Product Development Agreement). Wescor bought the 3.33 million shares on September 15, 2011.

FINANCIAL CONDITION

Corgenix exited Q3 with $1.0 million in cash and equivalents, up from $812k at 12/31/2011. The sequential increase in cash came from 245k inflow from operations, $110k used in investing activities and net inflow of $66k in financing activities. Excluding changes in working capital, operating cash flow was slightly positive for the nine months ending 3/12/2012. Debt remained relatively insignificant at quarter end.

VALUATION

Revenue
We model $9.1 million in revenue in fiscal 2012 and look for this to grow at a CAGR of about 15% through 2015, reflecting North America and International annual revenue growth of approximately 14% and 27%, respectively.

Income / EPS
We model net loss of approximately $419k (-$0.01 EPS) in 2012, roughly flat from 2011. This reflects our ~15% estimated growth in revenue which is offset by a contraction in gross margin related to a significant increase in lower-margin contract manufacturing revenue.

We expect Corgenix to gain leverage in operating expenses going forward and, coupled with strong revenue growth and incremental widening of gross margin, to result in EPS showing consistent year-over-year improvement. We look for EPS of $0.025 in 2015.

Our comparable cohort used to value CONX consists of several companies in the medical diagnostic testing space, some of which are considered direct competitors to Corgenix. Our valuation methodology uses the average of four

metrics; price-to-2015 estimated EPS, price-to-book value, price-to-sales (trailing 12 months), and enterprise value-to-sales (trailing 12 months).

Average valuation based on these four metrics is about $0.50/share which, based on CONX's current market price of approximately $0.12/share, indicates the stock is significantly undervalued. As such, we recommend accumulating the stock and are maintaining our Outperform rating on Corgenix.

Ticker	Est Long-term EPS growth	Implied 2015 P/E	Price / Book Value	Price / Sales (ttm)	EV / Sales (ttm)
IVD	NA	NA	1.9	1.2	1.1
DIA.MI	7.5%	8.7	NA	NA	2.6
OSUR	NA	NA	4.4	5.1	5.3
TRIB	14.6%	8.9	1.6	3.1	2.3
CEMI	48.0%	5.8	2.6	1.5	1.3
IDXX	12.7%	17.3	8.4	3.8	3.9
VIVO	16.4%	12.7	5.9	4.8	4.7
QDEL	1.5%	34.1	2.9	3.9	3.9
ABAX	13.1%	21.3	4.7	4.8	4.4
Average	**16.2%**	**15.5**	**4.0**	**3.5**	**3.3**

	Value using 2015 Comp P/E	Value using Comp Price/Book	Value using Comp Price/Sales	EV using Comp EV/Sales	Value using Comp EV/Sales	AVG
CONX	$0.39	$0.43	$0.62	$26.04	$0.58	**$0.50**

KEY PERSONNEL

Douglass Simpson, President/ CEO

Douglass T. Simpson has been the President of Corgenix since May 1998 and was elected a director in May 1998. Mr. Simpson joined Corgenix's operating subsidiary as Vice President of Business Development in 1992, was promoted to Vice President, General Manager in 1995, to Executive Vice President in 1996, to President in February 1998 and then to Chief Executive Officer in April 2006. Prior to joining Corgenix's operating subsidiary, he was a Managing Partner at Venture Marketing Group in Austin, Texas, a health care and biotechnology marketing firm, and in that capacity, served as a consultant to REAADS from 1990 until 1992. From 1984 to 1990 Mr. Simpson was employed by Kallestad Diagnostics, Inc. (now part of BioRad Laboratories, Inc.), one of the largest diagnostic companies in the world, where he served as Vice President of Marketing, in charge of all marketing and business development. Mr. Simpson holds B.S. and M.S. degrees in Biology and Chemistry from Lamar University in Beaumont, Texas.

