



Second Quarter 2009 Interim Earnings Report

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Q2-09 Overview - Revenue

- Achieved \$22.3M in revenue:
 - Consolidated revenue below expectations due to the following factors:
 - Delayed purchases of in-vitro diagnostics capital equipment by our US customers approaching ~\$1.5M, resulting in approximately 25% decline in consolidated instruments revenue over prior year's quarter
 - Many of the expected domestic orders were in the final stages of the selling cycle but capital funding was not released
 - Increased rental pattern: ~\$500k in equipment rentals whose instrument's revenue will be deferred over future periods
 - \$400K lower than expected sales in our Sample Processing Division
 - Strong international demand for both iChem VELOCITY and iRICELLS
 - 72 iQ Analyzers
 - 50 iChem Velocity analyzers
 - 56% of iChem VELOCITY shipments were part of an iRICELL
 - Record Consumables and Service revenue: \$12.9M, 58% of total sales
 - 8% increase over prior year's quarter
 - 3% sequential revenue growth

The financial estimates disclosed on this presentation are preliminary and subject to external auditor's review and, therefore remain subject to change.

Other financial indicators for Q2-09

- Estimated Consolidated gross margins for the comparable second quarter remain at 51% despite significant lower instruments volume, severance costs and VELOCITY introduction and retrofit costs
- Estimated Consumables and Service gross margins improved to 59% from 58%, after severance charges related to a reduction in force implemented in June 2009 (61% on pro forma basis)
- Cash and cash equivalents improved to approximately \$29 million from \$27 million in Q1-09
- Financial results include severance and terminations cost of approximately \$325k

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Q2-09 Overview - Earnings

- GAAP Earnings for the quarter expected to be in the range of \$0.05 to \$0.07 including the following non-recurring costs
 - Severance and termination costs
 - iChem VELOCITY Retrofit Cost Provision
 - Impact: \$0.03

Pro-forma earnings are expected to be \$0.08 - \$0.10 versus \$0.11 in guidance primarily due to the domestic instruments revenue shortfall

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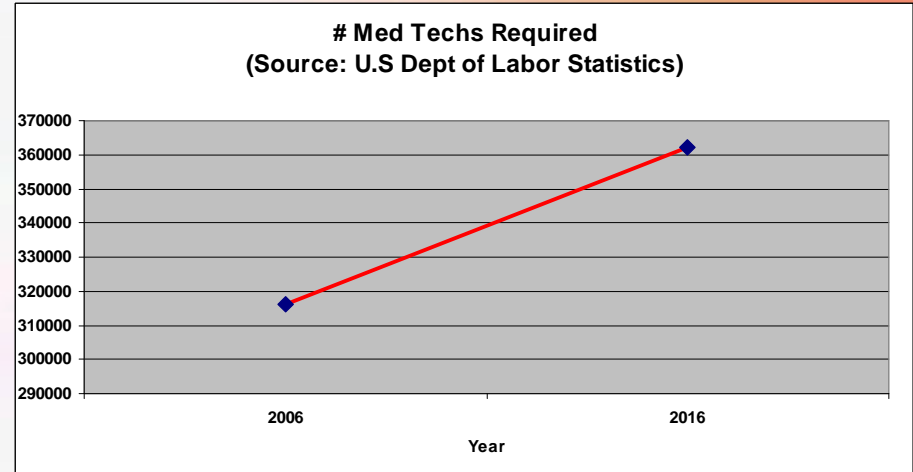
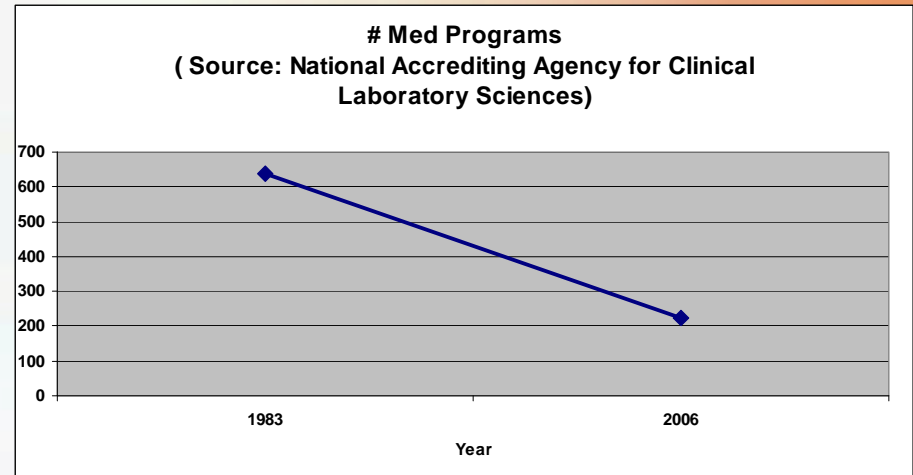
Q2-09 Overview – Action Plan

