



TMC

Partnering with small to mid-sized
biotechs and pharma companies to
accelerate drug development and maximise
value across the full product lifecycle

Why Partner with TMC?

How we help you bring transformative therapies to patients faster...



Integrated, end-to-end support delivered by cross-functional experts with extensive knowledge across regulatory, clinical, medical, pharmacovigilance and quality specialties.



Responsiveness of a specialist pharma services company combined with the strength of a full-service partner to help you navigate complexity and accelerate timelines while retaining control and maximising value.



Scalable global associates model provides the agility, insight and infrastructure to make your project happen — whether you're advancing a rare disease treatment or preparing for market entry in the EU and UK.



TMC Pharma Services

Providing end-to-end support — from early-stage planning through clinical execution and commercial launch



Therapeutic Agnostic



Rare Disease, Oncology















Therapeutic Agnostic



Strategic Support Across the Product Lifecycle

Right solution at the right stage — from early drug development to market access and commercialisation

TMC Business Unit	Pre-clinical	Phase 1	Phase 2	Phase 3	Early Access	Approved
 TMC Consulting						
 TMC Clinical						
 TMC Commercial						



TMC Consulting

Expert-led strategic and operational pharma consultancy services



Regulatory support



Pharmacovigilance



Clinical development



Medical affairs



Quality management



Reach critical drug development milestones compliantly, quickly and cost-effectively.



TMC Consulting

Expert-led strategic and operational pharma consultancy services

Gain regulatory clarity that aligns with your commercial goals. TMC Consulting brings deep functional expertise, integrated thinking and a flexible approach to guide you through complex decisions, mitigate risk and align your programme with regulatory and commercial objectives.

Get flexible support that scales with your team. Our experienced, senior specialists act as an extension of your team, partnering with you to deliver custom pharma solutions. These solutions are grounded in scientific rigour and clinical excellence, focusing on regulatory strategy, clinical development planning, market entry readiness and cross-functional integration across all stages of your product lifecycle.

Reach clinical trials — and market — faster. We bring agility and depth to offer you a smarter, faster and more collaborative path from early development to market. Through extensive expertise, robust planning and flawless execution, we help you hit your critical drug development milestones compliantly, on time and within budget to accelerate innovative treatment access for patients who need them most.



TMC Clinical

Early-phase clinical development for rare disease, complex oncology and advanced therapeutics



Accelerate timelines, reduce risk and move confidently towards global regulatory approvals.



TMC Clinical

Early-phase clinical development for rare disease, complex oncology and advanced therapeutics

Move efficiently from first-in-human to global pivotal trials. TMC Clinical partners with you to deliver early-phase clinical development solutions that help bring transformative therapies to patients faster. Our end-to-end clinical services span study design, trial execution, site management, data oversight and regulatory alignment — tailored to the unique demands of each molecule, modality and indication.

