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International Institute for
Business Information & Growth LLC PRESENTS:

5th Annual GLOBAL CLINICAL OUTSOURCING SUMMIT

Optimizing Cross-Border
Partnerships & Expanding Opportunities
Using eClinical Technologies

December 6 - 7, 2011

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Adam R. Chasse, Vice President, Corporate Development,
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Scott Connor, VP Marketing, **ACURIAN**

Ed Corsi, Manager Clinical Trials, Clinical Development,
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Janet Flisak, Clinical Program Leader, Oncology,
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Joseph Gardner, Senior Scientist, **PFIZER**

Pedro Gittens, Senior Director, Compliance and Regulatory
Affairs, **LUX BIOSCIENCES**

Kathy Goin, Global Trial Director,
ENDO PHARMACEUTICALS

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Kara Lee McWatters, Professor, **HUMBER COLLEGE
SCHOOL OF HEALTH SCIENCES** & Clinical Research
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Michael G. Minor, Sr. VP Strategic Planning & Proposals,
ICON LATE PHASE & OUTCOMES RESEARCH

Dhiraj Pathak, PhD, Cloud Infrastructure Services Leader, **PwC**

Ross D. Pettit, Vice President, Clinical Operations and Data
Management, **AMAG PHARMACEUTICALS, INC.**

Stefan Proniuk, Senior Director Product Development,
ARNO THERAPEUTICS

Evette Riegel, Senior Director Clinical Operations/Project
Management, **BEARDSWORTH**

Donald P. Rosen MD, Co-founder, **RADPHARM, INC.** and
Director, **ACR IMAGE METRIX**

Barry Sagotsky, President, **MAGNOLIA LANE CONSULTING**

Dr. Uwe Schneider, VP, Compound Development & Branding;
Head of Global Sourcing Management, **GRÜNENTHAL GMBH**

Michael Semo, Sourcing Manager /Virtual CMC Team,
ELI LILLY AND COMPANY

Elizabeth Shewell, Senior Director, Outsourcing, **INCYTE**

Mollie Shields-Uehling, President & CEO, **SAFE-BIOPHARMA**

Ven Thangaraj, Founder & Advisor, **RADIANT SAGE LLC**

Adair Turner, Regulatory Affairs Manager, **MISSION3**

Bryant Wales VP, Strategic Sourcing, **IMCLONE SYSTEMS**

Jane A. Wilcox, Associate Clinical Contracts Director,
Clinical Development Operations & Biostatistics,
SHIRE DEVELOPMENT INC.

Bradley T. Wyman, Director, Imaging and Technologies,
Precision Medicine, **PFIZER**

A WARM WELCOME FROM YOUR CONFERENCE CHAIR

Dear Colleague,

I am pleased to invite you to attend **iBIG's 5th ANNUAL GLOBAL CLINICAL TRIAL PARTNERSHIPS: OPTIMIZING CROSS-BORDER OUTSOURCING**, scheduled for December 6-7, 2011 in East Rutherford, New Jersey. We have planned an executive-level event that will focus on the unique issues involved in outsourcing clinical research & life sciences manufacturing to non-US locales and on the regulatory, logistical and technology-related issues that are empowering global partnerships.

To address these issues, an outstanding speaking faculty of senior-level executives along with key operational leaders who handle global outsourcing and alliance management has been secured for this conference. You will hear from representatives of large, medium, and small pharmaceutical as well as biotechnology and medical device companies as they share their insights on strategies for optimizing clinical operations and outsourcing.

