HIGHLIGHT

PatrickToxConsulting

Nonclinical Toxicology Expertise

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Who are you?

PatrickToxConsulting provides toxicology expertise with 25 years in drug development, specializing in biologics *(mAbs, bi-spe, ADC, etc…)*. I help biotech companies achieve regulatory success through safety insights and strategic guidance in preclinical development.

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What is your expertise?

Patrick is a European Registered Toxicologist with extensive experience in biologics, small molecules, and vaccines, gained at Pfizer, Novartis, Pierre Fabre and Sanofi. He brings strong expertise in preclinical development, regulatory toxicology, CRO's interactions and health authorities' expectations, ensuring high quality of consulting services.

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What are your key products or services?

I provide strategic support for nonclinical development plans preparation, monitoring of toxicology studies, writing of regulatory files and tox assessments for impurities from candidate selection up to registration. This approach allows you to optimize your resources and accelerate entry into clinical phases.

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Can you sum up your company in 3 words?

- Insight
- Safety
- Solutions

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Contact Information

Name: Patrick SYNTIN

Job Title: Toxicology Consultant

Phone: +33(0)6-24-20-70-10

E-mail: patricksyntin@patricktoxconsulting.com

patricktoxconsulting.com

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