Curriculum Vitae

Christina Balslev Rindshøj

Personal data
Born: 1972

Address
Bagsværd Hovedgade 163,
2. lejl. 2, 2880 Bagsværd
Tel: 23370572
E-mail:
christina.rindshoej@live.dk

Professional competencies

- Regulatory expertise
- Drug development excellence
- Organizational change management
- Business understanding
- Leading High Performing teams

Spare time

- Family
- Badminton
- Long walks in nature
- Building summer house in Rørvig

Global regulatory expert passionate to bring science and business together

- My continued excitement to work within the field of regulatory affairs after 20 years of experience is the fact that strategic regulatory knowledge is essential if you want to bring new drugs faster to market with profound competitive product features.
- In the spirit of regulatory expertise with insight into business opportunities, I have extensive experience in representing the regulatory perspective at governance boards and at meetings with executive stakeholders.
- I bring with me numerous experience interactions with various regulatory agencies around the world as FDA, EMA, PMDA, ANVISA, Health Canada, Saudi FDA and others to discuss aspects ranging from initiation of early clinical trials to ensure agreement on the clinical phase 3 development program.
- Curiosity and a focus to solve problems and finding new solutions which makes a difference is my real passion.
- I am also leader and people manager with an interest in the individual. My aspiration is to ensure high motivation and engagement among employees and create an openness and curiosity to other disciplines. My experience is that this inspires collaboration and a joint understanding of common goals. A working environment that fosters the right balance of working independently and collaborating is key to me.

Work experience

2020-now **Senior Director, Ozack ApS**
Global regulatory expertise within quality, non-clinical and clinical drug development and life-cycle management of regulatory licenses. Finding opportunities by applying a regulatory strategic mindset.

2006-2020 **Senior Director, Regulatory Affairs, Novo Nordisk A/S**
Leading the build-up of regulatory obesity expertise within Novo Nordisk together with 12 highly skilled project managers and professionals.
Responsible for the successful worldwide approval of Saxenda® having a sale in 2019 of 5.679 mill DDK.

2001-2006 **Regulatory Professional, LEO Pharma**
Responsible for planning and executing regulatory strategies for the quality and clinical improvements of marketed critical care products at LEO Pharma as Centyl®, Burinex® and Innohep®

2000-2001 **Project Manager, Danish Medicines Agency**
Administration of regulatory procedures for national extensions, variations, renewals and parallel import applications

Education

2009 Managing Medical Product Innovation (MMPI)
Scandinavian International Management Institute (SIMI)

2000 Master of Pharmacy
University of Copenhagen