Key issues to be addressed, especially through case studies, include:

- How to effectively share operational risk with sponsors & or CROs and develop strategic planning for win-win partnerships
- Full service versus functional (niche) outsourcing: Strategies for Global Clinical Outsourcing
- Cloud computing concepts applied in pharma research and development
- Using the latest technologies for patient recruitment, compliance, and retention
- Managing relationships in over 10, 20 or more countries
- How to effectively find the right suppliers with the proper skill set & knowledge in emerging markets
- Successful strategies sponsors & CROs use to protect intellectual property rights
- Enhancing effective, accurate and timely collection of trial data
- Innovative ways to use technologies to lower drug development costs
- Study preparation meeting – No SOWs allowed
- Best practices for selecting CROs, suppliers and sites in BRIC countries and beyond

This intimate, highly focused, and informative event is only a day-and-a-half away from the office yet you will hear from and network with many of the industry's most innovative and strategic leaders, as well as receive the latest thinking on cross-border partnering strategies. Take away the tools and techniques to tackle outsourcing challenges that are unique to doing business in a global setting in the biopharmaceutical and medical device industry.

I look forward to seeing you December 6-7 in East Rutherford.

Best regards,

Lissa Blake

Lissa Blake, Conference Organizer
iBIG

CONTINUING EDUCATION CREDITS

Continuing Education Credits (e.g. CLE, CPE, CME, etc.) may be available for iBIG conferences. Upon request conference attendees will be provided with a "Certificate of Attendance" and a copy of the conference agenda, showing topics, length of sessions, and name(s) and professional affiliation(s) of presenter(s) for each session. Attendees wishing to apply for continuing education credits for attending this conference may submit this documentation to the relevant organization in his/her state when applying for such credits.



Program Level: Overview
No Advance Preparation Required
No Prerequisites Required

Program is a "Group Live" offering
CPE Credits awarded: 13.5

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8:00 – 9:15 Registration & Networking

9:15 – 9:30
Welcoming Remarks/Introductions

9:30 – 10:20
State of the Industry

As Pharma sponsors reduce domestic workforces, dependence on outsourced relationships throughout the life sciences food-chain will increase. Outsourcing and strategic partnerships have never been more imperative to drive down costs and reduce drug delivery time. With increasing pressure to achieve a greater level of accuracy and skill on both the Sponsor and Provider front, it's more important than ever to drive higher levels of joint effectiveness with clinical partners by establishing strong alliances early on. Contract research is almost at 20% of total drug development spending, and it continues to grow at a rapid pace. Strategic partnerships with CROs are what drive clinical trials at the very core of clinical development; and with billions of dollars spent on development each year, there is much opportunity for clinical partners to meet and maximize one another's expectations and goals moving forward.

PANELISTS:  Michael G. Minor, Sr. VP Strategic Planning & Proposals, **ICON LATE PHASE & OUTCOMES RESEARCH**

 Mary Lynne Kupchella, Esq., Associate Director, Outsourcing Lead, Contracts, Compliance and Vendor Management, **PFIZER, INC**


 Ed Corsi, Manager Clinical Trials, Clinical Development, **TALECRIS BIOTHERAPEUTICS**


10:20 – 11:05
Gaining Control of Clinical Imaging Data


This session will explore the benefits of utilizing cloud based technologies to manage medical imaging data related to clinical trials. Cloud computing is changing the way users interact with applications and systems. The challenges with medical imaging revolve around image transfer, image volumes, image display, image storage, and workflows especially as the systems and image data are held on the cloud. The session will provide details and use cases on image transfer, image archival, workflow, blinded reads and collaboration as well as provide a cost benefit analysis of using cloud based technologies.

- Clinical Image Management System (CIMS)/ Image EDC perspective from a customer
- CIMS from an Imaging corelab and Radiology perspective
- CIMS from a use case perspective
- Cloud technologies for drug discovery in a high volume and regulated environment.

MODERATOR:  Ven Thangaraj, Founder & Advisor, **RADIANT SAGE LLC**

PANELISTS:  Keith Elliston, PhD, Vice President, Systems Biology, **CHDI**

 Bradley T. Wyman, Director, Imaging and Technologies, Precision Medicine, **PFIZER**

 Dhiraj Pathak, PhD, Cloud Infrastructure Services Leader, **PwC**

 Donald P. Rosen MD, Co-founder, **RADPHARM, INC.** & Director, **ACR IMAGE METRIX**

11:05-11:35 Morning Break & Networking

11:35 – 12:20
Move into the Fast Lane with eTMF

We will be covering the following in the session:

