

## THE PANCREAS IN MUCOVISCIDOSIS

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Introduction: Cystic fibrosis is an inherited-recessive disease with progressive chronic evolution, caused by a defect in the CFTR gene. The pancreas is one of the most commonly affected organs by disease, leading to pancreatic insufficiency and significant decrease in life expectancy.

Keywords: Cystic fibrosis, 70% pancreas, malnutrition, 60% mutation. 50% Purpose: To study the 40% genetic aspects in 30% affecting the pancreas 20% with cystic fibrosis by 10% presenting the following 0% clinical research.

**Material and methods:** A clinical research was conducted and analyzes the statistics of 49 pacients known with cystic fibrosis at the IMSP Institutul Mamei și Copilului. Aspects analyzed: patient age (PA), disease onset (DO), fecal elastase value (FEV), mutant gene class (MGC), body mass index (BMI).

## Results Mutant gene class (fig.1) Patient 0-5 5-10 10-15 15-20 20-25 25-30 30-35 Class III, V, VI age Class IV (years) 12 13 Pers. Class II Class The information above show the presence of CFTR gene Fig. 3 mutation in all pacients, hence the following data was recorded 10% Fecal elastase value (fig.2): low values of an elastase normal value 90% G 542 X 2184 ins 663 dup F508

## Conclusions:

- (1) Cystic fibrosis is a monogenic disease, the diagnosis of which is established, mainly during the first year of life.
- (2) The F508 delta mutation is the most common, respectively, class II remains the most affected.
- (3)Class I and II lead to a classic CF phenotype with pancreatic insufficiency.