

A late preterm pregnancy in a patient with cystic fibrosis (CF): case report

Cesare D'Orsi ¹, Rosaria Casciaro ², Francesca Cappozzo ¹, Elisa Calzolari ³, Carlo Castellani ^{2,*}

¹Department of Neurosciences, Rehabilitation, Ophthalmology, Genetics and Maternal and Child Health (DiNOGMI), University of Genoa, Genoa, Italy.

²Cystic Fibrosis Center, IRCCS Istituto Giannina Gaslini, Genoa, Italy.

³Department of Internal Medicine (DIMI), Allergy and Respiratory Diseases, IRCCS Policlinico San Martino, University of Genoa, Genoa, Italy.

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Background. Literature suggests that maternal and fetal pregnancy outcomes are satisfactory for cystic fibrosis (CF) affected women with well controlled clinical features. We present a case highlighting that even in CF patients in stable conditions the risk of severe clinical deterioration in the course of pregnancy is not to be overlooked.

Case presentation. A 25-year-old woman with CF and a history of a previous pregnancy ended in miscarriage. She presented a pre-pregnancy BMI of 21.1 kg/m², FEV1 60%, and was colonized by MSSA, therefore resulting in a low risk profile for pregnancy. The patient's general conditions progressively worsened, with four pulmonary infective exacerbations, a gradual decrease of pulmonary function, need of noninvasive oxygen support, and deterioration of nutritional status despite parenteral nutrition (weight be-

fore pregnancy 56 kg, at 36 weeks 54 kg). Anticipation of delivery was decided. A healthy female child was delivered by elective cesarean section at 36+0 weeks, Apgar 1' 9-5' 10, weight at birth 2850 g. She was discharged at 4 days with 12.6% loss of the birth weight and mild jaundice. In the following days feedings were switched to infant formula exclusive nutrition, due to lack of breast milk. The patient was discharged 9 days postpartum, after a 12-day i.v. antibiotic cycle, with improved general conditions and basal pulmonary function comparable to usual, pre-pregnancy, standards.

Conclusions. Even in CF women with a low risk profile before pregnancy, a careful monitoring of pregnancy and active collaboration between the CF center, anesthesiology and the Gynecological/Obstetrical Unit is highly recommendable.