- What is TMF?
- What is an eTMF?
- What is an eTMF system?
- Overview of the Trial Master File in Reference Model (RM)
- Benefits of eTMF based on the DIA RM to Stakeholders

MODERATOR:  Dirk Karsten Beth, President & CEO, **MISSION3**

PANELISTS:  Adair Turner, Regulatory Affairs Manager, **MISSION3**

 Pedro Gittens, Senior Directory, Compliance and Regulatory Affairs, **LUX BIOSCIENCES**

12:20 – 1:30 NETWORKING LUNCHEON

1:35 – 2:10
How to Effectively Share Operational Risk with CROs to Develop Win-Win Strategic Partnerships

The CRO competitive environment is evolving, as outsourcing demand from large pharmaceutical companies has picked up. Preferred partnerships between Big Pharma and CROs are becoming more common, with the benefits of increased efficiency, expanded developmental capabilities, and significant cost savings. Although partnerships have previously existed in various situations, the concept is only now becoming more prevalent throughout the industry. In addition to the substantial recurring revenue that partnership deals generate for CROs, any significant collaboration with Biopharma companies increases the likelihood of an eventual longer-term pact. CROs can concentrate on the various aspects of enterprise risk management, gaining appropriate insight across all functions where risks may threaten the enterprise.

MODERATOR:  Jane A. Wilcox, Associate Clinical Contracts Director, Clinical Development Operations & Biostatistics, **SHIRE DEVELOPMENT INC.**

PANELISTS:  Jason A. Gross, Vice President of Scientific and Medical Affairs, **CIPHER PHARMACEUTICALS INC.**

 Dr. Uwe Schneider, VP, Compound Development & Branding; Head of Global Sourcing Management, **GRÜNENTHAL GMBH**

 Bryant Wales VP, Strategic Sourcing, **IMCLONE SYSTEMS**

2:15 – 2:45 Refreshments & Networking

2:50 – 4:00

Study Preparation Meeting – No SOWs Allowed

- Bringing a global study team together to physically walk through processes – no assumptions
- A-Ha moments bring a team together
- Why we leave the Statement of Work at the door

SESSION LEADER:



Kathy Goin, MS, CCRA, Global Trial Director,
ENDO PHARMACEUTICALS

PANELISTS:



Kara Lee McWatters, Professor, **HUMBER COLLEGE SCHOOL OF HEALTH SCIENCES** & Clinical Research Consultant, **MCWATTERS CLINICAL RESEARCH CONSULTING INC.**



Lori Bainbridge, Director, Client Services,
JB ASHTIN



James Denmark, Founder & CEO,
MYCLIN

4:05 – 5:00

Finding the Right Suppliers: Identifying Proper Skill Sets & Knowledge-base in Emerging Markets

The choice of the right supplier is key and sometimes communication problems are an issue especially if emerging markets are taken into consideration. It is important to choose the right study geography and understand the different roles for emerging and established markets to roll out an outsourced study with more confidence. The CRO must have the right level of technical competence to deliver to the company's needs. The supplier must have the right equipment and/or facilities to deliver the project. And finally cost - the price has to be right.

MODERATOR:



Jacqueline Delfgaauw, PhD, Sr. Director, Head, Clinical Operations and Medical Monitoring US,
GRÜNENTHAL

PANELISTS:



Ross D. Pettit, Vice President, Clinical Operations,
AMAG PHARMACEUTICALS, INC.



Jacquie Mardell, Founder/Partner,
ANHVITA BIOPHARMA CONSULTING



5:00 – 6:30

EVENING NETWORKING RECEPTION:

Meet & Network with Your Fellow Outsourcing Colleagues

8:00 – 9:00

Networking & Registration

9:00 – 9:45

Managing Relationships in Over 10, 20 or More Countries

Pharmaceutical and biotechnology companies are increasingly attempting to expand the market for new drugs by applying for regulatory approvals in multiple countries simultaneously rather than sequentially as they have in the past. Many believe that contract research organizations with a global presence will continue to benefit from these trends.

