Effects of granulocyte colony-stimulating factor therapy for osteogenesis imperfecta: a case report

Paszko-Patej G.^{A,D,E,F}, Sienkiewicz D.^{E,F}, Kułak W.^{A,D,E,F}, Okurowska-Zawada B.^{E,F}, Wojtkowski J.^{E,F}, Kalinowska A.^{E,F}, Okulczyk K.^{E,F}, Sochoń K.^{E,F}, Dmitruk E.^{E,F}, Mirska A.^{E,F}.

Department of Pediatric Rehabilitation and Center of Early Support for Handicapped Children "Give a Chance," Medical University of Bialystok, Bialystok, Poland

A- Conception and study design; **B** - Collection of data; **C** - Data analysis; **D** - Writing the paper;

E- Review article; F - Approval of the final version of the article; G - Other (please specify)

ABSTRACT

Introduction: Osteogenesis imperfecta (OI) is a genetic disorder of increased bone fragility and low bone mass. OI type IV.

Materials and methods: We examined the safety and effectiveness of a low dose of analog granulocyte colony-stimulating factor (G-CSF) in a 15-year-old girl OI type IV. G-CSF 5 μ g/kg was given subcutaneously, for 5 days/month for 3, 6 and 12 months. Laboratory tests, including blood, biochemical tests were performed, in addition to clinical examination.

Results: Clinical examination revealed an increase of muscle strength in the upper and lower limbs between base line and day 6 and 12 months. We found no serious adverse events. Leukocyte levels remained below $38,000/\mu L$. Low dose G-CSF was safe and well tolerated by the patient.

Conclusions: A significant increase in muscle strength in this patient may indicate beneficial effects of G-CSF factor in this disorder. These results are inspiring and warrant further studies.

Keywords: Osteogenesis imperfecta; granulocyte colony-stimulating factor; muscle strength

DOI: 10.5604/01.3001.0010.1885