eConsent International Symposium

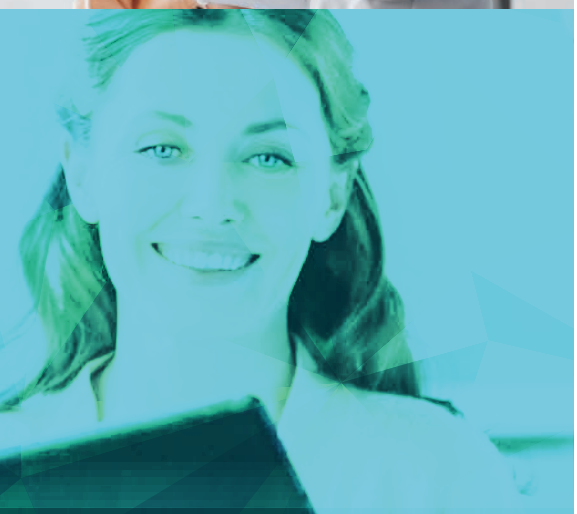
Online / Admission free

English and Japanese (simultaneous interpretation)

Register here >>>

Contact to:

e-mail: dit-dct@hosp.ncgm.go.jp
TEL: +81-(0)3-6228-0445



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National Center for Global Health and Medicine (NCGM)

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Practicing eConsent and Solving Issues in the era of the COVID-19

Date and Time: 2023.**3.3** (Fri) 15:00-17:30 (JST)

While interest in eConsent which uses videos and audio to enhance subjects' understanding of clinical trials and enables us to obtain consent remotely has been growing, the reality is that eConsent is not widely used in clinical trials. This symposium will share the current status and issues, and discuss directions toward solutions.

Moderator

Naoki Tomotsugu
Head of Executive Management, Department of International Trials, Center for Clinical Sciences, NCGM

Naoki Sakuma
Director, Electronic Standard for Medical Information Expert Committee, Drug Evaluation Committee, JPMA (Teijin Pharma)

Program

■ Welcome and opening remarks
Dr. Tatsuo Iiyama, Director, Department of International Trials, Center for Clinical Sciences, NCGM

I Common understanding of eConsent

- Current Status and Issues of eConsent in Drug Development
  Katsuhiko Yoshimoto Electronic Standard for Medical Information Expert Committee, Drug Evaluation Committee, JPMA
- Summary of the basic knowledge of eConsent –merit and demerit-
  Naoki Tomotsugu Head of Executive Management, Department of International Trials, Center for Clinical Sciences, NCGM

II Experience and Challenge for eConsent

- What we can do for Patient Centric IC
  Mami Takahashi Clinical IT Solution Architect, Clinical System and Informatics Group, Biometrics Dept, Chugai Pharmaceutical Co., Ltd.
- Making eConsent a habit in Clinical trial: Experiences and challenges in the EU and the US.
  Ramya Krishna Mokkapati Data management lead, Global Business Division, Remedy & Company
- Global experience and challenge in eConsent
  Hiroyuki Onoguchi Solution Sales Specialist, Direct sales, Medidata solutions K.K.

III Panel Discussion

- Why isn't eConsent more widespread?
- What challenges exist to implementing eConsent?
- What actions should we take with respect to these hurdles?

■ Closing Remarks
Mitsuhiro Kondo, Vice-Chairperson, Drug Evaluation Committee, JPMA