PANELISTS:



Renu Jain, Ph.D, RAC, Program Director,
GRIFOLS INC.



Joseph Gardner, Senior Scientist,
PFIZER

9:50 – 10:10

Implementation of PLM Software to Enable Integrated Collaboration and Vendor Oversight for CMC Activities

PRESENTED BY:



Michael Semo, Sourcing Manager / Virtual CMC Team,
ELI LILLY AND COMPANY

10:10 – 10:30

Morning Break & Networking

10:30 – 11:30

Full Service vs. Functional (niche) Outsourcing: Strategies for Global Clinical Outsourcing

Clinical outsourcing strategies are innovative and offer companies options to drive their time to market and decrease costs. These strategies have come in various forms and have been uniquely structured based on the needs of the sponsor company and service provider. While no one strategy works the best for everyone, a different model is appropriate at different times. Times have changed from full-service outsourcing, where a sponsor engages with global contract research organizations (CROs) to complete the services of a full clinical trial, or functional outsourcing, where various service providers provide a specific clinical function. The latter is a common model employed amongst many sponsors where "bundling" functional outsourcing has also been popular. The logic of functional outsourcing has many facets that compel a sponsor to choose this model versus full-service outsourcing - "jack of all trades, expert of none" and/or "not having all eggs in one basket."

PANELISTS:



Evette Riegel, Senior Director Clinical Operations/Project Management, **BEARDSWORTH**



Elizabeth Shewell, Senior Director, Outsourcing,
INCYTE



Jessica C. Lee, MPH, Senior Director, Clinical Development, **INOVIO PHARMACEUTICALS, INC.**

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11:35 – 12:15

Using the Latest Technologies for Patient Recruitment, Compliance & Retention

Current recruitment methods simply aren't bringing enough patients into clinical studies. New processes and technologies integrated with traditional methods are crucial to accessing new patient populations and helping to secure engagement. Several organizations are making dramatic strides in developing approaches and systems that work for them. This will address successful strategies for engaging patients including similarities and differences around the globe. This session will also address how to harness social media and the internet to recruit potential ePatients; increasing patient compliance by using new technologies as an engagement tool; reducing recruitment time with online screening and Electronic Health Records and more.

PANELISTS:



Scott Connor, VP Marketing,
ACURIAN



Janet Flisak, Clinical Program Leader, Oncology, Global Clinical Operations, **JOHNSON & JOHNSON**



Tia Carmon, Patient Research Coordinator,
MEDIGUARD.ORG

12:15 – 1:20

Best Practices for Selecting CROs, Suppliers, and Sites in BRIC Countries and Beyond

Proximity, expertise, language, culture and pricing are all considered when selecting a CRO. A good CRO should function as an extension of the Client's development group. Expertise, ease of communication, language and culture are most easily found in a local CRO. How do you effectively overcome the challenges of selecting the appropriate countries, sites and investigators? What are the cost differentials for site selection? This session discusses the need to carefully examine country specific regulations for doing business in BRIC countries and identify the challenges of outsourcing in these regions and possible solutions. What are the successful strategies sponsors & CROs use to protect intellectual property rights? What are the latest lucrative partnerships and why did they do so well?

PANELISTS:



Jeffrey N. Klein, Assistant General Counsel,
APTALIS PHARMA



Stefan Proniuk, Ph.D., MBA, Sr. Director Product Development,
ARNO THERAPEUTICS



Nancy Cardone, Business Development Manager,
CONFLUENT TRANSLATIONS



Joanne Jiang, PhD, MBA, VP, Business Development and Project Management, Co-Founder, **FOUNTAIN MEDICAL DEVELOPMENT**

1:20 – 2:20 **NETWORKING LUNCHEON**

2:20 – 3:00

Cloud Computing Concepts Applied in Pharma R&D

This session will share lessons learned and approaches in applying cloud computing concepts in a pharmaceutical research and development environment. Specific real world use cases will be highlighted. There will be discussion on future opportunities as well as current challenges in reaching further adoption including strategic issues such as security, technical integration, platform readiness, cost and service support. Topics will include value of cloud computing studies, cloud vendors/providers, non cloud solutions that provide what the cloud does, performance benchmarking, security models, and tools and frameworks for data analysis.

