







Medical devices are intended to improve health, even though materials used for the production of medical devices can contain – whether intended or not – substances or materials considered as nanomaterials or with the potential to cause adverse events, so-called substances of concern.

#### **OUR EXPERTISE**

Since 1989, SCC has specialised in assessing biological risks and finding the best solution for its customers. Our backbone is our scientific expertise bundled in the Regulatory Science department. Experts from various regulatory disciplines, such as chemists, physicists, food chemists, biologists, geo-ecologists, toxicologists, ecotoxicologists and environmental fate specialists, work together under one roof.

#### **BIOCOMPATIBILITY**

The international standards for assessing the biocompatibility of medical devices are known as the ISO 10993 series, with ISO 10993-1 describing the requirements for evaluation and testing within a risk management process and applicable to all medical devices.

This standard stipulates that for every medical device, the information on the chemical composition including potential impurities must be available and "leachables" should be evaluated by means of a chemical characterisation in line with ISO 10993-18.

Such chemical analysis shows the presence of any substance or material that may be absorbed by the patient and should be included in your risk assessment. This procedure helps you to eliminate unnecessary biological tests and focus on identified risks.

But what should you do, if specific substances of concern are present in your product?

# SUBSTANCES OF CONCERN

Materials containing substances with carcinogenic, mutagenic and reprotoxic (CMR) properties are increasingly coming into focus. There is growing political concern regarding the presence and use of endocrine disruptive substances (ED). The EU list of substances of very high concern (SVHC) is growing every 6 months.

While phthalates have been regulated for medical devices and guidelines are available, there is no clear guidance for other substances with potential risks to human health and the environment.



# **OUR SUBSTANCES OF CONCERN SERVICES**

Our experts can help you with:

- Estimating the likely regulatory costs based on a data gap analysis, identifying missing studies and finding waiving opportunities through detailed and systematic database and literature searches
- Developing tailor-made compliance strategies to meet your needs
- Generating in silico data and performing knowledge- and statistics-based QSAR models to predict the activity, properties and toxicity of new or incomplete characterised substances based on the knowledge of their chemical structure
- Planning, contracting and monitoring of laboratory studies using specialised laboratories
- Application of EU guidelines on the benefit-risk assessment of the presence of phthalates
- Establishment of allowable limits for leachable substances in accordance with ISO 10993-17 by our experienced toxicologists
- Dealing with specific requests to expert panels and competent authorities

As a result of our assessment, you will receive a detailed report signed by our certified and registered toxicologists, which is accepted by competent authorities and notified bodies.

# **NANOMATERIALS**

Based on the new classification rule 19 in the EU Medical Device Regulation 2017/745, many medical devices need to be classified in higher risk classes if they contain nanomaterials.

The current EU definition leads to ambiguity and causes testing difficulties when determining if medical devices contain or consist of nanoparticles.

In addition, the potential for internal exposure is not clearly defined within the MDR, leading to additional uncertainty.

How can you recognise if nanomaterials are present if there are no clear methods for their characterisation?

# **TESTING OF NANOMATERIALS**

If nanomaterials are present, biological testing should be performed in line with the recent technical report ISO/TR 10993-22, published in 2017. However, many labs failed to implement new test methods and several methods defined within other ISO 10993 standards need to be adapted in order to be suitable for nanomaterial testing.

How can you handle nanomaterials without approved test methods and sufficient guidance?





#### **OUR SOLUTIONS FOR NANOMATERIALS**

By choosing our services, you will benefit from our expertise in nanomaterials, focused on:

- Characterisation of nanomaterials according to the latest standards and guidelines
- Application of the nanomaterial definition and preparation of assessments to avoid higher classifications and additional tests
- Professional handling of questions regarding internal exposure
- Preparation of literature-based internal exposure assessments for your specific borderline products
- Assistance with selecting biological tests and monitoring services to handle the specific complexity of the test execution.

# MONITORING SERVICES FOR YOUR RAW MATERIAL INVENTORY

SCC has profound skills in monitoring the regulatory status of chemicals. We check your substance inventory against the latest scientific information and regulatory changes on a regular basis. We keep you informed on the most important developments via substance inventory status update services.

#### **YOUR BENEFITS:**

- Full-service support with respect to all issues concerning biocompatibility and environmental concerns relating to medical devices.
- Benefit from using our special tools and experience with databases and literature research.
- Use of advanced in silico methods to close existing data gaps.
- Using our knowledge of the impact of various substances, we carefully select the tests required. This strategy not only helps to save time and costs, but also helps to avoid unnecessary animal testing.
- We have been working with numerous national and international testing laboratories and know which ones can best meet your needs.
- Make sure your products are safe and convince notified bodies and competent authorities with our expert reports.

Whatever your specific needs, SCC helps to ensure your medical devices are fit for the future.

Please visit our website for more information:
<a href="https://www.scc-gmbh.de/medical-devices-substances-of-concern">https://www.scc-gmbh.de/medical-devices-substances-of-concern</a>

or contact one of our experts: <a href="mailto:scc@scc-gmbh.de">scc@scc-gmbh.de</a>