Xavier Luria, MD

Independent consultant, he founded DDR (Drug Development and Regulation, www.ddrmedic.com), located in Barcelona and London, where he is the Chair and Senior Consultant.

He was Head of Safety and Efficacy of Medicines at the European Medicines Agency (EMA) during 2005-2012, and previously held leadership positions in the biopharmaceutical industry for more than eighteen years, covering international clinical development, pharmacovigilance, biostatistics and medical affairs.

In addition to Dr. Luria's specialty in internal medicine, and pharmaceutical medicine and biostatistics (University Autonomous Barcelona), he has developed expertise in several specific therapeutic areas, carried out postgraduate qualification in clinical pharmacology, drug development and regulation (Tufts University School of Medicine, Boston), and he is a recognized expert on regulatory systems and benefit-risk assessment (modelling, development and methodologies) and has been lecturer at various universities in USA and Europe.

Furthermore, he is module chair at an IFAPP-King's College Postgraduate Course and lecturer at several other academic institutions in Europe and USA.

He is currently member of the advisory board of several biopharmaceutical and medical devices companies and serves as consultant and regulatory service provider to many others in Europe, Japan, US, Australia, Israel and Latin America.