Pharmaceutical Business and Drug Development Services

Marianne Svärd, Ph.D

Dr. Svärd offers expert services in pharmaceutical business development and early-stage drug development in collaboration with a network of leading consultants and contract research organizations.

Business development

- Project development plans
- Project evaluations (due diligence)
- Business plans
- IPR management

Outsourcing

Outsourcing of preclinical studies, preformulation studies, drug substance and drug product development, GMP manufacture of API and clinical trial material

Project management

Project management of early-stage drug development and clinical trial applications including coordination of:

- Preformulation work
- Preclinical safety and toxicological studies
- Formulation development (parenteral, topical, inhalation, oral preparations)
- Drug substance and drug product manufacturing for clinical trial
- Clinical study protocol development
- Documentation for CTA application

Also discrete project activities e.g. planning of preclinical studies and drug product development work are performed.

Experience

Marianne Svärd holds a PhD in Physical Chemistry / Surface and Colloid Chemistry from Lund University. Her professional experience includes more than 20 years in pharmaceutical R&D and production. Since 2003 Svärd operates as an independent consultant through Skogsmöllan AB and has managed or participated in several national and international clinical trial applications. For example, Dr. Svärd headed a project which, over a period of two years, took an investigational anti-cancer product from candidate drug to initiation of a phase I/II clinical trial.

Contact

Marianne Svärd M.Sc, Ph.D
P.O. Box 56
SE-247 26 Veberöd
Sweden

Mobile: +46 (0)70 528 59 87
Fax: +46 (0)46 859 86
E-mail: marianne@skogsmollan.se
www.skogsmollan.se