A 5-year-old girl was born with photophobia (A) and hair loss (B). Bilateral corneal neovascularization, superficial punctatekeratopathy (C), meibomian gland drop-out with thickened meibum (D), and eyelid hyperkeratosis (C) were noted. Physical examinationwas unremarkable except for extensive non-scarring atrichia of scalp and hyperkeratotic follicles. A heterozygous mutation c.1573\_1581del(p.P525\_R527del) in SREBF1 gene was detected, whereas family detection was negative.