

## 別添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
角田和繁	小口病、三宅病、錐体杆体ジストロフィ		眼底パーフェクトアトラス	文光堂	東京	2017	
松永達雄	遺伝性難聴の診断の進歩	山崎達也	医学のあゆみBOOKS 耳鼻咽喉科診療の進歩 40 のエッセンス	医歯薬出版	東京	2018	7-10
角田和繁	遺伝性網膜疾患の臨床診断		「あたらしい眼科」『網膜変性診療の未来予想図』	医学書院		2018	427-436
仁科幸子	未熟児網膜症 眼底検査法と写真撮影法 . リハビリテーション・ロービジョンケア . 類縁疾患	東範行	未熟児網膜症	三輪書店	東京	2018	
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後藤美和子	章 基本的診療 眼科問診	松永 達雄	先天性および若年性の視覚聴覚二重障害の原因となる難病の診療マニュアル				42-45
後藤美和子	章 特記すべき診療・療育・支援 成人への移行における課題と眼科的対応	松永 達雄	先天性および若年性の視覚聴覚二重障害の原因となる難病の診療マニュアル				161-163
守本倫子	学童期までの早産児、低出生体重児の神経・発達・感覚器の評価と対応	河野由美・水野克巳	低出生体重児の成長と発達のみかた	東京医学社	東京	2019	150-156
仁科幸子	小児や障害児に適した眼鏡 デザインと装用させるコツ	日本近視学会・日本小児眼科学会・日本視能訓練士協会	小児の近視 診断と治療	三輪書店	東京	2019	P139-142

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仁科幸子	先天白内障	大橋裕一・村上晶	眼科疾患 最新の治療. 2019-2 021	南江堂	東京	2019	p195

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守本倫子	小児難聴	ファルマシア	54	1035-1039	2018
守本倫子	小児で重要な聴覚・平衡覚疾患	日本医師会雑誌	147	150-151	2018
守本倫子	胎児期感染による先天性難聴	JOHNS	34	1521-1524	2018
Suzuki N, Mutai H, Miya F, Tsunoda T, Terashima H, <u>Morimoto N</u> , Matsunaga T	A case report of reversible generalized seizures in a patient with Waardenburg syndrome associated with a novel nonsense mutation in the penultimate exon of SOX10.	BMC Pediatr.	23	doi: 10.1186/s12887-018-1139-2.	2018

Morimoto N, Mutai H, Namba K, Kaneko H, Kosaki R, Matsunaga T	Homozygous EDNRB mutation in a patient with Waardenburg syndrome type 1	Auris Nasus Larynx	45	222-226	2018
Kurata K, Hosono K, Hayashi T, Mizobuchi K, Katagiri S, Miyamichi D, <u>Nishina S</u> , Sato M, Azuma N, Nakano T, Hotta Y	X-linked retinitis pigmentosa in Japan: Clinical and genetic findings in male patients and female carriers.	Int J Mol Sci. 2019, 20, 1518; doi:10.3390/ijms20061518			2019
Yoshida T, Katagiri S, Yokoi T, <u>Nishina S</u> , Azuma N.	Optical coherence tomography and video recording of a case of bilateral contractile peripapillary staphyloma.	Am J Ophthalmol Case Rep	13	66-69	2019
Hirayama J, Alifu Y, Hamabe R, Yamaguchi S, Tomita J, Maruyama Y, Asaoka Y, Nakahama K, Tamari T, Takamatsu K, Takamatsu N, Hattori A, <u>Nishina S</u> , Azuma N, Kawahara A, Kume K, Nishina H.	The clock components Period2, Cryptochrome1a, and Cryptochrome2a function in establishing light-dependent behavioral rhythms and/or total activity levels in zebrafish	Sci Rep. 2019 Jan 9:196. doi: 10.1038/s41598-018-37879-8.			2019
Hosono K, <u>Nishina S</u> , Yokoi T, Katagiri S, Saitsu H, Kurata K, Miyamichi D, Hikoya A, Mizobuchi K, Nakano T, Minoshima S, Fukami M, Kondo H, Sato M, Hayashi T, Azuma N, Hotta Y.	Molecular diagnosis of 34 Japanese families with Leber congenital amaurosis using targeted next generation sequencing.	Sci Rep. 2018 May 29;8(1):8279. doi: 10.1038/s41598-018-26524-z.			2018
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Takahashi M, Yokoi T, Katagiri S, Yoshida-Uemura T, <u>Nishina S</u> , Azuma N.	Surgical treatments for fibrous tissue extending to the posterior retina in eyes with familial exudative vitreoretinopathy.	Jpn J Ophthalmol	62(1)	63-67	2018
吉田朋世、 <u>仁科幸子</u> 、松岡真未、萬束恭子、赤池祥子、越後貫滋子、横井匡、東範行	Information and Communication Technology 機器の使用が契機と思われた小児斜視症例.	眼臨紀	11(1)	61-66	2018

