

Basile Trimbur, Medical and Scientific Officer at Kayentis, interviews Trishna Bharadia, Health Advocate & Patient Engagement Advisor, on **decentralized clinical trials and their impact on health disparities**.

Some of the topics covered in this interview are:

Can the gap between the number of patients reaching trials and the study recruitment needs be partially explained by trust issues between patients and the healthcare system? Can DCT help fix this situation? And how?

How would Trishna advise the industry so that we ensure that DCT do not increase the burden and become counter-productive?

Are DCTs a solution to reduce health disparities?

Listen to the full interview:

https://kayentis.com/wp-content/uploads/2022/12/dct-health-disparities-insights-from-a-patient-advocate_interview_kayentis_2022.mp3



Trishna Bharadia
Health Advocate & Patient Engagement Advisor



Basile Trimbur
Medical & Scientific Officer

[Learn more:](#)



Kayentis and ClinOne join forces to improve efficiency and patient experience in clinical trials

10 January 2023

Alliance merges products and services, gaining patients, sites and sponsors unique access to best-in-class eCOA,

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26 December 2022

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Kayentis has more than seventeen years of success stories with Big Pharma, and has been

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Increasing the engagement of study patients can be done in multiple ways: one way to

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Why using a simple televisit feature will benefit your clinical trial

If it is important to enable televisits for DCT trials, embedding the feature directly into the eCOA app brings multiple benefits for patients and sites

- BRINGING SIGNIFICANT COMFORT AND SUPPORTING PATIENT COMPLIANCE**
 - The Scientist feature enables assurance that the study will continue even if the patient does not visit the study site, with:
 - Reminders about patient behavior and patient care responses
 - A study accessible from the eCOA app on the user device
 - A particularly relevant action
 - The physical presence of the patient is not required on site any longer
 - The patient does not need to plan for a given study visit
 - The patient can still need to take medicine at the site schedule
- REDUCING THE STUDY SITE BURDEN**
 - Reduced flexibility for study visit
 - Reduced number of multiple visits
 - Simplified site patient coordination with only one visit
 - Reduced the patient
- ENSURING A HIGH LEVEL OF CONTROL**
 - The transparency of the process is shared by the patient
 - An early patient visit can provide valuable feedback
 - Assessment will also be able to bring control of the site and adjust the strategy according to the specific of the study site

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Televisits for more engaging clinical trials

23 November 2022

If it is important to enable televisits for DCT trials, embedding the feature directly into

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We spoke at DMB on “The shift to ePRO collection at home: Strategies to preserve data quality and re-think compliance”

9 November 2022

Date: November 22 Time: 11:15AM CET Location: Paris, France Presentation in French See the full

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