

WE NEED MORE OZACKIANS!

Are you **open-minded** and **brave enough** to navigate the wild waters through all stages of **drug development**? Do you want to make a real impact and learn something new **every day**? Then we have a unique opportunity for you!



Ann Christine
Korsgaard

Ozack is expanding to meet the urgent need for experienced regulatory affairs experts (5+ years) in small biotech companies that we collaborate with.

We are a team of five motivated, supportive, and highly engaged colleagues who aim to **make a difference every day** we go to work. No working day is the same as we are working with many different disease areas at various stages of drug development.

Work-life at Ozack is always colorful and we offer:

- Creating value through interaction with the **regulatory agencies**
- **Freedom** yet **commitment**
- **Sharing** expertise and **learning** new things each day
- Exciting mix of projects and clients
- Work **when and where you want** and yet belong to a small team
- Support from bright **young talented colleagues**
- Guaranteed monthly base salary, performance salary bonus

Your role involves:

- Being **responsible** for defining **regulatory strategies** for both early development and late-stage projects and preparing **scientific advice meeting packages** for both EU, US and other countries.
- Planning, compiling and submission of **applications for clinical trials and marketing authorizations in EU, US and other countries** and working closely with our customers to **support** any regulatory information needed.
- Defining possibility for **expedited opportunities** as Orphan Drug Designation, PRIME, Fast Track, Breakthrough Designation, etc., and take part in **achieving** these designations.

Confused? You won't be after a chat with us.

Contact me **+45 25403000** or **ack@ozack.dk** if you find the Ozackian life intriguing.

