













# How Does eConsent Optimize Research?

Electronic informed consent, or eConsent, is an exciting and empowering tool for clinical researchers. But why should you adopt it?

With the right solution, eConsent can

		
		
<p><b>Streamline study startup</b></p>	<p><b>Reduce or eliminate documentation errors</b></p>	<p><b>Improve participant engagement</b></p>
		
<p><b>77% say eConsent improves the process</b></p> <p>In a study at 13 sites in a phase III trial, 77% of sites said eConsent improved the entire consent process <sup>1</sup></p>	<p><b>214 FDA consent citations might have been avoided</b></p> <p>FDA Form 483s from 2011 - 2016 included 214 consent-related citations <sup>2,3</sup></p>	<p><b>69% of sites reported eConsent improves engagement</b></p> <p>Most (69%) sites reported eConsent is a helpful tool for improving subjects' engagement and in boosting their initial understanding of material <sup>1</sup></p>

Your solution should have the following features to take full advantage of eConsent:

		
<p><b>IRB Integration</b></p> <p>Reduce additional steps in the IRB review process</p>	<p><b>Error Locks</b></p> <p>Ensure all required fields are obtained prior to submission</p>	<p><b>Easy Communication</b></p> <p>Discuss questions with prospective participants</p>
<p><b>Real-Time Metrics</b></p> <p>Focus on the research with a solution that collects and reports enrollment metrics</p>	<p><b>Version Control</b></p> <p>Participants only sign the most current consent form</p>	<p><b>Off-Site Preview</b></p> <p>Allow participants to review the consent form at home prior to signing on-site</p>

## Is Your Organization Evaluating eConsent Solutions?

Quorum Review IRB is excited to introduce Q Consent, the only IRB-developed, cloud-based electronic informed consent solution that simplifies research studies, accelerates study startup, integrates with IRB processes, and improves participant engagement.

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