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Gene Mutations; Chapter 9. Lung Function Testing in Infants

Chapter 10. Assessment of Lung Function in Young Children with Cystic FibrosisChapter 11. Lung Function Testing in School-Age Children with

Cystic Fibrosis; Chapter 12. Thoracic Imaging in Cystic Fibrosis

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Chapter 30. A Biopsychosocial Model of Cystic Fibrosis: Social and Emotional Functioning, Adherence, and Quality of LifeChapter 31. Palliative and End-of-Life Care in Cystic Fibrosis; Index; Back Cover

<P>The median age of survival for those with cystic fibrosis has risen considerably in recent years. This text thoroughly examines the developments and breakthroughs which have led to this improvement in life expectancy. With a focus on the latest discoveries in the diagnosis and treatment of the disease, this book provides a comprehensive overview of the past, current and forthcoming advancements in cystic fibrosis research and clinical care.

## Sommario/riassunto