太刀川貴子, 武井正人, 清田眞理子, 斎藤雄太, 東範行, <u>仁科幸子</u> , 丸子一朗, 根岸貴志, 野田英一郎, 大熊康弘, 吉田圭, 藤巻拓郎, 松本直, 渡邊恵美子, 斎藤誠	低出生体重児における未熟児網膜症: 東京都多施設研究.	日眼会誌	112(2 )	103-113	2018
佐藤美保、加藤光広、田島敏広、川村孝、 <u>仁科幸子</u> 、根岸貴志、柿原寛子、初川嘉一、松村望、三木淳司、寺井朋子、横山利幸、森田由香、三原美晴、野村耕治、富田香、林思音、磯貝正智、堀田喜裕	中隔視神経異形成症の眼科診療に関する研究.	眼臨紀	11(5)	395-400	2018
<u>仁科幸子</u>	乳幼児の新しい視覚スクリーニング 簡便で正確な検査装置の導入	日本医師会雑誌	147(8 )	1628-1629	2018
吉田朋世、 <u>仁科幸子</u>	主訴と所見からみた眼科 common disease、内斜視	眼科	60	1157-1162	2018
Hayakawa I, Kubota M.	Digital Amputation by Congenital Insensitivity to Pain with Anhidrosis.	The Journal of Pediatrics		doi: <a href="https://doi.org/10.1016/j.jpeds.2019.01.023">https://doi.org/10.1016/j.jpeds.2019.01.023</a>	2019
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Minami S, Nara K, Mutai H, Morimoto N, Sakamoto H, Takiguchi T, Kaga K, <u>Matsunaga T</u>	A clinical and genetic study of 16 Japanese families with Waardenburg syndrome	Gene	708(1 )	86-90	2019

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山本修子、南修司郎、榎本千江子、加藤秀敏、 <u>松永達雄</u> 、伊藤文展、遠藤理奈子、橋本陽介、石川直明、加我君孝	東京医療センターにおける成人人工内耳症例の適応と有用性の検討	日本耳鼻咽喉科学会会報	122(8)	1118-1126	2019
松永達雄	視覚聴覚二重障害の診療	JOHNS	35 (9)	1377-1378	2019
松永達雄	遺伝情報をどう管理し、活用するか 耳科領域	JOHNS	35(10)	1452-1454	2019
Akiko Maeda-Katahira, Natsuko Nakamura, Takaaki Hayashi, Satoshi Katagiri, Satoko Shimizu, Hisao Ohde, Tatsuo Matsunaga, Kimitaka Kaga, Tadashi Nakano, Shuhei Kameya, Tomokazu Matsuura, Kaoru Fujinami, Takeshi Iwata, Kazushige Tsunoda	Autosomal dominant optic atrophy with OPA1 gene mutations accompanied by auditory neuropathy and other systemic complications in a Japanese cohort	Molecular Vision	25	559-573	2019
Nakamura N, Tsunoda K, Mizuno Y, Usui T, Hatase T, Ueno S, Kuniyoshi K, Hayashi T, Katagiri S, Kondo M, Kameya S, Yoshitake K, Fujinami K, Iwata T, Miyake Y	Clinical Stages of Occult Macular Dystrophy Based on Optical Coherence Tomographic Findings.	Invest Ophthalmol Vis Sci	Nov 1;60(14)	4691-4700	2019
角田和繁	網脈絡膜ジストロフィの遺伝学的病態解明および治療に向けた症例データバンクの構築	日眼会誌			2020
Fujinami K* † . Yang L*, Joo K* et al.	Clinical and Genetic Characteristics of East Asian Patients with Occult Macular Dystrophy (Miyake Disease): East Asia Occult Macular Dystrophy Studies Report Number 1.	Ophthalmology.	126(10)	1432-1444	2019

