

別紙 4

研究成果の刊行に関する一覧表レイアウト (参考)

書籍

著者氏名	論文タイトル名	書籍全体の編集者名	書 籍 名	出版社名	出版地	出版年	ページ

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Suzuki E, Shima H, Kagami M, Soneda S, Tanaka T, Yatsuga S, Nishioka J, Oto Y, Kamiya T, Naiki Y, Ogata T , Fujisawa Y, Nakamura A, Kawashima S, Morikawa S, Horikawa R, Sano S, Fukami M*	(Epi)genetic defects of MKRN3 are rare in Asian patients with central precocious puberty.	<i>Hum Genome Var</i>	6	7	2019
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Hattori H, Hiura H, Kitamura A, Miyauchi N, Kobayashi N, Takahashi S, Okae H, Kyono K, Kagami M, Ogata T, Arima T.	Association of four imprinting disorders and ART.	Clin Epigenetics	11(1)	21	2019
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