CNIC r	eceives a	donation to	investigate	Hutchinson-	Gilford
proger	ia syndro	ome			

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The Molecular and Genetic Cardiovascular Pathophysiology Laboratory headed by Dr Vicente Andrés at CNIC received a donation of 11,060 euros from the Alexandra Peraut Progeria Association for its research project into the Hutchinson–Gilford progeria syndrome (HGPS), an ultra-rare genetic disease that affects 1 in every 20 million people.

Source

 $\begin{tabular}{ll} \textbf{URL:} \underline{\textbf{https://www.cnic.es/en/noticias/cnic-receives-donation-investigate-hutchinson-gilford-progeriasyndrome} \\ \end{tabular}$