ABSTRACT

Rebecca Kush, Ph.D., President, CDISC (Clinical Data Interchange Standards Consortium, Inc), will present the status of industry standards to support the electronic acquisition, exchange, submission and archiving of clinical trials data and metadata to improve data quality and accelerate medical and biopharmaceutical product development. She will share use case examples and adoption rates of the production model standards, in addition to progress on new standards, such as clinical trial protocol representation. The collaboration between Health Level Seven and CDISC will be discussed in the context of the expanding mission of CDISC to ensure interoperable standards between clinical research and healthcare.

NOTE

No paper was made available for publication. Please contact the author directly.

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