PANELISTS:



Tim Immel, Chairman and CEO,
CLINVERSE



Richard Holland, Chief Business Officer,
EAGLE GENOMICS LTD.



Mollie Shields-Uehling, President and CEO,
SAFE-BIOPHARMA

WHO WILL ATTEND:

This conference is specifically designed for senior-level executives with titles and industry affiliations such as:

Heads, Executive Directors, Vice Presidents, Directors and Associate Directors in:

- Global Strategic Sourcing
- Outsourcing
- Clinical Operations
- Contract Management
- International Clinical Operations
- Risk Management
- Clinical Trial Projects
- Clinical Development
- Planning
- R & D
- Global Clinical Programs
- Alliance Management
- Corporate Counsel
- Government Affairs
- Project Management

Industries:

- Pharmaceutical
- Biotechnology
- Medical Device
- Contract Research Organizations (CROs)
- Contract Manufacturing Organizations (CMOs)
- Site Management Organizations (SMOs)
- Independent Review Boards (IRBs)

2011 KEY HIGHLIGHTS

- ✓ 90% NEW senior level speakers to present at the 5th Annual event— bringing NEW insights and NEW industry strategies with them
- ✓ In excess of 150 individual organizations were represented at this conference
- ✓ Over 45 leading industry experts on this event's speaker faculty
- ✓ Over 15 hours of content with 14 roundtable panel discussions
- ✓ Over 6 hrs of dedicated networking and face-to-face time with industry leaders
- ✓ Plus, networking breakfasts, breaks and drinks reception and an optional dinner for attendees

CORE FOCUS AREAS FOR 2011

- Applying cloud computing concepts in a pharmaceutical research and development environment
- Full Service vs. Functional (niche) Outsourcing
- Using the Latest Technologies for Patient Recruitment, Compliance & Retention
- Discover the latest risks and benefits of working in emerging markets in Asia, Central Eastern Europe and Latin America

WHY YOU NEED TO BE THERE

- ✓ Meet face-to-face with key pharma, biotech and medical device senior-level outsourcing professionals and decision-makers
- ✓ Obtain strategic operational plans of pharmaceutical and biotech sponsors who are conducting multi-regional clinical trials
- ✓ Attain competitive advantage of how to successfully implement cutting-edge outsourcing strategies for small, medium and large pharma
- ✓ Network with the best in the industry

VENUE INFORMATION:



SHERATON MEADOWLANDS HOTEL

2 Meadowlands Plaza
East Rutherford, NJ 07073
Phone: 201-896-0500
Fax: 201-896-9696
Web: www.sheraton.com/Meadowlands

HOW TO BOOK YOUR ROOM:

iiBIG has made arrangements with the Sheraton Meadowlands Hotel & Conference Center for a limited block of rooms for this conference at the discounted rate of only **\$155** per night. To take advantage of this discounted rate you must make your reservations no later than **Monday, November 6, 2011** as the discounted room block rate will expire on this date. When making your reservations, be sure to mention that you are a participant of the "iiBIG/ Global Clinical Trial Outsourcing Conference."

**TO RESERVE YOUR ROOM, CALL:
201-896-0500**

REGISTRATION TERMS & CONDITIONS:

Registration fee includes admission to all conference sessions; breakfasts, luncheons, refreshments, receptions; and one copy of the conference documentation workbook provided by iiBIG.

Cancellation of registration must be made in writing to iiBIG no later than 30 days before the conference. A \$199 service & handling fee will be retained for each cancellation. No refunds if cancellation is received less than 30 days before the conference; however, registration(s) may be transferred to another person by sending iiBIG the name and full contact details of the substitute registrant(s). In lieu of cancellation, registration(s) may be transferred to any other iiBIG conference held within twelve (12) months of the date of the original conference.

