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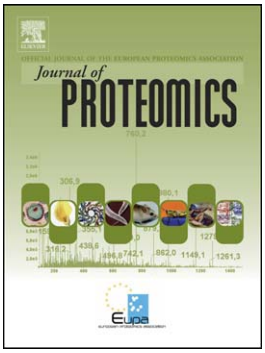

Identification of Mitochondrial Dysfunction in Hutchinson-Gilford Progeria Syndrome Through Use of Stable Isotope Labelling with Amino Acids in Cell Culture

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Identification of Mitochondrial Dysfunction in Hutchinson-Gilford Progeria Syndrome Through Use of Stable Isotope Labelling with Amino Acids in Cell Culture

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Running Title: Mitochondrial dysfunction in progeria

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