

Piramal Imaging GmbH (PI) belongs to the global Piramal Healthcare Group and is located in Berlin, Germany. Our vision is to build our position as an innovator and leading player in the Molecular Imaging field and to be perceived as a competent specialist and reliable partner in R&D, Product Supply and Marketing of PET tracers. We aim to create value for patients & physicians by improving early detection and characterization of chronic and life threatening diseases leading to better therapeutic outcomes and improved quality of life.

For our most advanced PET tracer florbetaben and our strong pipeline of tracers in clinical development we offer the following full-time position with immediate effect:

## CMC Operations Manager EU (m/f)

The CMC Operations Manager will report to the Head of Global Manufacturing. The position will be located in Berlin – the headquarters of Piramal Imaging GmbH.

## Main Tasks & Responsibilities:

- Setting up GMP- production (synthesis, quality control and release) at several Eurpoean contract manufacturing partner sites.
- Development of additional strategic partnerships in the EU.
- Continous monitoring and improvement of the production with regard to GMP compliance and performance.
- Preparation and coordination of clinical trial supply.
- Writing and compilation of submission relevant CMC-documents.
- Technical support for quality site audits.

## Skills:

- Ability to work effectively in a team environment
- Attention to detail and quality orientation
- Self-starter with a high degree of endurance, flexibility and mobility with respect
- Open-minded and motivating attitude
- Computer skills, in particular Word and Excel
- Fluency in English. Command of the German or French language would be a plus.

## Experience:

- University or academic degree in chemistry or pharmacy ideally complemented by
- Experience in working under GMP-standards.
- At least 5 8 years of experience in production processes of pharmaceuticals, preferably of parenteral drug products and in "contract manufacturing".



- Minimum of 5 years of experience in (clinical and/or manufacturing) quality assurance in a pharmaceutical international environment
- Experience especially in GCP and GMP matters and working knowledge of EU and/or FDA regulatory requirements
- Experience in conducting audits

If you want to be part of an entrepreneurial team and are prepared to assume a wide range of responsibilities and tasks, please submit your application documents electronically to:

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www.piramalenterprises.com

We are looking forward to receiving your meaningful application!