

Driven by our Promise

CSL Behring in Marburg



GGL Career Day 2017

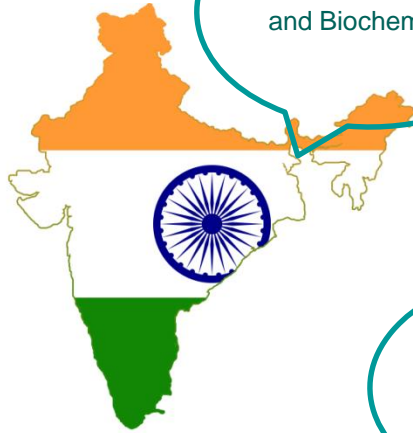
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CSL Behring

Content

- Myself
- What do I do whole day?
- Pro's and Con's

Me in short



Studied Biology
and Biochemistry

December 2014
Started working in
CSL Behring,
Marburg



2006 came to
Giessen for PhD

Postdoc in MPI,
Bad Nauheim



What do I do on a normal day?



- I plan and coordinate Stability Studies according to ICH Guidelines.
- Analyse the outcome of the studies.
- Prepare study reports and also documents for the registration of our products.
- Work in various project teams to implement new products or changes in manufacturing
- Manage a team of 3 colleague (Guiding them regarding their work as well as help them in their development)
- **Most importantly making sure that patients out there can trust Our Products**

What do I do on special days?

- I also prepare and participate in Audits (FDA, PMDA etc.)
- Work in deviation investigations
- Answer the Stability relevant questions from Regulatory Authorities
- Responsible for Climatic Chambers for Products
- Writing SOP
- Accompanying Change Control procedures



My group members (at least some of them)



Why work in the Pharmaceutical Industry

Pro's	Con's
Provides a structured career, training and opportunities for self-development and advancement	Unstable organisations with frequent reorganisations and periods of uncertainty
Competitive salaries and good terms of employment	Decreasing job security and fear of layoffs while companies strive to readjust
Increasing world-wide demand for new medicines and treatments	Highly competitive and often high levels of stress
Cutting edge of research with state-of-art facilities and opportunity to impact people's lives and society	Decisions are commercially- driven, not scientific. Research freedom perceived as less than in academia
Close teamwork and flexible working environment	R&D is long-term and most drugs fail long before getting to market