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... Good Clinical Practice (GCP) C.

The Declaration of Helsinki. D.

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5. When a short form is used for Informed Consent, the patient signs the short form **ONLY** and the receives

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a copy of both the summary and the short form.

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The Good Guidance Practice

regulation can be found at 21 CFR

10.115 21 CFR 10.115 (b) What is a

guidance document? (1) Guidance

documents are documents prepared

for FDA staff, applicants/sponsors, and

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the public that describe the agency's interpretation of or policy on a regulatory issue.

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anyone taken the most recent CCRP exam? I'm taking it in a few days and have no info aside from the manual. All the flash card sites and sample test questions I've found online are from the previous format.

New Updated CCRP Exam (SOCRA) :

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(Q&A) Follow . Most Read; Give
Answer; Does the patient sign the
short form only and he receives a copy
of both the summary and the short

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form when a short form is used for
Informed Consent? John Adney
Answered: Feb 24, 2017 § 50.27
Documentation of informed consent.

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