Conference attendees are solely responsible for costs to attend conference including – but not limited to – travel, lodging, and/or shipping and safe keeping of personal or business property. iiBIG encourages all conference participants to carry insurance to protect against the loss, damage or theft of any personal or business property.

DAY TWO WEDNESDAY, DECEMBER 7, 2011

3:00 – 3:20 Afternoon Break & Networking

3:20 – 4:00

How to Lower Drug Development Costs Through Effective Technology Use and Site Management

- Technology can be an important part of patient recruitment and retention
- EMR
- Social media
- Online screening
- Maintain patient compliance and engagement
- Human element can't be discounted – technology can IDENTIFY patients, but sites must manage studies well to fully leverage technology
- Recruitment planning (with contingencies)
- Metrics-based project management
- These skills are not found in abundance at most sites – how to remedy this?

PANELISTS:



Gretchen Goller, Senior Director, Therapeutic Expertise, Patient Access and Retention Services, **PRA INTERNATIONAL**



Adam R. Chasse, Vice President, Corporate Development, **RxTRIALS**

4:00 – 4:45

Site Activation: How to Get Beyond the Classic Pitfalls

Efficient site activation is vital to clinical trial success. With the various activities happening during site activation several challenges come into play again and again. It is essential to timely site activation to know what these typical challenges are, being able to work through them and avoiding them when possible. During this panel session, speakers will share their experiences specializing in site activation, and the best practices that have been implemented to handle these challenges of this wide-ranging process. What opportunities are being offered in managing global data for clinical trials; how should global pharmaceutical companies address the challenges and seize the opportunities from this increasingly important hub for clinical trials. How can pharmaceutical companies and sponsors stay abreast of clinical trial practices by improving efficiency and reducing cost in collecting and managing clinical data? How should CROs organize themselves to address sponsors' requirements and play the role of a strategic partner in reducing trial time and complying with regulatory requirements?

PANELISTS:



Heather Almonte, Associate Director, Clinical Operations, **OCTAPHARMA USA**



Richard Malloy, Assistant Director Quality Assurance, **VIOPHARMA**

4:45 – 5:15

Trust-Based Influencing to Create Collaborative Relationships

Critical to the success of any sponsor/CRO contract is the ability of the parties to work together as a team, to be open to each other's ideas, and to allow themselves to be influenced by the other party. Trust-based influencing has great potential for long-term success. We define "trust" as a willingness to be open to, and to take risks with, another individual over whom you have no direct control. As such, trust does not come easy; it takes time and effort. Trust-based influencing, then, requires three critical skills if it is to be effective:

- Being willing to listen to the other party and to acknowledge his/her issues as legitimate;
- Being clear about your own objectives and what you want from the interaction; and
- Being more interested in finding a solution that works for both parties than in "winning."

PANELISTS:



Ira Asherman, President, **ASHERMAN ASSOCIATES, INC.**



Barry Sagotsky, President, **MAGNOLIA LANE CONSULTING**

5:15

Conference Concludes: **Thanks for "Thinking iiBIG!"**

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Acurian is the global leader in clinical trial patient recruitment and retention solutions. Our predictive models, proven processes, and cutting-edge technology combine with years of experience to take the guesswork and uncertainty out of patient recruitment. Since 1998, Acurian has supported over 400 protocols for more than 60 companies worldwide.



Beardsworth – a full-service CRO specializing in business solutions for complex clinical trials with a particular expertise in oncology including newly emerging oncology vaccine technologies. Our experienced team of professionals averages 20+ years experience in clinical trial design, conduct, management, safety and reporting through FDA approval. Our patient-centric approach and cross-functional strength transform trial complexities into sound strategy and execution. As the founding partner of OncologyOne® – an alliance of best-in-class regional CROs, we provide global reach with local expertise for oncology trials. Celebrating 25 years in business, Beardsworth is a CCR registrant & designated Small Business. When you are looking for straight talk, accurate results, and a CRO partner invested in your project, Beardsworth provides the solutions – on time and within budget. Visit us at www.beardsworth.com.