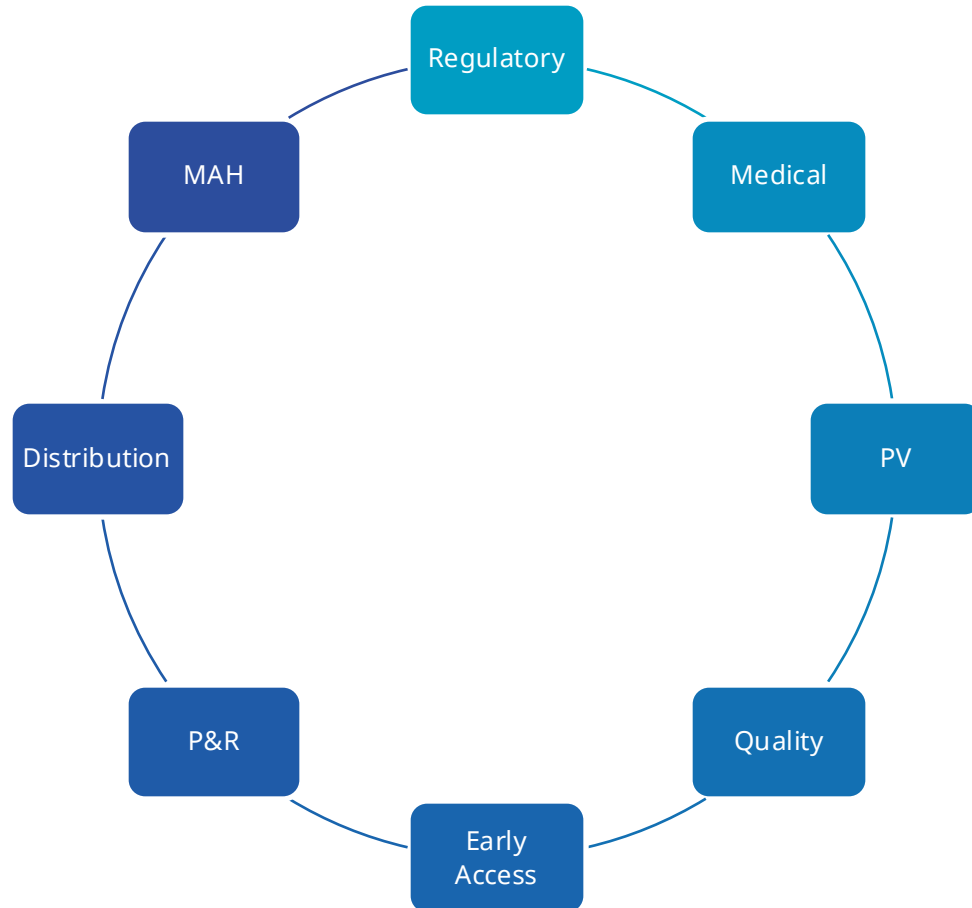
Specialist expertise for challenging, high-impact indications. We specialise in early-phase clinical trials for rare disease, complex oncology and advanced therapeutic modalities, offering tailored support to meet the unique scientific, regulatory and operational demands of these next-generation therapies.

Ensure speed, precision and quality across each phase. With a flexible and collaborative approach, our highly experienced teams integrate seamlessly with yours — combining responsiveness with full-service capabilities to ensure speed, precision and quality across your drug development programme as you move towards global regulatory approvals.



TMC Commercial

Flexible, full-service drug commercialisation solutions in the EU and UK



**Build your own commercial presence
or out-license to a partner,
depending on the path that best fits
your strategy.**



TMC Commercial

Flexible, full-service drug commercialisation solutions in the EU and UK

Launch and scale confidently across the EU and UK. TMC Commercial enables you to launch and scale innovative therapies across the EU and UK with speed and confidence. As your trusted partner, we provide a full-service drug commercialisation solution, including regulatory compliance and marketing application support, medical affairs, pharmacovigilance, pricing and reimbursement strategy, distribution and quality assurance.

Keep control of your asset without the operational burden. Whether as a temporary bridge or a long-term solution for biotechs and pharma companies with limited infrastructure, we have the specialist expertise to act as your Marketing Authorisation Holder (MAH). This means you benefit from our operational capabilities while maintaining the strategic flexibility you need to retain control over your products — without the need to out-licence.

Make informed, long-term decisions that maximise patient access. Our flexible approach, deep expertise and commitment to seamless execution accelerate your product's path to market and help you make informed, long-term strategic decisions that maximise impact and patient access.



Client Testimonials

Your strategic partner in advancing patient health

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“Thank you... and I must take a moment and specifically let you know how well regarded you are here and how grateful we are to be working with you.”

COO, Global Biotech

“Very happy to have you and the TMC Team on our side as our partner! This is a really important project for us, and I can’t think of anyone else I would want in my corner — thank you!”

CSO, Global Biopharma

“I cannot even begin to thank you for all your hard work over the last few weeks driving toward DBL. Every time there seemed to be a light at the end of the tunnel, we added another hurdle! You handled absolutely everything we threw at you brilliantly!”

Head CPM, Biotech

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TMC

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Set up a meeting with our team today to learn more about our capabilities and how we can help you accelerate drug development and maximise value across the full product lifecycle.