

Technolas Perfect Vision GmbH is one of the world's leading global ophthalmic companies, which focuses exclusively on refractive surgery. Our advanced and innovative products and solutions define the future of the refractive industry for the benefit of ophthalmologists and patients.



For our Headquarter in **Munich** we are looking for a

Specialist Regulatory Affairs (f/m)

As a member of our Global Regulatory Affairs Department you will be responsible for:

- National as well as international (mainly EMEA) product registrations
- Supervision of external consultants
- Contact person for Notified Bodies and Competent Authorities regarding product approvals
- Development of approval strategy
- Responsible for the "Technical File" of our products according to MDD93/42 EEC
- Risk Management according to ISO 14971

Skills and Experience we are looking for:

- More than 2 years experience in regulatory affairs of medical devices
- Fluent in English
- Dipl.-Ing. FH/University or equivalent

If you feel that you have what it takes to be successful in this role, we look forward to your application to job@technolaspv.com

TECHNOLASTM
P E R F E C T V I S I O N

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