

Progeria o síndrome de Hutchinson-Gilford

IMAGEN 1. MUTACIÓN GENÉTICA DE LA PROGERIA

Ageing mutation

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Progeria is a rare genetic condition in which children appear to age prematurely. The mutation is in the *LMNA* gene, which codes for a protein called lamin A. This acts as a scaffolding on the inner side of the cell nucleus

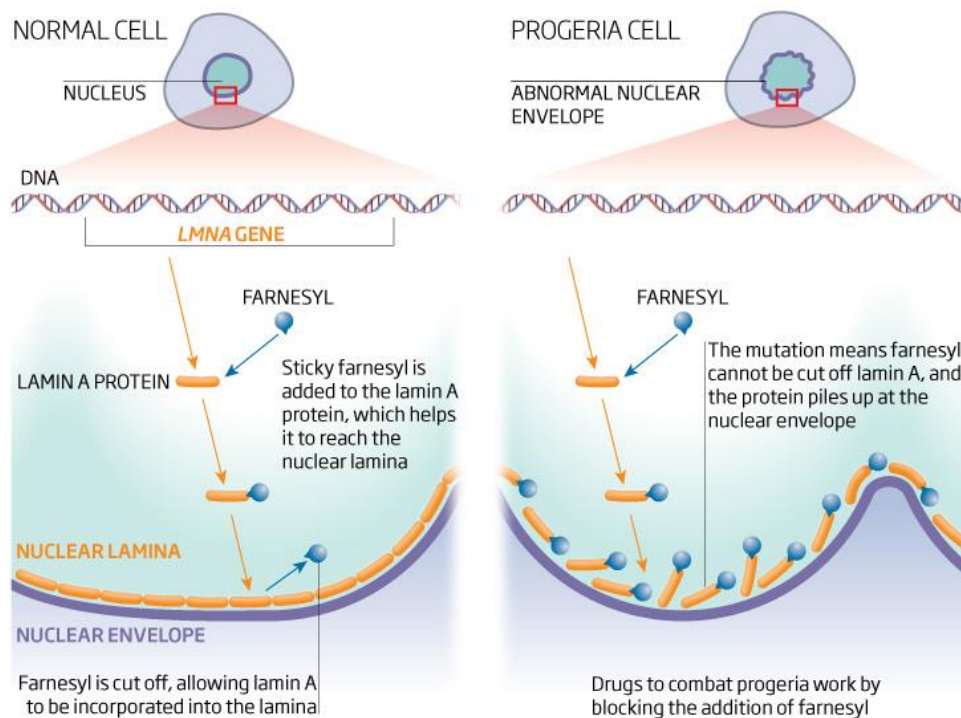


IMAGEN 2. SIGNOS DE LA PROGERIA.

