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Replies to Inquiries to FDA on Good Clinical Practice | FDA

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Good Clinical Practice (GCP) | NIHR

GCP Questions, FDA Answers: Transfer of Study Records. August 12, 2019. The FDA's Office of Good Clinical Practice responds to inquiries on a variety of trial-related subjects, providing answers on the agency's official regulations as well as best practices. The following is one such question and answer excerpted from the CenterWatch publication, GCP Questions, FDA Answers.

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MAGI - Clinical Research Conference 2020

In any clinical trial, good clinical practice (GCP) must be consistently maintained. According to the Food & Drug Administration (FDA), GCP is “an international ethical and scientific quality standard for the design, conduct, performance, monitoring, auditing, recording, analyses and reporting of clinical trials.”

Preparing for Your Next Good Clinical Practice Audit

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