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	Titolo	Fetal morph functional diagnosis / / Hideaki Masuzaki, editor
	Pubbl/distr/stampa	Singapore : , : Springer, , [2021] ©2021
	ISBN	981-15-8171-1
	Edizione	[1st ed. 2021.]
	Descrizione fisica	1 online resource (IX, 354 p. 108 illus., 60 illus. in color.)
	Collana	Comprehensive gynecology and obstetrics
	Disciplina	618.1
	Soggetti	Radiology Gynecology Gene therapy
	Lingua di pubblicazione	Inglese
	Formato	Materiale a stampa
	Livello bibliografico	Monografia
	Nota di contenuto	Part I Ultrasound examination 1.Ultrasonic Screenin 2.Abnormal findings in Ultrasound examination Part II_ Genetic tests 3.  Screening tests 4.Diagnostic test Part III_Genetic Disorders 5.  Mendelian Diseases 6.Abnormalities in fetal brain(congenital hydrocephalus etc) 7.Muscular Dystrophy 8.Skeletal dysplasia 9.Genito-urinary tract abnormality 10.Genomic imprinting disorders (including mesenchymal placental dysplasia) 11.Genetics in Multiple gestation 12.Fetal Therapy Part IV_Chromosoma disease 13.  Autosomal disease 14.Sex chromosome-linked diseases Part V_Genetic counseling 15.fetal anomaly and genetic counseling 16.Soft marker test (NT, Nasal bone etc) and genetic counseling 17.  NIPT and genetic counseling 18.Trisomy and genetic counseling 19.Sex chromosomel abnormality and genetic counseling 20.  Chromosome structural abnormalities and genetic counseling 21.  Chromosome Mosaic and Genetic Counseling 22.Gene disorders and genetic counseling Part VI_Technical 23. G-banding 24. FISH 25. PCR 26. Microarray and Next generation sequencing 27.  How to get the licenses for prenatal diagnosis.
	Sommario/riassunto	This book explores the recent clinical and research findings in the field of prenatal screening and diagnosis. It presents new devices and tests such as real-time 3D ultrasound, ultrafast fetal MRI, and next-

generation sequencing and discusses genetic counseling and fetal therapy. Written by pioneering scientists, the book is divided into six themed parts: ultrasound examination, genetic tests, genetic disorders, chromosomal diseases, genetic counseling, and techniques, presenting carefully prepared original data. This thought-provoking, instructive and informative book is intended for geneticists, obstetricians, pediatricians, genetic counselors and nurses. Although the incidence of congenital abnormalities such as structural, chromosomal and genetic disorders is very low, it is important to have accurate information on their incidence and likely outcome, and on the screening and diagnosis of congenital abnormalities during pregnancy care. This book provides valuable insights into prenatal screening, genetic counseling and fetal diagnosis.