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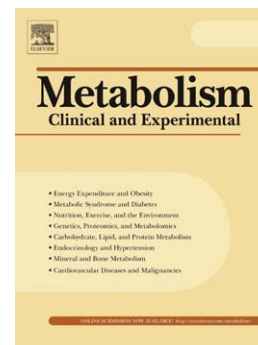
Exome sequencing reveals a *de novo* *POLD1* mutation causing phenotypic variability in mandibular hypoplasia, deafness, progeroid features, and lipodystrophy syndrome (MDPL)

Sahar Elouej, Ana Belezza-Meireles, Richard Caswell, Kevin Colclough, Sian Ellard, Jean Pierre Desvignes, Christophe Bérout, Nicolas Lévy, Shehla Mohammed, Annachiara Desandre-Giovannoli

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