As a regulatory consultant NUPUR SHARMA combines knowledge of quality and regulatory requirements with interest in the medical device industry. She has experience of working in regulatory analysis, 510k, classification rationale and product registrations in various markets. She has experience in managing client relationships, needs assessments and mapping. NUPUR's approach is to ensure that solutions support the strategy while it is transmitted into practical and workable solutions.

NUPUR´s key roles include:

- Collecting data from multiple sources on consumers, competitors and the marketplace.
- Consolidating information into actionable items, reports, and presentations.
- Compiling and analyzing bulky data using modern and traditional methods to collect it.
- Perform valid and reliable market research SWOT analysis
- Provide competitive analysis on various companies, identify market trends and pricing/business models.
- Strong communication and presentation skills
- Understand the client requirements and provide the deliverables on time.
- Well acquainted with regulatory authorities like USFDA, CDSCO, EMEA, Health Canada, TGA.

QUALIFICATIONS

Quality and Regulatory Affairs for medical devices

Tools & Methods

- Scientific Research – analysis and data management
- Literature search like Pubmed, Clinical trials.gov
- 510 (k) database

LANGUAGES

- Swedish – basic
- English – fluent
- Hindi – mother tongue - fluent

EDUCATION

- Bachelor in Pharmacy
- Master in Pharmacology, Chandigarh College of Pharmacy, Punjab

Courses & certificates

- MDR training, SIS

EMPLOYMENTS

- Mediteq Svenkebo AB, 2022
- Fluicell AB, Scientific Sales Specialist, 2019-2022
- SME, Lifescience & medical devices, Cheers Interactive, 2016-2018
- Product Executive, Symbiotic Drug & Diabetic care, 2016

Mediteq
Kanalstråket 1
SE-433 76 Jonsered,
Sweden

+46 31 774 25 00
+46 708 72 96 68
helen@mediteq.se
www.mediteq.se

+46 72 875 04 91
nupur@mediteq.se