William Critchfield, SVP Operations & Finance/CFO

William H. Critchfield has been Senior Vice President Operations and Finance and Chief Financial officer since April 2011, was the Senior Vice President Finance and Administration and Chief Financial Officer of the Company since April 2006, and was Vice President and Chief Financial Officer from December 2000 to April 2006. Prior to joining Corgenix, Mr. Critchfield was Executive Vice President and Chief Financial Officer of U.S. Medical, Inc., a Denver, Colorado based privately held distributor of new and used capital medical equipment. From May of 1994 through July of 1999, he served as President and Chief Financial Officer of W.L.C. Enterprises, Inc., a retail business holding company. From November 1991 to May 1994, Mr. Critchfield served as Executive Vice President and Chief Financial Officer of Air Methods Corporation, a publicly traded company which is the leading U.S. company in the air medical transportation industry and is the successor company to Cell Technology, Inc., a publicly traded biotechnology company, where he served in a similar capacity from 1987-1991. From 1986 through September 1987 he served as Vice President of Finance and Administration for Biostar Medical Products, Inc., a developer and manufacturer of diagnostic immunoassays. In the past, Mr. Critchfield also served as Vice President of Finance for Nuclear Pharmacy, Inc., formerly a publicly traded company and the world's largest chain of centralized radiopharmacies.
Mr. Critchfield is a certified public accountant in Colorado. He graduated magna cum laude from California State University-Northridge with a Bachelor of Science degree in Business Administration and Accounting.

Luis Lopez, M.D.

Luis R. Lopez, M.D., served as the Chief Executive Officer and Chairman of the Board of Directors of Corgenix from May 1998 until April 2006 when his title was changed to Chairman of the Board of Directors and Chief Medical Officer. In July 2009, Dr. Lopez stepped down from his role as Chairman. From 1987 to 1990, Dr. Lopez was Vice President of Clinical Affairs at BioStar Medical Products, Inc., a Boulder, Colorado diagnostic firm. From 1986 to 1987 he served as Research Associate with the Rheumatology Division of the University of Colorado Health Sciences Center, Denver, Colorado. From 1980 to 1986 he was Professor of Immunology at Cayetano Heredia University School of Medicine in Lima, Peru, during which time he also maintained a medical practice with the Allergy and Clinical Immunology group at Clinica Ricardo Palma in Lima. From 1978 to 1980 Dr. Lopez held a fellowship in Clinical Immunology at the University of Colorado Health Sciences Center. He received his M.D. degree in 1974 from Cayetano Heredia University School of Medicine in Lima, Peru. He is a clinical member of the American College of Rheumatology, and a corresponding member of the American Academy of Allergy, Asthma and Immunology. Dr. Lopez is licensed to practice medicine in Colorado, and is widely published in the areas of immunology and autoimmune disease.

Ann Steinbarger, SVP Sales & Marketing

Ann L. Steinbarger has been the Senior Vice President of Sales & Marketing since April 2011, was the Senior Vice President of Operations from April 2006 to April 2011 and was the Vice President of Sales and Marketing from May 1998 to April 2006. Ms. Steinbarger joined Corgenix's operating subsidiary in January 1996 as Vice President, Sales and Marketing with responsibility for its worldwide marketing and distribution strategies. Prior to joining Corgenix, Ms. Steinbarger was with Boehringer Mannheim Corporation, Indianapolis, Indiana, a $200 million IVD company. At Boehringer from 1976 to 1996, she served in a series of increasingly important sales management positions. Ms. Steinbarger holds a B.S. degree in Microbiology from Purdue University in West Lafayette, Indiana.

Taryn G. Reynolds, VP Facilities and IT

Taryn Reynolds has been a Vice President of Corgenix since May 1998. Mr. Reynolds joined Corgenix's operating subsidiary in 1992, serving first as Director of Administration, then as Managing Director, U.S. Operations. He has served as Vice President, Operations and in 1999, became Vice President, Facilities and Information Technology. Prior to joining Corgenix, Mr. Reynolds held executive positions at Brinker International, MJAR Corporation and M&S Incorporated, all property, operational and financial management firms.

