Curriculum Vitae

Name	**Gustavo Arroyo**
Current Job Title	Coordinator Clinical Research Unit, and Coordinator Scientific Committee at CASMU-IAMPP.
Department	
Date	2012 November 01.
Education and Qualifications	
Degree or Qualification	Medical Doctor Post graduate studies: 1) Internal Medicine with special dedication to Diabetes, 2) Clinical and Therapeutic Pharmacology, 3) Toxicology and Therapeutic
Name of Institution	School of Medicine
Date Obtained	1977 February 15 1)1980 June 15, 2) 1981 July 15, 3)1982 April 15
Degree or Qualification	1) Assistant Professor of Internal Medicine, 2) Assistant Professor of Toxicology and Therapeutic 3) Associate Professor of Pharmacology
Name of Institution	Medical University Hospital, Medical University Claeh
Date Obtained	1) 1980 March 01, 2) 1981 September 01
Positions held at Roche	
Job Title	Clinical Study Coordinator, Medical Director
Department/Site	Pharma Medical Departement
Start/End Dates	1999 April 01, 2012 October 01.
Positions relevant to role / GCP Experience held outside Roche	

Name of Company	Abbott Laboratories
Job Title	Medical Director
Start/End Dates	1991 May 01 / 1999 March 31
Name of Company	Squibb Farmacéutica Uruguaya
Job Title	Medical Marketing Director
Start/End Dates	1986 May 01 / 1989 May 01
Name of Company	Hoechst AG
Job Title	Medical Director, Marketing Director, Sales Force
Start/End Dates	1973 Januar 01 / 1986 May 03
Membership in Professional Societies or Organizations relevant to role / GCP	
Name of Organization	Foro Latinoamericano de Comités de Ética en la Investigación en Salud (FLACEIS)
Year Joined	2001 September 01
Role	Member
Name of Organization	Cámara de Especialidades Farmacéuticas y Afines (CEFA) Organización de las Normas Técnicas de Investigación Clínicas en Humanos para el Mercado Común del Sur (MERCOSUR)
Year Joined	1991 November 08
Role	Member
Additional Relevant Information	
Languages	Spanish, English, German
Publications	90 nationals and 12 internationals
Faculty Membership	

Other, as appropriate	Achievement Awards: Excellence in applaying protocol writing S.O.P., 1994 November, Chicago, U.S.A.
	Coordinator of 43 nationals and 17 internationals Clinical Phase III and IV studies
	Training courses: 139 nationals and 61 internationals: e.g: Global Drug Safety SOP's, Roche Basel, December, 2003, San Pablo, Brasil Clinical Investigator GCP Roche laboratories, July 2002, Montevideo, Uruguay AIMS, Roche Laboratories, 2001 February, Buenos Aires, Argentina Good Clinical Practice, Roche Laboratories, 2000 November, Mexico Good Clinical Practice in the Conduct of Clinical Trials Training Programm Abbott Laboratories 1995 August Chicago Pharmacology in Humans, Walter Reed Medical Center, 1993 March, Washington, U.S.A. Clinical Pharmacology, Inter American, 1986 November, Buenos Aires, Argentina Controlled Clinical Trials, Insitut National de la Santé et de la Recherche Medicale, 1988 June, Buenos Aires, Argentina Markon Training Course, 1983 November, London, England Pharma Development and Marketing, Hoechst AG, 1982-1984, Frankfurt Main, Germany: