Bridge Associates International LLC

"Partners for Quality and Manufacturing Excellence"

Christian-Peter Christiansen, Ph.D.

Managing Partner

Christian-Peter has more than 30 years of experience in pharmaceutical quality operations for global pharmaceutical firms. Peter has lead both site and regional/corporate quality organizations covering all aspects of Quality Operations, including quality control, quality assurance, microbiology, regulatory compliance, and quality support activities. Peter heads up the European office for Bridge Associates in Frankfurt, Germany.

Among Peter's areas of expertise are pharmaceutical analytical technology, quality operations management, and cost and efficiency optimization. Peter has published widely in the analytical arena, including chromatographic analysis, asbestos analysis, and most recently, process analytical technology (PAT). He is currently active in bringing PAT on-line in several major companies, and remains instrumental in the continuing dialog with regulatory agencies world-wide to implement PAT into the product releasing process.

Peter holds a doctoral degree in Pharmacy from the University of Marburg in Germany. He is a member of the Working Group for Production, Quality, and the Environment of Ethical Pharmaceutical Manufacturers (Verband forschender Arzneimittelhersteller (VfA) and a member of both the German and the European Pharmacopeial Commissions.

Curriculum Vitae

Experience since 2005

Managing Partner, Bridge Associates International, LLC
Princeton, NJ USA

Responsible for cGMP, manufacturing and management consulting, servicing global firms world-wide, specializing in Quality Control, PAT, Quality Operations Management.

2000 – 2004

Head of Quality Operations/Drug Products, Germany
Aventis Pharma Deutschland GmbH, Frankfurt1Main,
Germany

Accomplishments and responsibilities:

Excellent Management skills in big pharma quality operations. Experience with world-wide regulatory authorities, specializing in German authorities (BfArM).

Responsible for testing and release of Commercial Products dealing with quality control by chemical, physico-chemical and pharmaceutical analyses of bulk drug substances and finished dosage forms including evaluation of pharmaceutical technical complaints according to cGMP (KontrolI Leiter)

1995-1999

Head of Quality Operations Drug Products, Frankfurt
Hoechst AG -Hoechst Marion Roussel, FrankfurtfMain,
Germany

1991-1994

Group Leader, Biosynthetic Marketed Products,
Pharmaceutical Quality Control Department Hoechst AG, Frankfurt/Main, Germany

1977-1991

Laboratory Head, Finished Dosage Forms, Marketed Products,
Pharmaceutical Quality Control Department Hoechst AG, Frankfurt/Main, Germany

1975- 1977

Industrial Pharmacist, Finished Dosage Forms, marketed,
Pharmaceutical Quality Control Department, Hoechst AG, Frankfurt/Main, Germany

Education

Dr. rer. nat., Pharmacy,
University of Marburg, Marburg (Lahn), 1974

Certification as Pharmacist, Darmstadt, 1971

Pharmaceutical Examination,
University of Marburg, Marburg (Lahn), 1971

Graduate and undergraduate level studies in pharmacy,
University of Marburg, Marburg/Lahn, 1968- 1974

Publications: Available upon request.

LANGUAGES: German, native
English, fluent