



regenold GmbH

Your partner for development, regulatory
and market access.

Introduction to the regenold organisation

- Founded in 1994 by Dr. Jürgen Regenold
- Offering a full suite of regulatory services from development to market access and life cycle management
- Strong international presence with > 50 % sales outside of Germany
- Proprietary network, **regulanet**[®] which today covers more than 90 countries with over 120 domain and regional experts
- Established separate medical device division, **CE plus** in 2009
- Started developing our presence in data science and analytics in 2015
- Offering legal manufacturer and representative services through our subsidiary, **NEXTEC medical GmbH**
- Today regenold has over 100 employees



Our service offering

Our service range in pharma and medical devices is complemented by Legal Representation services and IT/Data Sciences

- Full-Range Regulatory Services
- Legal Manufacturing & EU Representative Services (NEXTEC)
- Data Science/Validated IT/Data Access

Pharma

- Medicines (Rx & OTC)
- Biopharmaceuticals
- Orphan Drugs
- Vaccines

Devices/IVD

- Active Medical Devices/ Software/Apps
- Non-Active Medical Devices
- IVD & Companion Diagnostics

Other

- Food Supplements
- Cosmetics
- Chemicals
- Biocides



Our Key Differentiators

Experience

Throughout the years we have helped over **2000 clients** with more than **10'000 products** and implemented **+100 QM Systems**.

Expertise

Our Subject Matter Experts possess **extensive practical experience** with backgrounds from industry as well as regulatory bodies.

Communication

We offer **short response times** & dedicated contact points and we maintain **close communication** with Authorities and notified bodies.

Flexible Roles

Flexible roles such as Consultancy, Advisory Board, Regulatory Affairs Manager, Quality Manager, Project Manager etc..

State-of-the-art

Proven experience with **breakthrough technologies** including the use of AI / Digital Biomarkers.

Active member of **numerous networks**, working groups and standard committees.

One-Stop Shop

We provide tailored solutions for **all phases of the product life-cycle** including acting as Legal Manufacturer as well as European Representative.

Pharma Services

- Strategic Advice
- Pharmaceutical Development
- Preclinical Development
- Clinical Development
- Project Management
- Regulatory Strategy & Implementation
- Pharmacovigilance
- Data Science & Analytics
- Auditing
- Market Access
- Portfolio Analysis
- Due Diligence
- Quality Management & Compliance



We offer
customised
solutions with a
focus on innovative
development plans
in view of market
access

We cover the full range of regulatory services in medical devices and IVD's with deep product expertise in most categories



Medical Device & IVD Services

Technical Documentation

- Compilation
- Maintenance

Assessments

- Gap Assessments
- Due Diligence
- Clinical
- Biocompatibility

Quality Management System

- Implementation
- Maintenance
- Auditing

Market Access

- International Registrations in collaboration with **regulanet**[®]

Regulatory Strategies

- Design & Development
- Classification, CE marking

Post-Market Services

- Post-Market Surveillance
- Device Vigilance
- Regulatory Intelligence

Service portfolio of NEXTEC

- Our subsidiary NEXTEC is providing regulatory affairs services and takes over legal responsibility for medical devices and in-vitro diagnostics (MDR 2017/745 and IVDR 2017/746)
- “Legal” manufacturer service (MDR/IVDR Article 10)
- EU REP services (MDR/IVDR Article/11)
- Contract development for Medical device Software and Combination products (MDR/IVDR and EN ISO 13485:2016)



Data Science & Analytics complementing our regulatory service range

- Experienced team and validated AI platform (SAS & open source) geared towards custom development
- Targeted towards development of tailored solutions in GxP and medical device area
- Strong partnership with SAS with technical and business development support
- Regenold AI solutions span R&D through commercial, some of our focus areas are:
 - Digital diagnostics
 - Business analytics for critical insights into drug and device development stages
 - Automation of regulatory processes
 - Market analysis & decision support
 - Clinical decision making

We address the following needs:



A private company
with clients' interests at heart

We are passionate about regulatory –
that is the reason why we provide
a full range of development,
regulatory and market access services

Ability to solve complex combination issues by coverage of
both pharma and medical devices

