Medical Devices meet Biocidal Products

knoell

worldwide registration

Make the best out of your data by killing two birds with one stone. Leverage your core data set to cover your products under the Medical Device Regulation (MDR), in addition to your global biocidal products portfolio (disinfectants, antimicrobials).

Our global regulatory services include

- Classification & labelling evaluation and review
- Strategic support with borderline cases
- Strategic consulting for active substances, products or Biocidal Product Families (BPFs) (incl. treated articles and borderline cases)
- Covering disinfectants (PT 1–5) under the Biocidal Products Regulation (BPR)
- Efficacy testing strategy development under MDR and BPR
- Expert support for a broad set of services
- Assessment of Endocrine disrupting (ED) properties
- Effects and exposure assessment
- Dietary safety and residues
- Literature search & review
- Interaction with authorities and/or notified bodies

Founded in 1996, knoell is an independent global service provider for biocides, chemicals, cosmetics, crop protection, food contact materials, medical devices and veterinary medicine with sites in Europe, Asia and the Americas.
For further information visit www.knoell.com or call us.

knoell Germany GmbH
Tel +49 621 718858-0
meddev@knoell.com

www.knoell.com

1911-0111