



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

Why Partner with TMC?

We can help you bring transformative therapies to patients faster



Fill key skills and knowledge gaps with flexible, scalable support

- Experienced, cross-functional team
- Expert solutions across product lifecycle
- Spanning regulatory, clinical, medical, pharmacovigilance and quality



Overcome global complexities to move forward efficiently and compliantly

- Global Associates Model
- Agile, scalable worldwide delivery framework
- Combining global reach with local expertise



Do more with limited resources (without compromising quality or speed)

- Strong cultural alignment
- Understand the challenges of doing more with less
- Tailored approach to maximise your resources



Reduce risk in critical partnerships and have confidence that we'll deliver

- Proven track record of keeping commitments
- Exceptional client loyalty
- High level of repeat business



TMC Pharma Services

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



Therapeutic Agnostic



Rare Disease, Oncology

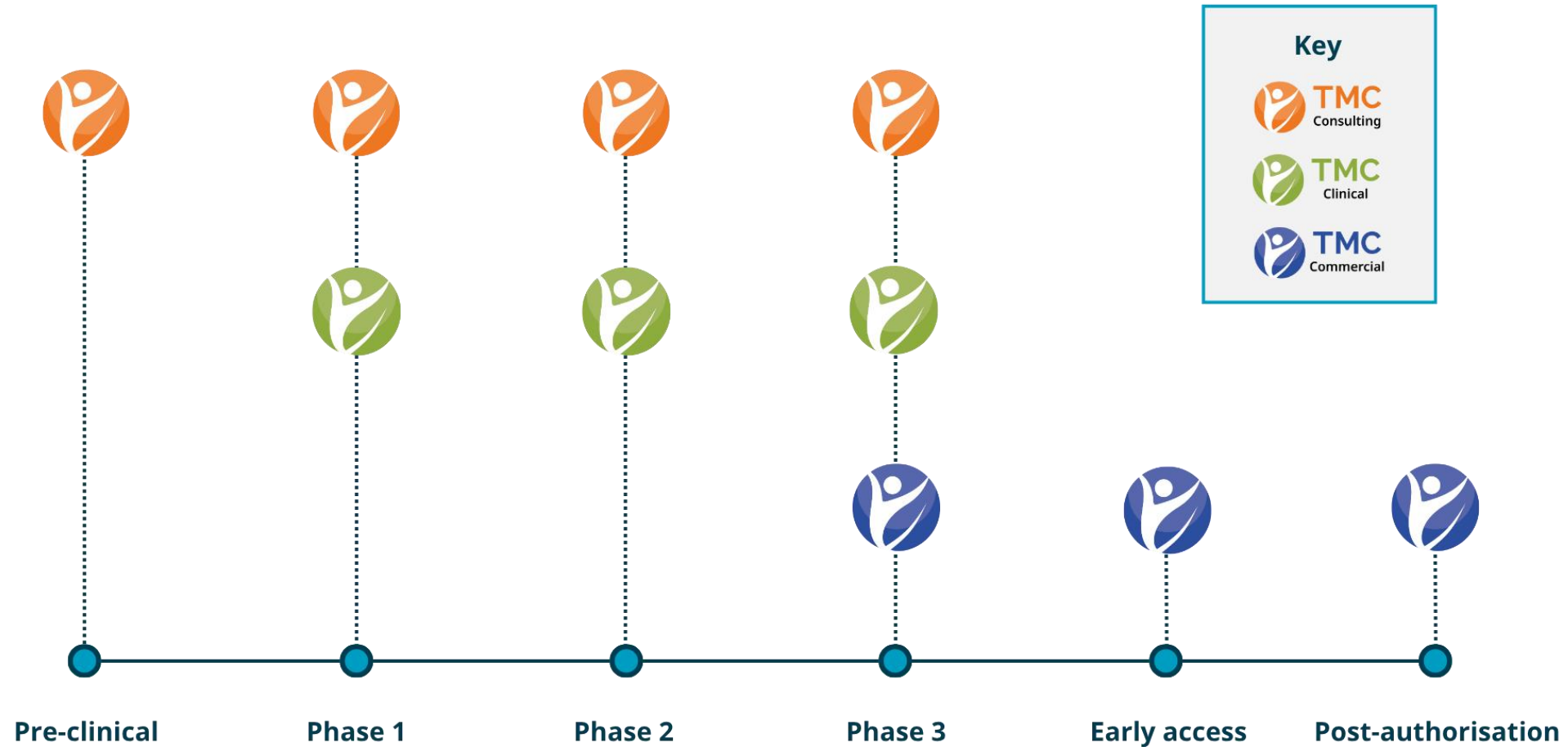


Therapeutic Agnostic



Strategic Support Across the Product Lifecycle

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



TMC Consulting

Expert-led strategic and operational pharma consultancy services

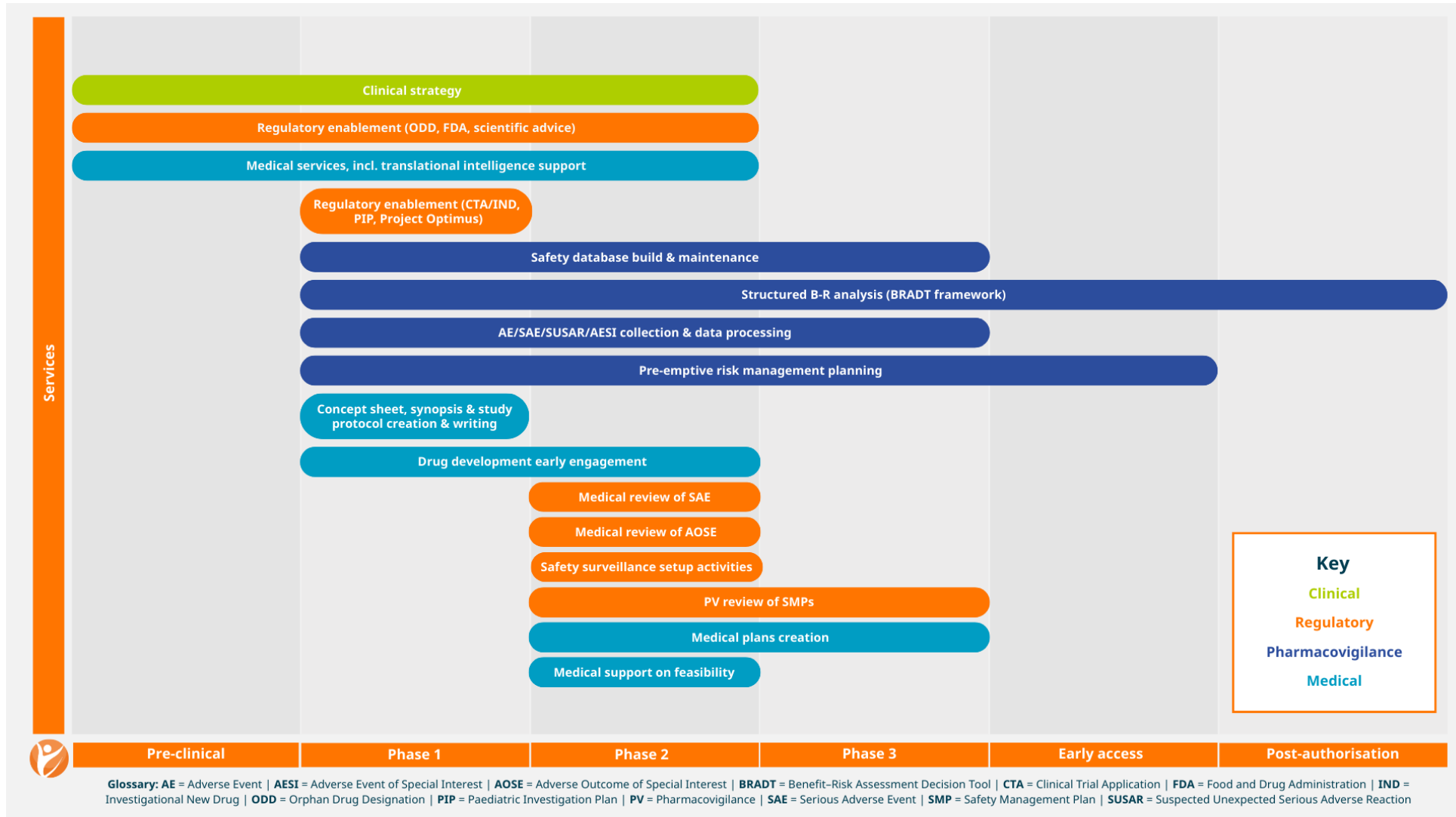


