

1. Record Nr.	UNINA9910830181503321
Autore	Sharpe Neil F
Titolo	Genetic testing [[electronic resource]] : care, consent, and liability // Neil F. Sharpe, Ronald F. Carter
Pubbl/distr/stampa	Hoboken, N.J., : Wiley-Liss, c2006
ISBN	1-280-28812-4 9786610288120 0-470-36082-8 0-471-74889-7 0-471-74888-9
Descrizione fisica	1 online resource (622 p.)
Altri autori (Persone)	CarterRonald F
Disciplina	616.042 616/.042
Soggetti	Human chromosome abnormalities
Lingua di pubblicazione	Inglese
Formato	Materiale a stampa
Livello bibliografico	Monografia
Note generali	Description based upon print version of record.
Nota di bibliografia	Includes bibliographical references (p. 453-566) and index.
Nota di contenuto	Genetic Testing; Contents; Contributors; Foreword; 1 Genetic Counseling and the Physician-Patient Relationship; Overview; Introduction; Core Concepts; Trust, Genetic Testing, and the Physician-Patient Relationship; Genetic Counseling and the Physician-Patient Relationship; Nondirectiveness; Web Resources; Genetic Counseling Approach to Genetic Testing; Introduction to Genetic Counseling; Historical Perspective and the Role of Nondirectiveness in Genetic Counseling; Practice of Genetic Counseling Today; Genetic Counselor Goals; Important Cultural Considerations in Genetic Counseling Factors That Promote Effective Genetic CounselingGenetics in Mainstream Medical Care: The Emerging Milieu; Categories of Genetic Testing: Issues to Consider; Summary; 2 Communication; Overview; Need To Know; Standards; Watch Out For; Obligations; Methods; Terms, Tone, and Follow-up; Culture and Communication in the Realm of Fetal Diagnosis: Unique Considerations for Latino Patients; Miscommunication and Prenatal Genetic Care; Idioms and Jargon; Suggestions; Cultural Sensitivity; Genetic Consultation; Opportunistic Observation; Translation and Second-Hand Information; On-Site

Translators

Genetic Consultation Interview after Consultation; Family and Friends as Translators; Confidence and Trust; Woman Who Declined Amniocentesis; Woman Who Accepted Amniocentesis; Discussion: Communication, Miscommunication, and Genetic Counseling; Conclusion; Communication: Clinical Diagnosis; History; Background and Diagnostic Criteria for NF1; Molecular Genetics; Issues; Introduction; Client Characteristics: Culture and Intellect; Educational Tools and Facilitators; Decision Making: Considerations and Approaches; 3 Psychological Aspects; Overview; Introduction; Scenario 1; Scenario 2
Need to Know Potential for Psychological Harm; Potential for Psychological Benefit; Anxiety and Perceived Risk: Pretest Status of Patient; Recognizing Psychological Stress; Coping with Stress Induced by Genetic Counseling and Testing; Watch Out For; Factors Affecting Perception; Standards and Interpretation; Psychological Aspects of Genetic Testing For Adult-Onset Hereditary Disorders; Introduction; Process of Genetic Testing; Pretest Genetic Counseling Phase; Posttest Counseling Phase: Disclosure of the Genetic Test Result; Test Result Indicating Presence of Genetic Mutation
Negative Test Result Indicating Absence of Genetic Mutation Receiving an Ambiguous Genetic Test Result; Impact of Genetic Testing on the Family; Societal and Ethical Issues Linked to Psychosocial Outcomes; Clinical Implications; Summary; 4 Duty of Care; Overview; Need to Know; Standard of Care; Watch Out For; Duty of Care; 5 Family History; Overview; Introduction; Clinical Scenario; Need to Know; Family Pedigree; Watch Out For; Inaccurate or Incomplete Information; Prenatal and Neonatal Testing; Family History in Adult-Onset Disorders Importance of Family History in Approaches to Common Chronic Diseases of Adulthood

Sommario/riassunto

A complete review of the issues with specific recommendations and guidelines. With over 1,000 tests commercially available, genetic testing is revolutionizing medicine. Health care professionals diagnosing and treating patients today must consider genetic factors, the risks and limitations of genetic testing, and the relevant law. Genetic Testing: Care, Consent, and Liability offers the only complete, practical treatment of the genetic, clinical, ethical, and legal issue surrounding genetic testing. The authors present present protocols, policies, and models of care that are currently in use, a
