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Handbook of Cystic Fibrosis

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Handbook of Cystic Fibrosis



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The dysfunction of CFTR results in the creation of solid mucus in the lungs and digestive tract, leading to pulmonary and gastrointestinal manifestations. The incidence of cystic fibrosis in European countries and the United States ranges between 1 in 2,000 and 1 in 25,000. Cystic fibrosis can be an inherited condition in which a mutation in the gene coding for the cystic fibrosis transmembrane conductance regulator (CFTR) causes lack of function. This concise manual provides clinicians and other related health care professionals with an important reference tool to the background of cystic fibrosis, and the management and treatment of the disease. The latest suggestions are examined and current and emerging treatments are talked about in the latter chapters.



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