EDPB Advises on Lawful Grounds for Processing Personal Data in Clinical Trials

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With the coming into effect of the General Data Protection Regulation (GDPR), those conducting clinical trials in the EU face a complex set of rules ranging from lawful grounds for processing and transparency to restrictions on data transfers and secondary uses. To assist with this task the European Commission is in the process of adopting a Q&A document on which it has sought the advice from the European Data Protection Board (EDPB).

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