

CV George Datukishvili



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Summary of Qualifications

Professional Experience

Pharmaceutical Enterprise "ABiPharm" the Qualified person. (QP)
The Representative of International Quality Consulting Inc. (Germany) in Caucasus
The Representative of "Klimaoprema"DD (Croatia) in South Caucasus GMP Consultant

Experience/qualifications

25-years experience in pharmaceutical industry;
13-years work experience in internal processes of quality audit;
19-years experience in GMP training;
13-years experience of implementing validation procedures at a pharmaceutical enterprise.

Professional Experience

Responsibilities and achievements	Project manager		
	2012- 2017	« GM Pharmaceuticals »	Georgia
	2013-2016	« Pharma Thech »	Armenia
	2016	«Hoffman La Roche»	Georgia
	2014-2015	« Riyad Pharm »	Azerbaijanian
	2017-2018	"National laboratory of genetics"	Georgia
	2018-2019	" New Hospitals"	Georgia

	<p>Duties include:</p> <ul style="list-style-type: none"> • Concept design of the new Enterprise, including quality control and microbiological laboratories. • Basic design and management of the implementation facilities under the GMP Requirements. • Implementation of the GMP standard in Georgia: GM Pharmaceuticals, ABiPharm; In Armenian companies «Pharma Tech,» «Medical Horizons,» Esco Pharm» • Implementation of GDP standard in Hoffman La Roche Georgia
	<p>Director</p> <p>2010-2012 Pharmaceutical Enterprise “AbiPharm” Tbilisi</p>
	<p>Executive Manager</p> <p>1999-2010 Pharmaceutical Enterprise “GMP” Tbilisi</p> <p>Duties Include:</p> <ul style="list-style-type: none"> • Monitoring the work of the staff at the Department of Production, Quality Control and Assurance, along with the work of the personnel at Quality Control Laboratory; • Organization and Management of logistic issues; • Staff qualification; • Supervising the process of self inspection. • Representative of Supreme Administration of Quality Control • Part of the commission working on the certification of ISO 9001:2000 standard relevance at the company

	<p>Production Manager</p> <p>1996-1999 Pharmaceutical Enterprise “Levicha-Geo”</p> <p>Duties Include:</p> <ul style="list-style-type: none"> • Member of the commission working on drafting and implementing the plan of company construction on the part of “Levicha-Geo”. • Being a production manager was accountable for production organization, setting up GMP, the issues related to production management, the coordination of writing documents, production planning, logistics; • Certification of GMP standard relevance (the state standard of the Czech Republic) and took over the responsibility for quality system and production relevance;
	<p>Head of the Pharmaceutical Enterprise</p> <p>1993-1996 “Gardis”</p> <p>Duties included:</p> <p>Tasks related to production Management and Logistics;</p>

Teaching Experience

<p>2001-present</p> <p>Duties included:</p> <p>Giving lectures concerning the system of Quality control.</p>	<p>Ltd “PSP”</p>	<p>Georgia</p>
<p>Duties included:</p> <p>Coordination of the initial training and periodic teaching, fostering qualification and professional growth of staff engaged in production and technological processes, control and quality assurance, the system of logistics.</p>	<p>Ltd “GMP”</p>	<p>Georgia</p>

Publications/Newspaper Articles

“Elaboration and Standardization of Immune Modulating Effect of Plant Composition”
“Creation and Pharmacologic Study of Complex Anti ulcer Herbal medicine”
“Anti ulcer and Antiphlogistic Activeness of Complex Herbal Drug “Polyplant K”
“Elaboration of Plant Collection for Treatment of the gastrointestinal Tract”
“Medication for treating Stomach Ulcer”
“Invention of a Medical Herbal Composition for treating Stomach Ulcer”
Effect of a New Antiulcerogistic Herbal Drug” (Tibetan Medicine; the State and Perspectives of the Research”)
“The study of the Extraction Process of Biologically Active Substances in plant composition comprising three ingredients” (Magazine “Pharmacia” #2, 1997 p.21)

Projects/Consulting/Coaching

ISO 9001-2000 IMPLEMENTATION	GM Pharmaceuticals, PSP, Caparol Georgia, Abipharm.
GMP IMPLEMENTATION AND CONSULTING	<ul style="list-style-type: none"> • Lechiva (czech republic); • GM pharmaceuticals (georgia); • ATM uzbekistan • Doridarmon-uzbekistan • Soprodin (algeria)-solid pharmaceutical dosage forms manufacturing plant • Biopoluse-batumi-georgia • Abipharm tbilisi-georgia • Pharma thech-armenia • Riyad pharm-azerbaijan

COACHING	<ul style="list-style-type: none"> • Georgian drug agency, ministry of healthcare of georgia • Gmp inspectors training for georgian drug agency • Tbilisi state medical university - masters degree program for pharmacists
AUDITS	GMP, GLP, GCP, ISO 9001-2000, ISO 19025, ISO 22000- HACCP

Education

Education	<p>UNITED STATES DEPARTMENT OF COMMERCE, SABBIT TECHNOLOGY COMMERCIALIZATION PHARMACEUTICALS</p> <p>25.01-23.02 2003</p> <p>Diploma of General Profile Pharmacist (with Honors)</p> <p>09. 1983 – 06. 1988 Tbilisi State University of Medicine, the Department of Pharmaceutics</p>
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Trainings

March, 1997	<i>“The management of Pharmaceutical Production-the Formation of Organizational Structure, Issues Related to Logistics”;</i> <i>The Czech Republic, Company “Lechiva”;</i>
October, 1997	<i>“The System of Quality Assurance; Storage Economy”</i> <i>The Czech Republic, Company “Lechiva”;</i>
March, 1998	<i>“Production Planning, the organization and management of Production Processes”;</i> <i>The Czech Republic, Company “Lechiva”</i>
11.04.- 20.04. 2001	<i>“GMP Requirements as the basis of Quality Assurance for Medicinal Products”</i> Educational Centre “State Scientific Centre for Antibiotics”;
11. 03. - 16 03. 2002	<i>“The Quality Control of drugs in accordance with GMP Standards”;</i> Educational Centre “State Scientific Center for Antibiotics”;
18. 03. – 23. 03. 2002	<i>“The Organization and Management of Production Processes In Line with GMP Standards”, “Normative and Technological Documents”;</i> Educational Centre “State Scientific Center for Antibiotics”

02-06.09. 2002	<i>“Quality Control; Toll Manufacturing and Analysis; Complaints and Recalls; Self Inspections; Validation Water treatment Systems; Cleanroom Construction and Validation; Sterile Manufacturing; Active Pharmaceutical Ingredients; Inspection and Audit”.</i> Heathside Information Services Ltd.
25.01. – 22.02. 2003	United States Department of Commerce, SABIT Technology Commercialization: Pharmaceuticals.
12. 07. – 16.07.2004	<i>“The Self Inspection Planning and Management within the Pharmaceutical Enterprise”.</i> “SKIF”

Languages

Georgian (Native)

Russian (Fluently)

English (Advanced)

Skills

Driving License

Interests

Tourism, folk music, wine industry