Don’t Miss the CDISC EU Interchange!
Presentations (FDA, IMI, ISO, C-Path and more),
CAB Meeting, Education and Networking!
16-20 April, Stockholm, Sweden
Last Chance for Online Registration - 6 April!

The CDISC vision is to inform patient care and safety through higher quality medical research.

Registration for the conference is currently open through the following link: http://www.cdisc.org/interchange?a=2660#2660.

Plenary Speakers for the conference will feature:
- Dr. Charles Cooper, U.S. Food and Drug Administration (FDA/CDER)
- Mary Ann Slack, U.S. Food and Drug Administration (FDA/CDER)
- Professor Marie Lindquist, Uppsala Monitoring Center
- Dr. Bernard de Bono, speaking on Innovative Medicines Initiative projects
- Lisa Spellman, Secretariat, International Standards Organization (ISO)
- Enrique Aviles, Critical Path Institute
- CDISC Leaders (Dr. Rebecca Kush and Wayne Kubick)
- Opening Remarks by CDISC Board Chair, Paula Brown Stafford

Presentation Topics
- FDA use of CDISC Standards – Challenges and Progress
- IMI Projects and Therapeutic Area Projects
- New Protocol Representation Model Tools for non-technical users
- CDISC Strategic Goals and Technical Roadmap
- CDISC Standards – Case Studies, Progress and Implementation
- Reports from the Global CDISC Coordinating Committees and User Networks
- SHARE, Metadata, BRIDG

Educational Courses – Certified CDISC Training
- SDTM Theory and Application
- ODM Implementation
- Protocol Representation
- ADaM Implementation
- CDASH Implementation
- Controlled Terminology
  PLUS - Legacy Data Conversion Workshop and BRIDG Deep Dive Workshop

Don’t miss the CDISC Advisory Board (CAB) meeting on Tuesday (17 April) 4:30-6:00, with Private Q&A session with FDA, ISO, IMI, CDISC Board members and CTO.
Networking Boat Ride on the Bay of Stockholm!

Last but not least, do not miss our luxurious evening event on a boat ride on the inlet to Stockholm accompanied with dinner and refreshments. Seating is limited so register now to guarantee your seat! Tickets are offered online through either of the following links: http://www.cdisc.org/interchange?a=2660#2660 or http://www.event.com/events/cdisc-interchange-europe-2012/event-summary-ec2945e764e44519b869972aa28558e7.aspx.

If you have read this far and still don’t know why you should attend this Interchange, here is some advice from our Program Chair from the E3C.

“CDISC is driven by our stakeholders: Standards accepted for progress, CDISC reliable as a partner”

Attending the CDISC Interchange conferences is always a big event. You will be able to:
- Get a full picture about CDISC standards and how they help you to streamline your Clinical Research activities and speed up biopharmaceutical development
- Network with more than 200 industry experts, regulators and scientists
- Find out how to get involved in different CDISC working areas like the content development for therapeutic areas
- See the latest available technologies and services around the CDISC standards at a great exhibition

Contribute to the CDISC vision of informing patient care and safety through higher quality medical research.

After 6 April, only offline registration will be open!

Further questions on the CDISC Europe Interchange should be forwarded to Diana Harakeh (dharakeh@cdisc.org).