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Kameya S*, Fujinami K,* Ueno S et al.	Phenotypical Characteristics of POC1B-Associated Retinopathy in Japanese Cohort: Cone Dystrophy With Normal Funduscopic Appearance.	Invest Ophthalmol Vis Sci.	60(10)	3432-3446.	2019
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Katagiri S, Hayashi T, Nakamura M,..., Fujinami K, et al.	RDH5-related fundus alipunctatus in a large Japanese cohort.	Invest Ophthalmol Vis Sci.	In press	In press	In press
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Pontikos N,..., Fujinami K, Fujinami-Yokokawa Y et al.	Phenogenon: Gene to Phenotype Associations for Rare Genetic Diseases.	PLoS One.	In press	In press	In press
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<u>加我君孝</u>	Auditory Neuropathy	JOHNS	36	61-66	2020
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柳澤瞳、今井裕弥子、 <u>守本倫子</u>	2012年から2013年の風疹流行に伴う先天性風疹症候群4例の経過と言語指導について。	小児耳	40 ( 3 )	271-277	2019

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Yoshida T, Katagiri S, Yokoi T, <u>Nishina</u> S, Azuma N	Optical coherence tomography and video recording of a case of bilateral contractile peripapillary staphyloma.	Am J Ophthalmol	13	66-69	2019
Hirayama1 J, Alifu Y, Hamabe R, Yamaguchi S, Tomita J, Maruyama Y, Asaoka Y, Nakahama K, Tamari T, Takamatsu K, Takamatsu N, Hattori A, <u>Nishina</u> S, Azuma N, Kawahara A, Kume K, Nishina H.	The clock components Period2, Cryptochrome1a, and Cryptochrome2a function in establishing light-dependent behavioral rhythms and/or total activity levels in zebrafish.	Sci Rep		Jan :196; doi: 10.1038/s41598-018-37879-8.	2019

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中尾志郎、 <u>仁科幸子</u> 、八木瞳、田中慎、吉田朋世、横井匡、東範行	外直筋鼻側移動術を施行した動眼神経麻痺の一例.	眼臨紀	13 (2)	105-110	2020
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林思音、 <u>仁科幸子</u> 、森隆史、清水ふき、南雲幹、臼井千恵、杉山能子、八子恵子	三歳児眼科健診における屈折検査の有用性:システムマティックレビュー.	眼臨紀	12 (5)	373-377,	2019
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<u>仁科 幸子</u>	乳幼児の視覚スクリーニング	日本の眼科	90 (10)	1291-1292,	2019
<u>仁科 幸子</u>	乳幼児の視覚スクリーニング	東京小児科医会報	38 (1)	63-69	2019
<u>仁科 幸子</u> ・佐藤美保	序説 弱視と斜視のカレントトピックス	あたらしい眼科	36 (8)	971-972	2019
吉田朋世・ <u>仁科幸子</u>	急性後天性共同性内斜視. 特集 弱視と斜視のカレントトピックス	あたらしい眼科	36 (8)	995-1001	2019
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<u>仁科 幸子</u>	レーバー先天盲. ~知つておきたい稀な網膜・硝子体ジストロフィー	オクリスタ	75 (6)	31-37	2019
<u>仁科 幸子</u>	手持ちフォトスクリーナー装置、乳幼児期の眼鏡・コンタクトレンズ	チャイルドヘルス	22 (6)	21-23, 47-49	2019

Hayakawa I, Abe Y, Ono H, <u>Kubota M</u>	Severe congenital RYR1-associated myopathy complicated with atrial tachycardia and sinus node dysfunction: a case report.	Ital J Pediatr	45	doi: 10.1186/s13052-019-0756-1	2019
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6 . 研究成果による特許権等の知的財産権の出願・登録状況  
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