

**R&D AND COMMERCIAL LAW SUPPORT FOR HEALTH**

**We help healthcare organisations draft and negotiate commercial agreements, navigate procurement risk and comply with R&D and privacy rules.**

<ul style="list-style-type: none"> <li>• <b>Research services</b> <ul style="list-style-type: none"> <li>- Contracts for research services.</li> </ul> </li> <li>• <b>Use of facilities</b> <ul style="list-style-type: none"> <li>- Letting someone use your stuff.</li> </ul> </li> <li>• <b>Materials transfers</b> <ul style="list-style-type: none"> <li>- MTAs and related compliance issues.</li> </ul> </li> <li>• <b>Biobanks / tissue collections</b> <ul style="list-style-type: none"> <li>- Human Tissue Act and privacy laws.</li> <li>- Terms of access/supply.</li> </ul> </li> <li>• <b>Research approvals</b> <ul style="list-style-type: none"> <li>- RECs, HRA, Animals, Pathogens.</li> </ul> </li> <li>• <b>Collaboration agreements</b> <ul style="list-style-type: none"> <li>- Collaborations with businesses, academic institutions and charities.</li> </ul> </li> <li>• <b>Clinical trial agreements</b> <ul style="list-style-type: none"> <li>- UK and foreign trials.</li> <li>- Traditional and decentralised/digital.</li> <li>- Investigator and sponsor relationships.</li> </ul> </li> <li>• <b>Clinical trial procurement</b> <ul style="list-style-type: none"> <li>- Contracts for the things needed for a clinical trial (including IMP supplies).</li> <li>- CRO agreements.</li> </ul> </li> <li>• <b>Clinical trial compliance</b> <ul style="list-style-type: none"> <li>- Is it a clinical trial?</li> <li>- Clinical Trial Authorisation</li> <li>- Recruitment</li> <li>- Privacy and transparency</li> </ul> </li> </ul>	<ul style="list-style-type: none"> <li>• <b>Data licences</b> <ul style="list-style-type: none"> <li>- Research or commercial use of a data set.</li> </ul> </li> <li>• <b>Public procurement support</b> <ul style="list-style-type: none"> <li>- Advice on Procurement Act 2023.</li> </ul> </li> <li>• <b>Supply chain flow downs</b> <ul style="list-style-type: none"> <li>- Information.</li> <li>- Net zero.</li> </ul> </li> <li>• <b>Pilots</b> <ul style="list-style-type: none"> <li>- Testing out new solutions.</li> </ul> </li> <li>• <b>Compassionate use</b> <ul style="list-style-type: none"> <li>- Medicines and medical devices.</li> </ul> </li> <li>• <b>Manufacturing</b> <ul style="list-style-type: none"> <li>- Pharmaceuticals and medical devices.</li> <li>- Custom devices / advanced therapies.</li> <li>- Specials and compounding services.</li> </ul> </li> <li>• <b>Contract management</b> <ul style="list-style-type: none"> <li>- Difficult supplier conversations.</li> <li>- Amendment and termination.</li> </ul> </li> <li>• <b>Health law statutory interpretation</b> <ul style="list-style-type: none"> <li>- Opinion on what the law really means.</li> </ul> </li> <li>• <b>Templates</b> <ul style="list-style-type: none"> <li>- Develop or improve contract templates</li> <li>- Harmonise/simplify.</li> </ul> </li> <li>• <b>Artificial intelligence</b> <ul style="list-style-type: none"> <li>- Developing and on-boarding AI solutions.</li> </ul> </li> <li>• <b>Privacy and consent:</b> <ul style="list-style-type: none"> <li>- Privacy policies and consent forms.</li> </ul> </li> </ul>
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**Competitive project and day rates (interim). Save costs compared to law firms.**

**Partner-level advice for a fraction of the price.**

**Single contact for contract, life sciences, R&D, procurement, and privacy.**

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