Galenicum was founded in Barcelona in 2003 with the mission of improving the quality and affordability of pharmaceutical products. This purpose has remained at the core through all these years, while the company has grown to become a global ecosystem of several businesses within the pharmaceutical, food and nutritional supplement sectors. Galenicum covers the entire industry value chain and has commercial agreements in place that span more than sixty countries worldwide. Galenicum continues its adventure to become a specialty global pharmaceutical company, bringing further innovation to products in the years to come. Our tagline Believe in life, expresses the spirit of overcoming we aim to bring across all patients, as well as our team. All our effort is thanks to our support of a highly committed qualified team.

Do you want to learn more? Visit Galenicum at https://www.galenicum.com/en/

Join us!

We are looking for a passionated and committed trainee who wants to develop his/her career in the generic drugs industry inside our R&D Team in San Agustín de Guadalix, Madrid. He/She will be in charge of following responsibilities:

Roles and responsibilities:

- Coordination with CMOs for the transfer of the developed products on an industrial scale.
- Prepare and review development protocols, reports and manufacturing documentation.
- Participation in the answer of deficiency letters, dossier audits and clients/collaborators queries received for own developed products.
- Collaborate in troubleshooting and optimization of developed products and coordinate any technical problem related with the technology transfer activity.
- Participation in the regulatory strategy of the finished products. Processing quality module of the registration dossier.

Suitable Skills:

- Self-sufficient, critical, disciplined, hardworking, creative, motivated, proactive, dynamic, methodical, organized and decisive.

- Strong analytical and problem-solving skills.
- Excellent organizational, planning and time management skills in order to design, monitor and enforce work plans.
- Highly responsible and capable to work under pressure to meet deadlines.
- Ability to prioritize multiple tasks and execute projects on time in a fast-paced environment.
- Good team player and collaborative working style.
- Detail oriented professional with good communication skills.
- Supportive and able to build effective working relationships. throughout the organization.
- Excellent written and oral communication skills.
- Ability to increase technical knowledge and apply new skills.

Education:

- High level of search and comprehension of scientific documentation.
- Experience MS Office.
- Ability to use and generate documentation in English.
- Life sciences degree and master.

Previous Experience:

- No previous experience required.

Please send your CV at: recruiting@galenicum.com