- Laid-off 19 full time equivalents which would generate \$450K net savings in the second half of 2009 and over \$1.0 million in 2010
- Increasing emphasis in domestic sales funnel to compensate for potentially lower close rate
- Implemented cost containment measures in all areas

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Demand for Lab Automation Should Continue and Help the Recovery of Instruments Sales in 2010

- According to the Wage & Vacancy survey by the American Society of Clinical Pathology, 50% of all laboratories in the US have difficulty hiring lab personnel
- Average lab worker age is 50+ years
- Baby boomer population is driving the demand for more and higher complexity lab tests in the US



Adapting to Challenging Market Conditions

- Providing multiple acquisition options for IRIS products
- Limited promotional discounts on high margin options
- Selective international promotions/ concessions
- Cost improvements and containment programs implemented

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2H-2009 Outlook

- Expect a stronger second half in revenue and earnings
 - Historical seasonality pattern: very strong fourth quarter
 - iChem VELOCITY learning curve should be over early 2H-09
 - Higher iChem VELOCITY consumable pull through in 2H-09
 - Higher R&D spending in the second half to support 3GEMS and NADiA clinical studies etc.
- Updated full year guidance to be provided July 30

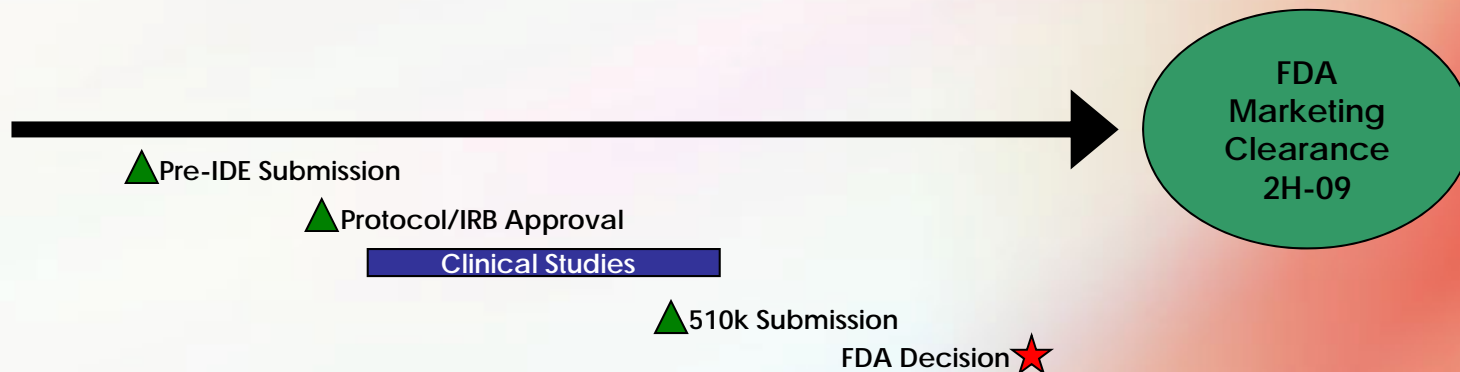
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iChem®VELOCITY™ Launch Status

- Successful re-launch of iChem Velocity in Q2-09
 - Customer satisfaction is high
 - Most distributors have re-initiated their full scale launch and field retrofit program
 - Shipped 50 analyzers bringing the total to approximately 150 units shipped since September 08
 - Beginning to experience the pull through of iChem test strips
- Pending 510(k) Clearance: IRIS responses are being reviewed by the FDA and an official response is expected imminently

NADiA ProsVue Status as of 07.21.09

- Adopted all FDA recommendations from their latest Pre-IDE review. Protocol and proposed claim are final.
- The clinical trials will be initiated as soon as ALL IRB's are approved by the following institutions:
- Duke University (Lead Investigators) IRB Aug 09
- Sloan Kettering IRB Aug 09
- Eastern Virginia Medical School – IRB approved
- University of Washington – IRB approved





Q & A