CollPlant Biotechnologies

WE'RE HIRING!

RA/QA Associate

CollPlant Biotechnologies is developing technologies in the fields of regenerative medicine, 3D bioprinting of tissues and organs and medical aesthetics.
We are looking for a talented and highly motivated RA/QA Associate to join CollPlant.

Job description

- Ensure QMS compliance to EU Medical Devices Regulation (MDR) requirements
- Prepare product Technical Files for MDR certification
- Monitor new developments in QA/RA e.g. standards and regulations, assess impact, plan and complete mitigating actions
- Deliver training on the QMS to existing and new staff
- Assist in Document Review and Control
- Perform Internal Audits
- Investigate process failure and implement CAPA
- Provide required QMS training

Qualification requirements

- Minimum 2 years working in the Medical Device Industry with experience in Class IIb, III implantable devices
- Working within an ISO 13485 QMS under the Medical Devices Directive (MDD)/ (MDR) is essential
- Knowledge and experience of the medical device CE marking process and MDD/MDR is essential
- BS.c and above degree in sciences
- High English skills (writing & speaking)

Join us: careers@collplant.com

CollPlant Biotechnologies Pioneering Regenerative Medicine