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Cystic Fibrosis and Life Expectancy "A critical review of studies on the effect of cystic fibrosis on the life expectancy of CF patients." \*Massive Life Expectancy -- Why Should we ask the question?. Author(s) National Center for Cystic Fibrosis Partitioning for Survival: A Web-Based Approach Capstone Biosciences Associates:. The pre-operative global functional assessment (GFA) score predicts post-operative global function in children with thoracic and abdominal congenital heart disease. 1.2. 2015Medscape Medical NewsFamilial cystic fibrosis (CF) is the most common fatal genetic disease of the Caucasian population (1). The gene mutation responsible for this autosomal recessive disease is located on the long arm of chromosome 7, specifically the cystic fibrosis transmembrane conductance regulator (CFTR) gene, which codes for a 1304 amino acid protein. CF is a life-threatening disease that affects multiple organs. CF causes lifelong morbidity, mortality, and morbidity. Most of the individuals with CF are affected by recurring severe respiratory infections, which cause recurrent episodes of pulmonary disease and life-threatening complications. Though the disease can have significant effects on the affected individual, it is equally devastating to the patient's family, who become primary caregivers and must deal with a condition that will change their lives. The mutations in the CFTR gene typically cause a multidrug-resistant disorder, although there are rare exceptions. For example, the missense mutation, which causes replacement of phenylalanine for cysteine at position 508 (F508C), causes a decreased permeability of ion channels, reduced absorption of salt and water, and a lessened hyperabsorption of polypeptides across the intestine. About 85% of individuals with CF are homozygous for the F508C mutation. CF is the most common lethal hereditary disease in Caucasians, affecting about 1 in 2500 live births (Figure 1). More than 50% of individuals with CF are diagnosed during the first year of life. Though these patients survive into their second or third decade, the majority will experience life-threatening respiratory infections beginning in early childhood that are associated with progressive lung disease. In 2011, the median age of death of individuals with CF was 30 years of age (Figure 2), with half of the deaths occurring between the ages of 20 and 29 years. Most individuals with CF have pancreatic insufficiency (cystic

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