SAXOCON
Co-owned by Technical University of Denmark DTU

Safe equipment and systems

All parts of a production line that come into contact with drug substances or drug products must be designed, tested, and documented to not alter the safety or quality of the finished product. SAXOCON provides you with everything you need to establish and document compliance with regulatory requirements for pharmaceutical manufacturing systems including:

- The selection and characterisation of construction materials
- A risk analysis in accordance with European and US Pharmacopeia standards
- Extractable test strategies
- A review of protocol and monitoring of extractable studies
- A toxicological risk assessment of results

Why choose us?

SAXOCON services for manufacture of pharmaceutical products give you access to:

- A multidisciplinary team of highly skilled toxicologists, material scientists, and supply chain professionals
- Extensive experience in pharmaceutical product safety assessment
- Safety evaluations based on proprietary information obtained from our world-wide network of materials suppliers
- State-of-the-art solutions according to international standards and regulatory guidelines
- Smart, cost-effective test strategies that bring your products to market in a timely manner

Delivery

SAXOCON compiles all necessary documentation for you to receive a Good Manufacturing Practice file as a written report.

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