## €IQV **TECHNOLOGIES**

# **Digital Patient Suite**

Improve the patient experience and increase the efficiency of your virtual clinical trials

IQVIA Technologies helps you maintain trial continuity and effectively run decentralized, hybrid and traditional clinical trials while providing patients with a seamless experience from enrollment through trial completion. The Digital Patient Suite is part of IQVIA Technologies Orchestrated Clinical Trials (OCT), the end-to-end, patient-centric platform that provides an unparalleled data infrastructure, seamless connectivity and intuitive design to **drive smarter**, faster trials.

#### **YOUR CHALLENGES**

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Accelerating the adoption of full or hybrid virtual clinical trials



Recruiting and retaining patients

Keeping patients informed and engaged





Reducing patient burden



Implementing operational changes necessitated by the COVID-19 pandemic



Improving data collection and quality



Working with increased sources of data including devices



Increasing the speed of study start-up



Achieving data transparency and real-time insights



Minimizing administrative burden on site and investigative personnel



Keeping up with rapidly evolving regulatory requirements



Reducing cycle times



Managing trial supplies



Maintaining audit readiness

### **THE SOLUTION**

IQVIA Technologies Digital Patient Suite eases the burden of clinical studies for patients, sites and sponsors. By capturing the patient's voice we are able to intelligently infuse it throughout the patient journey and provide sponsors with access to high-quality, reliable data insights to maximize trial efficiency and maintain trial continuity while reducing costs and cutting cycle times. The Digital Patient Suite is a scalable solution composed of six technology offerings that help solve the key challenges facing patients, sites and sponsors.

Patient Portal	Engage and educate patients and give them access to their data throughout the whole clinical trial
Virtual Trials	Enable decentralized trials, increase patient safety and retention, and improve data collection and quality
eCOA	Optimize data collection and reliability and protocol compliance while enhancing the patient experience
eConsent	Empower patients to manage consent remotely and make informed decisions; provide sites and sponsors with greater transparency and compliance
Cenduit IRT	Optimize trial supply management, accelerate study start-up and simplify site and investigator experiences
Connected Devices	Allow greater access to critical, reliable patient data from medical devices and wearables



#### THE BENEFITS OF THE DIGITAL PATIENT SUITE

FOR PATIENTS	FOR SITES	FOR SPONSORS
<ul> <li>Tools to keep patients better informed</li> <li>Easy to use self-reporting in real-time</li> <li>Intuitive dashboard of results, telehealth and virtual visits</li> <li>Seamless journey from enrollment to study completion</li> </ul>	<ul> <li>Virtual options reduce site burden by managing patient engagement and retention</li> <li>Technology helps accelerate study start-up and optimize drug dispensation and reconciliation</li> <li>Data collection is easier and more accurate through connected devices and electronic reporting</li> </ul>	<ul> <li>Decentralized trials have built-in efficiencies in recruitment and retention</li> <li>Provides improved data quality and real-time insights and efficiencies</li> <li>Rapid study design process cuts time to first-patient-in</li> <li>Electronic audit trail allows for long- term direct-to-patient engagement</li> </ul>

Discover how IQVIA Technologies Digital Patient Suite works in conjunction with the other OCT suites to help optimize clinical trials. Visit <u>IQVIA.com/OCT</u> to learn more.

