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A gain-of-function RAC2 mutation is associated with bone-marrow hypoplasia and an autosomal dominant form of severe combined immunodeficiency

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Contributions: CLP and MC supervised and designed the project, CLP analyzed and interpreted data and drafted the manuscript. AO and CT performed the biochemical studies and drafted part of the manuscript. HS, AG and CDS performed some experiments and analyzed data. PR and YC performed the protein homology modelling and drafted part of the manuscript. AF, DM, CP, JLC and MC provided patient care. CP, JLC, IA and MC helped to draft the manuscript.