Peponaki Charoula
Pharmacist, MSc

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PROFESSIONAL EXPERIENCE

Pharmathen SA

Technology Transfer Director May 2018 – present

Key Responsibilities:

- Leads Supply Chain Development Industrialization activities ensuring GMP compliance and to facilitating the timely release of product.
- Coordinates plant issues and tasks with respect to scaling up processes, new equipment and procedures, equipment and process validation, manufacturing of development, stability, and product validation batches.
- Coordinates the preparation of the manufacturing site for the manufacture of the new products.
- Communicates as necessary with the various manufacturing plants, other departments, project team members.
- Coordinates and monitors technology transfer of process formulation from lab to (pilot) plant in coordination with R&D team
- Seeks and negotiates for new alliances with contract manufacturers when necessary.

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Head of Product Industrialization, 2006 –April 2018

Key Responsibilities:

- Leads Supply Chain Development Industrialization activities ensuring GMP compliance and to facilitating the timely release of product.
- Coordinates plant issues and tasks with respect to scaling up processes, new equipment and procedures, equipment and process validation, manufacturing of development, stability, and product validation batches.
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Pharmanel SA

Production Manager & Plant Director, 1995 –2006

Key Responsibilities:

- Providing leadership for the manufacturing plant.
- Maintains and coordinates day to day finished drug product manufacturing and packaging activities to support the growth of company's Manufacturing with increased drug product lines and multiple shift operations.

Chrispa Pharmaceuticals
QC Manager, 1994 - 1995

Key Responsibilities:

- Follows Company's Quality Policy and complies with current operational standards and legislative requirements Provides guidance to all involved departments of the facility.
- Follows Company's Quality Policy and complies with current operational standards and legislative requirements.
- Follows the rules of hygiene and control of health status as defined in the relevant Policies and Standard Operating Procedures
- Performs all QP duties as provided by the Decision of the President of the Management Board of EOF.
- Evaluates analytical data of APIs, Excipients, packaging materials and finished products.

- Ensures that all necessary testing is appropriately performed.
- Approves specifications, sampling instructions, control methods and other quality control procedures.

Chemica Pharmaceuticals
RA / QA Officer, 1990 - 1994

Key Responsibilities:

- Responsible for submitting applications for marketing authorizations and for lifecycle management of the MAs (renewals/variations).
- Acting as a liaison person with Greek regulatory authorities/clients
- Assuring compliance to the GMP standards and ISO requirements

EDUCATION

• Degrees:	MSc in Industrial Pharmacy Athens University	(2002)
	BSc in Pharmacy Aristotle University Thessaloniki	(1988)
• Foreign Languages :	English (Fluently)	
• Computer Skills :	Microsoft Office, SAP	

PATENS

EP149 1199B1 Pharmaceutical compositions of alendronate sodium trihydrate and process for the reparation thereof