FINANCIAL MODEL

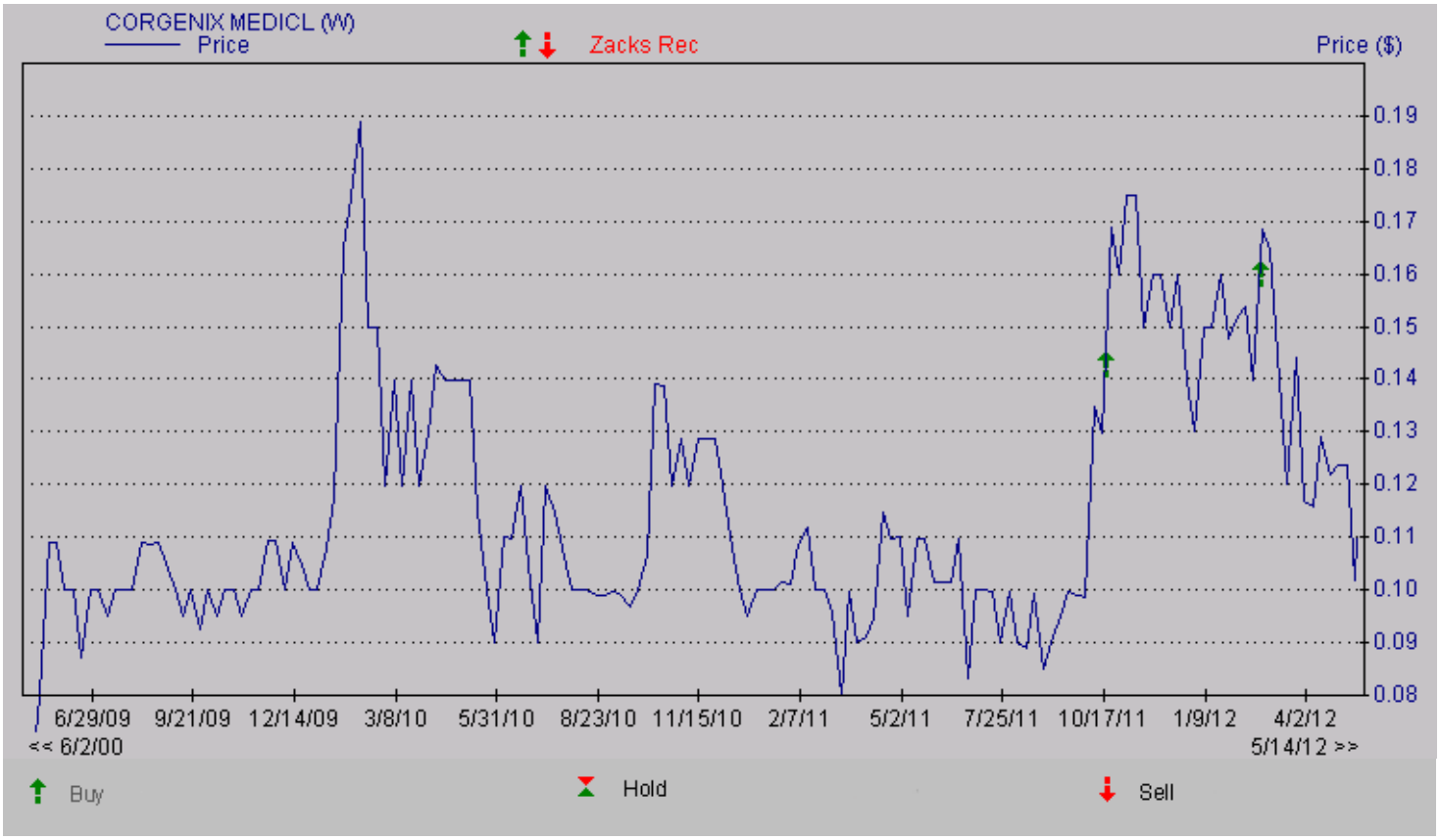
Corgenix Medical Corporation

	2011 A	Q1A	Q2A	Q3A	Q4E	2012 E	2013 E	2014 E	2015 E
North America sales	$6,483.2	$1,875.9	$1,733.9	$2,464.1	$1,883.0	$7,956.9	$9,397.1	$10,543.5	$11,640.0
y-o-y growth	*8.1%*	*28.1%*	*22.2%*	*58.5%*	*-7.9%*	*22.7%*	*18.1%*	*12.2%*	*10.4%*
International sales	$1,458.4	$302.3	$331.7	$253.3	$280.0	$1,167.3	$1,578.2	$1,947.5	$2,377.9
y-o-y growth	*-35.5%*	*-41.1%*	*-9.5%*	*-7.6%*	*-7.9%*	*-20.0%*	*35.2%*	*23.4%*	*22.1%*
Revenue	$7,941.6	$2,178.2	$2,065.6	$2,717.4	$2,163.0	$9,124.2	$10,975.3	$12,491.0	$14,018.0
YOY Growth	*-3.8%*	*10.1%*	*15.7%*	*48.6%*	*-7.9%*	*14.9%*	*20.3%*	*13.8%*	*12.2%*
Cost of Goods Sold	$4,012.4	$1,131.0	$1,128.1	$1,704.8	$1,233.9	$5,197.7	$6,281.3	$6,898.2	$7,386.8
Gross Income	$3,929.2	$1,047.2	$937.5	$1,012.6	$929.2	$3,926.5	$4,694.0	$5,592.8	$6,631.2
Gross Margin	49.5%	48.1%	45.4%	37.3%	43.0%	43.0%	42.8%	44.8%	47.3%
Sales / Mktg	$1,536.9	$509.3	$448.4	$510.5	$402.3	$1,870.6	$1,975.6	$2,223.4	$2,355.0
% S&M	*19.4%*	*23.4%*	*21.7%*	*18.2%*	*18.6%*	*20.5%*	*18.0%*	*17.8%*	*16.8%*
R&D	$194.5	$90.1	$75.2	$166.7	$90.0	$422.0	$355.0	$377.0	$402.0
% R&D	*2.4%*	*4.1%*	*3.6%*	*6.1%*	*4.2%*	*4.6%*	*3.2%*	*3.0%*	*2.9%*
