



QA/QC specialist

The QA/QC Department in Sandoz is looking for an independent QA/QC specialist to our Nordic Head Office located in Copenhagen.

With your natural drive and will to cooperate and change, you get the opportunity to influence the future development of the Nordic team. The QA/QC department is responsible for all quality control and quality assurance tasks are handled according to GMP and to Novartis quality standards in the Nordic Area.

Your primary responsibilities

You will act as QA/QC specialist in the QA/QC Department on existing and future products, and your primary responsibilities and tasks are:

- Maintenance of the stability program including review of stability data
- Product Quality Review including Risk Assessment.
- Elaboration and maintenance of specifications and methods
- Performing audits of contract takers and API suppliers.
- Keeping updated with the European Pharmacopoeia and related publications.
- Release products as QP

Your primary professional qualifications

- It is mandatory that you have a Masters degree in Pharmacy or other Life Sciences
- 3-5 years experience from working in the pharmaceutical industry
- Preferably work as Qualified Person
- Excellent skills in English – both written and verbal

Your primary personal qualifications

- Responsibility and accountability comes natural to you
- Robust analytical skills with “completer/finisher” mindset
- You approach life with a look out for opportunities rather than limitations
- You work style is efficient and focused, and you remain calm and in control even when you are busy

Our offer to you

A challenging job in an international organisation, with a high degree of speed and independence. You will work both as a generalist and as an expert with a high degree of professional competency and responsibility. You will get lots of opportunities to grow your professional and personal skill base through a variety of independent and team based tasks.

For more information please contact QC/QC Manager Nordic, Marianne Schwarz on telephone +45 40231919 or Head of Regulatory Affairs, Nordics Malene Metz Moerch on telephone +45 213 697.

Apply for the position on the following e-mail: job.sandoz@sandoz.com

- Deadline for applying for the position is January 31, 2010.

Sandoz is a world leading manufacturer of generic pharmaceuticals and is part of the Novartis Group. Sandoz in the Nordics employs 70 people at our Nordic Headquarter in Copenhagen. It is Sandoz's vision to develop and manufacture non-patent protected high quality pharmaceuticals, offering them to patients and society at affordable prices, and thereby making health accessible to all.