



October 5, 2021
EPS Corporation

News Release

Web Seminar Held for Pharmaceutical Companies and Academic Institutions Considering the Introduction of DDC*1

The EPS Corporation (Head Office: Shinjuku-ku, Tokyo; President and Representative Director: Akira Sasa; hereinafter “EPS”) held Day 4 and Day 5 of its “Startup Guide to Quickly Grasp the Ins and Outs of DCT*2 – Take Your First Step Toward DCT!” seminar series focusing on the topic of DDC on August 24 and August 25, 2021.

よくわかるバーチャル治験
DCTを取り巻くツールの
スタートガイド
～バーチャル治験、DCT導入への一歩を踏み出す!～

ePRO/eCOA
eConsent、オンライン診療
DDC (Direct Data Capture)

Day1	2021年7月19日	(月)	11:00-11:45
Day2	2021年7月20日	(火)	11:00-11:45
Day3	2021年7月21日	(水)	11:00-11:45
Day4	2021年8月24日	(火)	11:00-11:45
Day5	2021年8月25日	(水)	11:00-11:45

参加費無料

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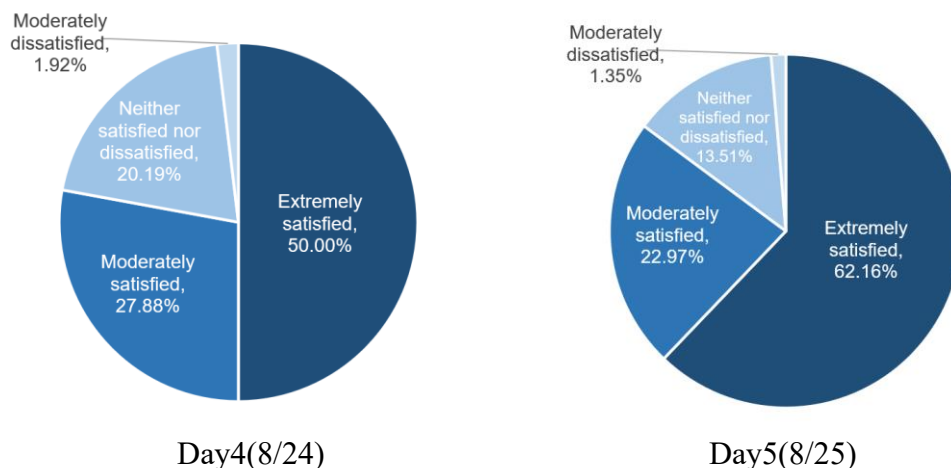
大達 由香里
イービーエス株式会社
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データマネジメント部 リーダー

This seminar generated a large amount of interest in both virtual/online clinical trials as well as DDC with 465 people joining the seminar on Day 4 and 467 people on Day 5.

Over those two days, this basic DDC web seminar provided an overview of DDC and its core concepts, the pros and cons of introducing DDC, regulatory and industry trends, and a wide range of other information.

As a result of a more practical and comprehensive look at DDC and an introduction to real-world examples founded in the cumulative knowledge and experience about clinical studies gained by the EPS Data Management Department, a participant satisfaction survey showed a total of 77.88% on Day 4 and 85.13% on Day 5 were either “Very Satisfied” or “Satisfied” with this web seminar.

Q: How satisfied were you with the web seminar today?



EPS will also hold a “Follow-up Q&A Seminar” on October 26, 2021 to cover the many questions received throughout this entire web seminar series from Day 1 to Day 5.

This follow-up seminar will consist of three sessions: ePRO^{*3}/eCOA^{*4}, eConsent^{*5}/Online Medical Examinations, and DDC. EPS plans to answer many of the questions it received on each of these topics at each session.

Anyone who has not yet applied to join this seminar should feel free to register using the following link (only Japanese):

<https://event.on24.com/wcc/r/3407554/5E8C496ECBF6D7B2FA8486896CE2BFFC>

EPS is also advocating *Virtual Go* as a framework to promote virtual/online clinical trials as one activity to make virtual/online clinical trials commonplace. In the future, EPS will continue to hold regular web seminars on this framework.

EPS *Virtual Go* Framework

As a leading company supporting clinical study work, EPS is advancing the *Virtual Go* framework to promote virtual/online clinical trials using its wealth of expertise and experience in data science, which is backed by an extensive track record.



Virtual Go is a generic name for all EPS services used to promote virtual/online clinical trials.

Virtual Go always strives to satisfy two core ideas: “virtual/online clinical trials in which patients do not have to visit medical institutions” and “virtual/online clinical trials in which CRAs^{*6} do not have to visit medical institutions.” To do this, EPS handles everything from the proposal to

operation and management of various schemes to advance virtual/online clinical trials right for the trial design and type of disease. This includes eConsent, online medical examinations, DDC, eCOA, the use of wearable devices, shipment of investigational product, home-visit nursing, and specimen collection.

About EPS Corporation (<https://www.eps.co.jp/en/>)

EPS launched its business in 1991 as a CRO^{*7} to comprehensively support clinical studies with the focus on clinical trials and PMS^{*8}. The company proposes new models to satisfy customers' needs by leveraging its data science expertise and digital technology cultivated through its extensive track record. These proposals are founded in its Trial GATE concept, which acts as the gateway for all promotion functions of clinical trials.

*1: DDC (Direct Data Capture)

*2: DCT (Decentralized Clinical Trial)

*3: ePRO (Electronic Patient-Reported Outcomes)

*4: eCOA (Electronic Clinical Outcome Assessment)

*5: eConsent (Electronic Consent)

*6: CRA (Clinical Research Associate)

*7: CRO (Contract Research Organization)

*8: PMS (Post Marketing Surveillance)

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