PwCs Pharma and Life Sciences

Four cutting-edge digital regulatory and tax solutions







Keep up to date with regulatory developments

Overwhelmed by an avalanche of pharma and life science regulation?

The Regulatory Radar gives you a customised overview of all the regulatory initiatives launched by the health authorities in the different countries relevant to you.



It's a cutting-edge digital solution enabling you to collect, document and archive key regulatory data that have an impact on your business in areas such as tax, operational and regulatory strategy. The radar analyses regulatory initiatives and alerts you if there's a potential need for action. You need never miss out on new regulations concerning clinical or commercial trials again.

- Save costs and resources
- Consolidated supervision
- Tailormade solution matching your corporate structure
- Clear prioritisation
- Profit from our knowledge and longstanding experience
- Free up time for key strategic issues

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Meet demands for more transparent regulatory and tax data

The centralised European database for pharma (EU GMP Annex 6 & 16) and medtech products (revision of medical device regulation in the EU) heralds a new era of tighter control and surveillance. Companies will now have to be much more transparent on their regulatory and financial information, including regulatory and tax data, for both commercial and clinical trials.



Are you ready for such a high degree of transparency?

We can help take your company to the next level and prepare for the next digital audit with digital solutions that enable you to aggregate and reconcile your tax and regulatory data, and compare it with public data if required.

- Meet the new mandatory disclosure rules
- Ensure compliance and transparency (audit trail)
- Save time and money
- Structure your information

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The establishment of the EudraGMDP centralised European database and EUDAMED will force pharma and medtech manufacturers to be much more transparent about their regulatory and tax-relevant data.



Now that you have to file your clinical trial and commercial data electronically, can you be sure of doing so in a lean and secure way?

Discover how our blockchain solution, with a focus on the relevant tax topics, can help.

Our digital services help you meet the new digital reporting obligation related to EudraGMDP and EUDAMED. We prepare you for the next digital wave in pharma, life sciences and medtech, empowering you to digitise the transaction workflow and reporting requirement by linking blockchain technology and business processes.

. Digitised supply chain

a tax focus

- . Data traceability and security
- Data transparency and immutability
- Easy verification and exchange of information

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A contract in a day for pharma and life sciences

There's so much to be gained from automating and digitising your routine legal processes. The technology's out there. We help you implement it in the way that works best for your set-up.



We'll install a legal document engine for clinical trials and commercial agreements, and help you automate and optimise your clinical trial and commercial contracting. We can also show you how to improve the interface with clinical research organisations (CROs), investigators and hospitals, as well as automating your process for contract drafting, negotiating and signing.

- Improve your clinical and commercial contract cycle time
- Implement a lean and efficient process with the right mix of technology
- Simplify your processes with smart templates
- Increase visibility and transparency on contracts
- Enhance collaboration between stakeholders such as study teams and legal experts

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"One of the biggest challenges to medicine is the incorporation of information technology in our practices."

Samuel Wilson

Let's find the right formula for your business.