

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
森崎裕子	Ehlers-Danlos症候群、Marfan症候群	「小児内科」 「小児外科」 編集委員会	小児疾患診療のための病態生理2	東京医学社	東京	2021	304-308
仁科幸子	眼疾患	臨床遺伝専門医制度委員会	臨床遺伝学小児領域	診断と治療社	東京	2021	124-126
仁科幸子	視覚器の発達	" 秋山千枝子・五十嵐隆・					
岡明・平岩幹夫"	小児保健ガイドブック	診断と治療社	東京	2021	96-98		
松永達雄	クロマチンリモデリング因子異常症、CHARGE 症候群	副島英伸、秦健一郎 編集.	遺伝子医学Mook 36号 エピゲノムで新たな解明が進む「先天性疾患」	メディカルドゥ	東京	2021	151-157
松永達雄	ワルデンブルグ症候群	「小児内科」 「小児外科」 編集委員会 共編.	小児内科2021 vol. 53 小児疾患診療のための病態生理2	東京医学社	東京	2021	203-205
松永達雄	外耳奇形	大森孝一、野中学、小島博己・編集.	標準耳鼻咽喉科・頭頸部外科学 第4版	医学書院	東京	2022	52-53
松永達雄	中耳奇形	大森孝一、野中学、小島博己・編集.	標準耳鼻咽喉科・頭頸部外科学 第4版	医学書院	東京	2022	54-56
小崎里華	標準的医療説明 インフォームド・コンセントの最前線	内科系学会社会保険連合	標準的医療説明 インフォームド・コンセントの最前線	医学書院	東京	2021	264-266
黒澤健司	遺伝学的検査	秋山千枝子、五十嵐隆、岡明、平岩幹夫	小児保健ガイドブック	診断と治療社	東京	2021	197-200
黒澤健司	遺伝子診断と遺伝カウンセリング	土屋弘行	今日の整形外科治療指針 第8版	医学書院	東京	2021	235-236
齋藤伸治	CTCF関連神経発達症	副島英伸、秦健一郎	エピゲノムで新たな解明が進む「先天性疾患」	メディカルドゥ	大阪	2021	164-167

岡本伸彦	ヒストン修飾異常症 Rubinstein-Taybi症候群、エピゲノムで新たな解明が進む「先天性疾患」】		遺伝子医学MOOK	メディカルドゥ	大阪	2021	126-131
岡本伸彦	Coffin-Siris症候群とBAF複合体		遺伝子医学MOOK	メディカルドゥ	大阪	2021	96-101
岡本伸彦	Smith-Lemli-Opitz症候群	小児疾患診療のための病態生理2 改訂第6版	小児内科	東京医学社	東京	2021	296-299
西恵理子	13トリソミー症候群	小児疾患診療のための病態生理2 改訂第6版	小児内科	東京医学社	東京	2021	209-213
西恵理子	小児期の診察とマネージメント、「Dysmorphologyの診かた」		臨床遺伝専門医テキスト、臨床遺伝学小児領域	診断と治療社	東京	2021	44-49
Soejima H, Ohba T.	Chapter 11 Genomic Imprinting Disorders (Including Mesenchymal Placental Dysplasia)	Masuzaki H (ed.)	Fetal Morph Functional Diagnosis	Springer	Singapore	2021	149-168
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東元健、副島英伸.	第3章先天性疾患 3. ヒストン修飾異常症 1) Sotos症候群	副島英伸、秦健一郎編集	遺伝子医学MOOK 36 エピゲノムで新たな解明が進む「先天性疾患」	メディカルドゥ	大阪	2021	105-110
渡邊 淳	第4章 遺伝性疾患	青笹克之(監修), 加藤光保(編集), 金井弥栄(編集), 菅野祐幸(編集)	解明病理学第4版 病気のメカニズムを解く	医歯薬出版	東京	2021	79-93
渡邊 淳	13-15 先天性結合組織病	矢崎義雄(編集), 小室一成(編集)	内科学 第12版	朝倉書店	東京	2021	III-416-419

加藤光広	二分脊椎, 脳形成異常	水口雅、山形崇倫	クリニカルガイド 小児科 専門医の 診断・治療	南山堂	東京	2021	846-852
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