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

Chantal Lagresle-Peyrou, Aurélien Olichon, Hanem Sadek, Philippe Roche, Claudine Tardy, Cindy Da Silva, Alexandrine Garrigue, Alain Fischer, Despina Moshous, Yves Collette, Capucine Picard, Jean Laurent Casanova, Isabelle André, and Marina Cavazzana

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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.