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Autore	Tereskerz Patricia M
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Descrizione fisica	1 online resource (282 p.)
Altri autori (Persone)	EdelmanRobert <1936-> McKinneyRoss
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Soggetti	Biology - Research - Law and legislation - United States Medicine - Research - Law and legislation - United States Clinical trials - Law and legislation - United States Human experimentation in medicine - Law and legislation - United States
Lingua di pubblicazione	Inglese
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Livello bibliografico	Monografia
Note generali	Description based upon print version of record.
Nota di bibliografia	Includes bibliographical references and index.
Nota di contenuto	Research malpractice & negligence -- Duty of care : understanding the legal differences between medical treatment and medical research -- Establishing standard of care & violation of standard of care -- Informed consent in clinical research -- Liability issues for institutional review boards (IRBs) and data safety monitoring boards (DSMBs) -- Legal aspects of financial conflicts of interest in clinical trials -- Disclosure of clinical trial information : legal ramifications of withholding study results -- Clinical trials & insider trading -- Clinical trials and criminal law -- Clinical trial contracts.
Sommario/riassunto	This book provides a comprehensive resource for medical professionals on the various legal aspects involved in conducting clinical research. It encompasses legal and ethical issues such as duty of care, research malpractice and negligence, standards of care, informed consent, liability issues for Institutional Review Boards (IRB), conflicts of interest,

insider trading and the disclosure and withholding of clinical trial results. It will also provide legal guidance on research contracts, setting up clinical trials and common legal pitfalls encountered in medical research.
