# Molekularmedizinische Grundlagen von hereditären Tumorerkrankungen (Molekulare Medizin) (German Edition)

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Durch den molekularbiologischen Erkenntnisfortschritt und die damit zusammenhängenden Methodenentwicklungen sowie die kürzlich gelungene Entschlüsselung des menschlichen Genoms ergeben sich auch für die Krebsforschung und insbesondere für die Früherkennung hereditärer Tumore durch eine umfassende Gendiagnostik neue Möglichkeiten für die Erfassung von genetischen Risikofaktoren, die zur Krebsentstehung prädisponieren. In 19 Kapiteln werden von ausgewiesenen Experten die heute erkennbaren molekularen Ursachen von hereditären Krebserkrankungen dargestellt. Die Themen reichen von malignen Veränderungen des Auges, der Haut und der Brust bis zu Tumorerkrankungen der Bauchspeicheldrüse, der Niere und der Prostata. In 5 Kapiteln werden gastrointestinale Erkrankungen in einem gesonderten Abschnitt beschrieben.

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