



## Services

Regulatory Concepts	European Authorized Representative	Legal Representative
<ul style="list-style-type: none"><li>• CE Marking</li><li>• Quality Assurance/ Good Manufacturing Practice</li><li>• International Submissions</li><li>• Registration in Italy</li><li>• Clinical Trial Services</li><li>• Process Validation &amp; Product Safety</li></ul>	<ul style="list-style-type: none"><li>• Acting as <b>European Authorized Representative</b> for medical device and in-vitro diagnostics manufacturer located outside the European Union.</li></ul>	<ul style="list-style-type: none"><li>• Acting as <b>Legal Representative</b> for clinical trials of medicinal products according to Directive 2001/02 EC.</li></ul>