**Access cross-functional expertise
to accelerate your drug
development timeline**



TMC Consulting

Expert-led strategic and operational pharma consultancy services



TMC Clinical

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

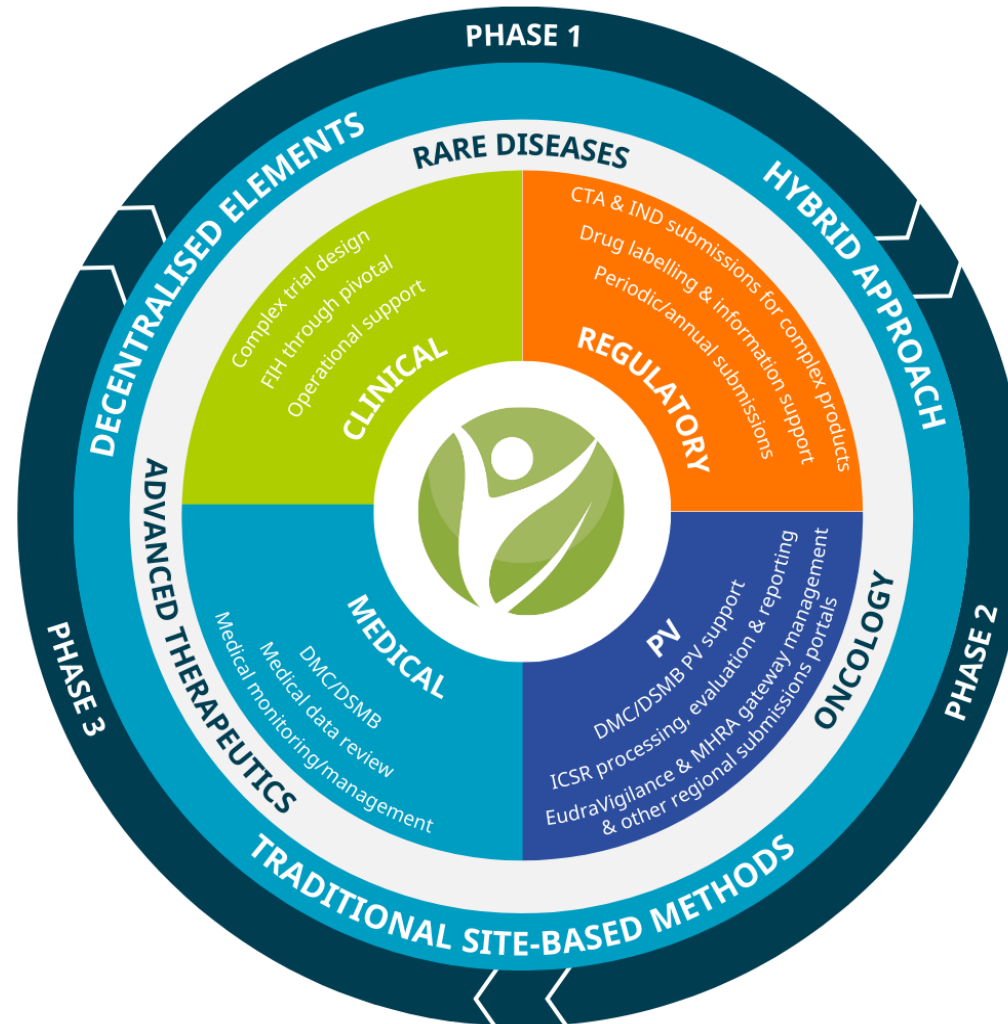


Conduct seamless early-phase clinical trials for advanced therapeutics



TMC Clinical

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



TMC Commercial

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

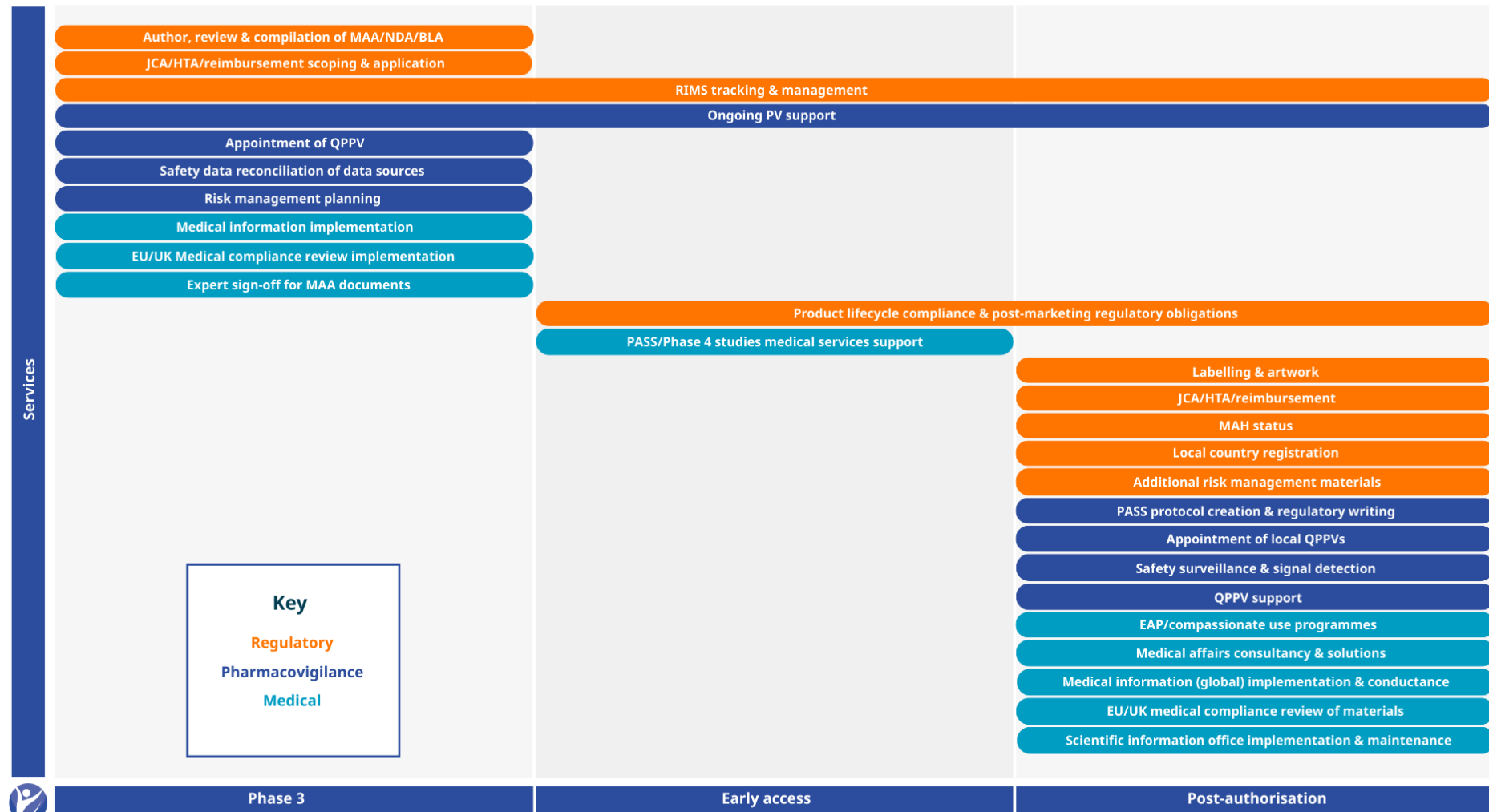


**Gain EU and UK market access
quickly and confidently**



TMC Commercial

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



Glossary: BLA = Biologics License Application | EAP = Early Access Programme | HTA = Health Technology Assessment | JCA = Joint Clinical Assessment | MAA = Marketing Authorisation Application | MAH = Marketing Authorisation Holder | NDA = New Drug Application | PASS = Post-Authorisation Safety Study | PV = Pharmacovigilance | QPPV = Qualified Person Responsible for Pharmacovigilance | RIMS = Regulatory Information Management System



Client Testimonials

Your strategic partner in advancing patient health

“

“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 Client

“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 Client

“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 Client

”





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.