From : **LSCP** Global Management Team
To : Global Customers & Friends
Subject : Substantial Competence Addition at LSCP: Dr. Peter Schiemann as new Associated Partner – for immediate release

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Uerikon/Zurich, Switzerland, January 28, 2015

The **LSCP** **L**ife **S**ciences **C**onsulting **P**artners Management Team is pleased to announce the adherence of a new Associated Partner in the context of our extended Information Management and eHealth transformation initiative to our global professional network for the Life Sciences Industry and Regulators.

Dr. Peter Schiemann, MBA

Specialties

Risk Management Expert in Clinical Trials and Drug Safety Processes. Experienced Project Manager and Management Consultant.

Summary of experience:

Peter brings proven skills of managing a wide range of complex Biopharma projects, from small to large, in multicultural contexts to our customers:

- Starting in research in 1992 while working on his PhD thesis, Peter was managing a research grant including a small team of lab technicians. In addition, he initiated the follow up grant for that particular research project.
- Peter left research to join Roche in Switzerland in 1996 to lead the environmental monitoring and cleaning validation group in the newly opened packaging plant.
- After successfully implementing the processes he moved on joining Roche's clinical supply group to lead the clinical packaging and labeling plant, streamlining the processes, upgraded the machinery park and established strategic collaborations with contractors in order to cope with the increasing workload.
- 2 ½ years in that position, he then joined PricewaterhouseCoopers' R&D Strategy consulting group, leading several projects with large and small Pharma/Biotech companies in Europe and the US helping them to establish more robust strategies and processes, e.g. supply chain, document management, R&D strategy and supporting organization, etc.
- Rejoined Roche US in 2003 as planning manager in the Clinical QA group, he established a risk based approach to planning QA activities and forged a strategic alliance with a CRO in auditing clinical trial centers.
- After moving back to Roche Switzerland end of 2004, Peter started the Quality Risk Management (QRM) project. In 2007 the successful implementation of QRM started, continued in 2008 and was completed in the major areas in 2009. Mid 2009 I was appointed Global Head of the Quality Risk Management Group in Clinical QA.
- He left Roche end of 2011 to co-found Widler & Schiemann Ltd, a Management Consulting firm specialized to support clients in clinical development.

He is based in the Basel/Brussels area.

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"The LSCP Global Management Team is very pleased to have Peter joining and becoming active member of our global group. As our customers start to move into large transformation programs and projects, he will be an important additional asset we can offer and which makes him an excellent additions to the Team!"

A very warm welcome to Peter!

For a personal/group contact or full CV:
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