G&A	$1,847.7	$424.9	$470.7	$505.3	$488.0	$1,888.8	$2,063.4	$2,123.5	$2,298.9
% G&A	*23.3%*	*19.5%*	*22.8%*	*18.6%*	*22.6%*	*20.7%*	*18.8%*	*17.0%*	*16.4%*
Disposal of assets	$486.1	$17.2	$0.0	$0.0	$0.0	$17.2	$0.0	$0.0	$0.0
Operating Income	($136.0)	$5.7	($56.7)	($169.9)	($51.2)	($272.1)	$300.1	$868.9	$1,575.2
Operating Margin	*-1.7%*	*0.3%*	*-2.7%*	*-6.3%*	*-2.4%*	*-3.0%*	*2.7%*	*7.0%*	*11.2%*
Total other income	($257.1)	($55.9)	($28.6)	($15.2)	($27.2)	($127.0)	($96.0)	($43.0)	$24.0
Pre-Tax Income	($393.1)	($50.2)	($85.4)	($185.1)	($78.4)	($399.1)	$204.1	$825.9	$1,599.2
Taxes (benefit)	$0.0	$0.0	$0.0	$0.0	$0.0	$0.0	$0.0	$0.0	$0.0
Tax Rate	*0.0%*	*0.0%*	*0.0%*	*0.0%*	*0.0%*	*0.0%*	*0.0%*	*0.0%*	*0.0%*
Preferred dividends	$21.5	$5.1	$5.1	$4.9	$5.1	$20.2	$0.0	$0.0	$0.0
Income to common	($414.6)	($55.3)	($90.4)	($190.1)	($83.5)	($419.3)	$204.1	$825.9	$1,599.2
Net Margin	*-5.2%*	*-2.5%*	*-4.4%*	*-7.0%*	*-3.9%*	*-4.6%*	*1.9%*	*6.6%*	*11.4%*
EPS	($0.01)	($0.00)	($0.00)	($0.00)	($0.00)	($0.01)	$0.00	$0.01	$0.02
YOY Growth	*674.1%*	*NA*	*NA*	*NA*	*NA*	*NA*	*-131.0%*	*304.7%*	*93.6%*
Diluted Shares O/S	40,153	41,517	45,047	45,997	46,000	44,640	70,000	70,000	70,000

Brian Marckx, CFA

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