別紙４

研究成果の刊行に関する一覧表

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
井原健二	他の疾患・条件に伴う糖尿病	日本小児内分泌学会編	小児内分泌疾患の治療	診断と治療社	東京	2022年	244-246

雑誌

発表者氏名	論文タイトル名	発表誌名	巻号	ページ	出版年
井原健二	遺伝病と遺伝相談：近未来の小児科のあり方	小児科 (臨時増刊号)	63(13)	1506-1512	2022
Kato H, Koshizaka M, Kaneko H, Maezawa Y, Yokote K.	Lifetime extension and the recent cause of death in Werner syndrome: a retrospective study from 2011 to 2020.	Orphanet J Rare Dis.	17(1)	226	2022
Kinoshita D, Kato H, Kaneko H, Ishikawa T, Teramoto N, Tsukagoshi A, Maeda Y, Minamizuka T, Hayashi A, Shoji M, Sawada D, Funayama S, Koshizaka M, Ogata H, Kubota Y, Mitsukawa N, Takemoto M, Yokote K, Maezawa Y.	Case of Werner syndrome with significant improvement of refractory skin ulcer despite fibroblast cellular senescence.	Case Reports Geriatr Gerontol Int.	23(3)	239-241	2023
Nagai T, Yokouchi H, Miura G, Koshizaka M, Maezawa Y, Oshitari T, Yokote K, Baba T.	Optical coherence tomography findings in three patients with Werner syndrome.	BMC Ophthalmol.	22(1)	448	2022

Sugawara S, Okada R, Loo TM, Tanaka H, Miyata K, Chiba M, Kawasaki H, Katoh K, Kaji S, Maezawa Y, Yokote K, Nakayama M, Oshima M, Nagao K, Obuse C, Nagayama S, Takubo K, Nakanishi A, Kanemaki TM, Hara E, Takahashi A.	RNaseH2A downregulation drives inflammatory gene expression via genomic DNA fragmentation in senescent and cancer cells.	Commun Biol.	5(1)	1420	2022
Miller DE, Lee L, Galey M, Kandhaya-Pillai R, Tischkowitz M, Amalnath D, Vithlani A, Yokote K, Kato H, Maezawa Y, Takada-Watanabe A, Takemoto M, Martin GM, Eichler EE, Hisama FM, Oshima J.	Targeted long-read sequencing identifies missing pathogenic variants in unsolved Werner syndrome cases.	J Med Genet.	59(11)	1087-1094	2022
Kaneko H, Kawase C, Seki J, Ikawa Y, Yachie A, Funato M.	Intellectual disability and abnormal cortical neuron phenotypes in patients with Bloom syndrome.	J Hum Genet	in press		2023
Okada Y, Funato M, Takeda S, Kaneko H.	Duchenne muscular dystrophy diagnosis using fibroblast-derived myotube cells.	Pediatr Int	doi: 10.1111/ped.15151.		2022
Kaneko H, Takemoto M, Murakami H, Ihara K, Kosaki R, Motegi SI, Taniguchi A, Matsuo M, Yamazaki N, Nishigori C, Takita J, Koshizaka M, Maezawa Y, Yokote K.	Rothmund-Thomson syndrome investigated by two nationwide surveys in Japan	Pediatr Int	doi: 10.1111/ped.15120.		2022