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Novel TINF2 gene mutation in dyskeratosis congenita with extremely short telomeres: A case report

prepared by the authors

Verónica Judith Picos-Cárdenas, Saúl Armando Beltrán-Ontiveros, José Alfonso Cruz-Ramos, José Alfredo Contreras-Gutiérrez, Eliakym Arámbula-Meraz, Carla Angulo-Rojo, Alma Marlene Guadrón-Llanos, Emir Adolfo Leal-León, Dora María Cedano-Prieto, Juan Pablo Meza-Espinoza

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