

A New Clinical Scenario: the Presence of Neutrophil Anti-Cytoplasm Antibodies in the Combination of Pulmonary Fibrosis and Emphysema (CFPE)

Jose Salvador García-Morillo*, David Blanco Alba and Santiago Rodríguez Suárez

Rare and Autoimmune Disease Unit, Internal Medicine Service, Virgen del Rocio University Hospital, Seville, Spain

***Corresponding author:** Dr. Jose Salvador García-Morillo, Rare and Autoimmune Disease Unit, Internal Medicine Service, Virgen del Rocio University Hospital, Seville, Spain, Tel: (+34) 678 421 588; E-mail: salvaymar@gmail.com

Received: July 23, 2020; **Accepted:** August 06, 2020; **Published:** August 13, 2020

Copyright: ©



Figure 1: Chest X-ray. HCRT tórax base. HCRT tórax vertice.

Smoking is assumed as the main etiological factor, as it is a constant in all cohorts studied [8], when it occurs in an individual with a genetic predisposition (the relationship with mutations of the SFTPC gene of the C protein of the pulmonary surfactant is known [9] as well as mutations of the TERT, TR and NAF1 genes of telomerases) [10]. In