Customised solutions
tailored to individual client
needs

Long-standing relationships
with regulatory bodies

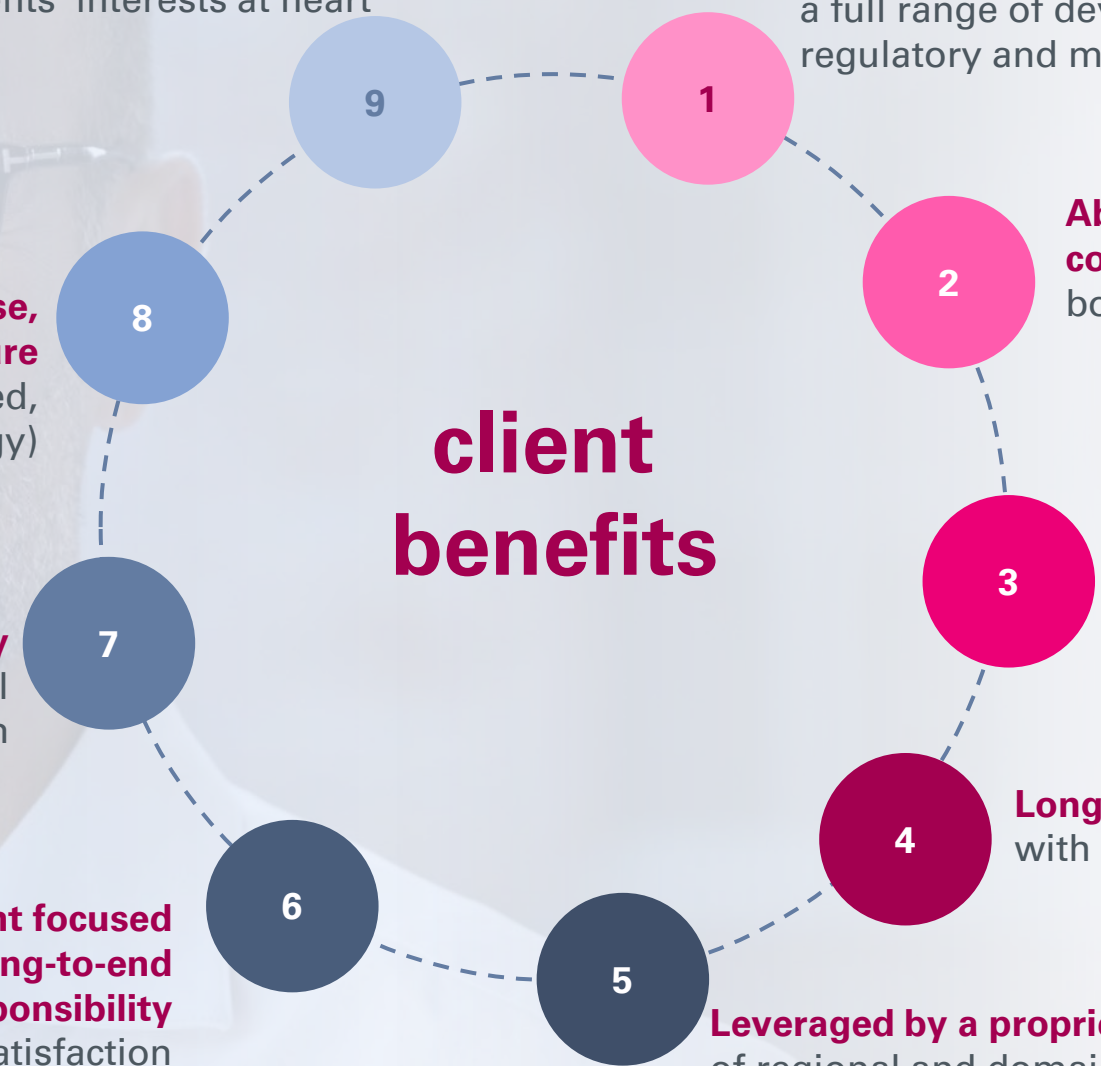
Leveraged by a proprietary network
of regional and domain experts
(regulanet®)

Project management focused on beginning-to-end responsibility
to ensure client satisfaction

Regulatory responsibility services e.g. in PV, QM, legal
representation

We provide expertise, people, and infrastructure
(e.g. data science, validated,
latest technology)

client benefits



Thank You!

Thank you.



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