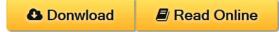


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New Clinical Genetics, third edition

By Andrew Read, Dian Donnai



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the widespread use of next-generation sequencing as a routine diagnostic tool
the checking of a patient's whole exome for the cause of their problem
noninvasive prenatal diagnosis by next-generation sequencing of free fetal DNA in the maternal circulation

•a new integrated treatment of epigenetics

•mosaicism, 'RASopathies' and disorders of the spliceosome are described in new *Disease boxes*

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