

Closing Remarks

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The Journey Continues...



- Integrating data from multiple sources into a standardized and centralized repository is the solid foundation upon which we can generate novel quantitative and analytical tools and solutions
- Effective and efficient clinical trial design with better patient stratification, and use of established and validated efficacy endpoints
- Accounting for and promoting diversity in future clinical trials
- Better understanding of what matters most to patients and what are the most clinically meaningful measures for tracking disease progression and treatment efficacy



CRITICAL PATH

Thank you!

Pharmaceutical Industry

AbbVie Inc.

Biogen

Boehringer Ingelheim Pharmaceuticals Inc.

Eisai

Eli Lilly and Company

F. Hoffman La Roche

IXICO plc.

Janssen Research & Development LLC Merck, Sharp & Dohme Corporation Novartis Pharmaceuticals Corporation Oxford Brain Diagnostics Takeda Pharmaceuticals Unlearn.AI, Inc vTv Therapeutics



Government and Regulatory Agencies

European Medicines Agency (EMA) National Institute on Aging (NIA) National Institutes of Health (NIH) National Institute of Neurological Disorders and Stroke (NINDS) U.S. Food and Drug Administration (FDA)

Non-profit research Organizations

Alzheimer's Association Alzheimer's Drug Discovery Foundation Alzheimer's Research UK CHDI Foundation UsAgainstAlzheimer's

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