Clinverse, Inc. provides the industry's first fully automated, global, SaaS-based financial management network for investigator payments. Our flagship product, ClinPay™, efficiently automates and facilitates the entire investigator payment process from contract execution through site payment. It provides complete transparency for Sponsor, CRO's, and Investigators offering near real time transaction and payment information for all involved in the network. We are integrated with various e-Clinical systems to receive study data which triggers payment milestones in ClinPay eliminating duplication of data entry. Clinverse's patent pending technology platform supports payments in 130 currencies, is fully customizable, and provides a very cost effective solution for investigator payments.



Confluent Translations' mission is to go beyond translation to honest understanding through ongoing dialog with clients to ensure success. We are ISO 9001:2008 certified dedicated to providing quality work in a timely manner. Our name reflects the company's customer centric philosophy by merging the client and Confluent in a fluid process to obtain a positive outcome. Good translation and cultural training can make everything a company does fall into place internationally. You, your end customers, partners, and employees will appreciate the joy of engaging with people, products and processes in their first language. That's the most respectful and profitable way to conduct business globally. "Reliable and responsive" are words we hear every day from our clients. Visit www.confluenttranslations.com or call Nancy Cardone at 412 539 1410 x 101 to make sure your message is clearly delivered.



Fountain Medical Development Corp. is a contract research organization (CRO) offering a full range of ICH GCP compliant clinical research services for multinational clients conducting clinical research in China. The management team of Fountain has decades of combined experience working with the world's leading CRO's and drug developers. In the Chinese CRO market, we fill the void of a service provider that balances high quality with moderate cost. Our extensive global experience in multiple therapeutic areas is unmatched by other local CRO's, and our lower operational cost allows us to pass significant savings on to our clients. www.fountain-med.com

- We have very strong regulatory team and clinical operation team in China.
- Our clinical operation team provide phase II-IV clinical trial service, operation in China, Hong Kong, Taiwan and Korea.
- We have our own central laboratory, and we already got the certificate by CAP (College of American of pathologist).
- We also have our own phase I center, the only one managed by CRO in China.
- Our management team have many years of experience working for leading CRO or Pharma companies in US.



Impact Corelab, Inc., headquartered in Sacramento, Calif., provides a comprehensive portfolio of clinical imaging corelab services, which combines a purpose-built cloud-based clinical image management platform with intensive radiological expertise to improve process transparency and control over complex imaging-based trials delivering significant time and cost savings. The company is a wholly owned venture of Radiological Associates of Sacramento, a premier provider of specialized radiological services founded in 1917, and Radiant Sage LLC, a leader in cloud-based software solutions for clinical trial image management. Impact Corelab can be found on the Internet at: www.impactcorelab.com



Mission3 is the premier Regulatory Information Management software provider for the Life Sciences industry. Our innovative software solutions help Life Sciences companies of all sizes handle their electronic data management, regulatory submission planning and publishing, product registration tracking, virtual data rooms and electronic trial master files. Mission3 recently added GlobalTMF, an eTMF product that allows clinical trial companies to significantly reduce the start-up cost and efficiently manage trials. GlobalTMF is an easy but powerful document management that encompasses everything related to a clinical trial document's lifecycle, which promotes greater compliance with your SOPs, improves process efficiency and drives control of study cost and quality.



Radiant Sage LLC provides on-demand, web-based Clinical Trial Image Management Solutions for organizations involved in drug discovery and research. Specifically designed for clinical trials, the company's solutions, Core-Lab-in-a-Box™ and RadVista Viewer, enable rapid trial start and increased overall efficiency of the collection, distribution, and processing of images within a few weeks and requiring no capital infrastructure investments. The Software-as-a-Service (SaaS) delivery model enables sponsors to retain control of their clinical trials by leveraging cost-effective, efficient and accurate image management solutions. Radiant Sage is headquartered in Boston, Mass. and can be found on the Internet at: www.radiantsage.com

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