

Kayentis

Dedicated to eCOA and patient engagement

Kayentis goes one step further in the integration of electronic Informed Consent to improve patient convenience and sponsor efficiency

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The announcement in May 2016 of the partnership between Kayentis and SecureConsent was just the first step in the integration of eConsent into Kayentis eCOA solutions. Today, Kayentis and SecureConsent® strengthen their strategic alliance to enhance the benefits of this integration.



eConsent benefits already available in Kayentis eCOA solutions

- **Kayentis - a global, innovative provider of eCOA solutions - and SecureConsent® - a leading provider of eICF solutions** announced in May 2016 that they have built a Strategic Alliance to optimize the provision of eCOA and eICF for all stakeholders in a clinical study/programme. This alliance enables both solutions to be made available on a single tablet device, allowing the streamlining and acceleration of the setup process as well as the integration of services during study follow-up.

To read: 'A step further in improving clinical trials: an electronic informed consent fully integrated into the eCOA solution'

- **Key advantages of SecureConsent® with Kayentis:**

- * One single tablet device for both solutions.
- * The choice of both Samsung and Android makes it very robust and easy to maintain; OS upgrades are controlled.
- * Kayentis tablets have worldwide 3G capacity based on an alliance with SFR/Vodafone; this ensures that the SecureConsent online solution works regardless of Wifi availability at study sites.
- * Logistics are shared and managed by Kayentis.
- * Tier 1 support is managed by Kayentis; teams are already trained.

And more steps are to come to improve patient engagement!

- **Kayentis/SecureConsent®: an “evolving/dynamic” alliance**

This strategic alliance is continually developing. The aim is to continue to strengthen eConsent integration to continually improve patient engagement and data quality.

- **eConsent benefits under review for 2017 include:**

- * SecureConsent® training performed by Kayentis.
- * Greater integration - SecureConsent® key information will become available in Clin'Form, allowing data management activities to be performed by the Kayentis team (e.g. reconciliation of subject numbers with the randomization tool).
- * Single Sign On for study sites.
- * Joint archiving procedures.

eConsent is a significant step forward in clinical trials and is key to the improvement of patient engagement in eCOA solutions as well as being beneficial to all stakeholders:

- * **Benefits for the Subjects: one single device.**
- * **Benefits for the Sites: simpler setup and handling, a single helpdesk.**
- * **For the Sponsors: accelerated setup, guaranteed connectivity, significant cost savings.**

Guillaume JUGE, CEO - Kayentis - November 30 2016.

// Learn more about SecureConsent® //

*// Contact us for more information about eICF integration in your eCOA study
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