

Your Checklist for Evaluating eConsent Partners

Utilizing improved systems and tools are necessary in the ever-changing clinical trial landscape. eConsent is a powerful tool that can help you efficiently manage patient recruitment, maximize retention, mitigate risk of data loss, and increase ROI. When deciding to leverage the benefits of eConsent, careful consideration of partners is important as vendors should be held to high standard given the sensitive nature of eConsent clinical trial data. During your eConsent evaluation, ask if potential partners have:

- An easy, DIY document creation process
- No programming required
- The ability to edit the compliance process at any time
- The ability to capture electronic signatures and/or upload wet signatures
- Clear requirements for e-signatures or initials to advance pages on the forms
- Capacity to add videos, graphics, quizzes, and FAQs
- The ability Deploy changes in real time
- Consent paths for Legally Authorized Representatives or witnesses
- Tools to enable roles and permissions to limit access to authorized personnel
- Encryption, both at rest and in transit, to protect PI
- Secure documents with passwords and remote backup
- Reporting on enrollment in real time