

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
大園恵一	大理石骨病		今日の整形外科治療指針	医学書院	東京	2016	251.
芳賀信彦	FGFR3異常症(軟骨無形成症、軟骨低形成症、タナトフオリック骨異形成症), II型コラーゲン異常症(先天性脊椎骨端異形成症、Kniest骨異形成症、Stickler症候群1型)。	日本小児整形外科学会	小児整形外科テキスト、第2版	メジカルビュー社	東京	2016	255-256
芳賀信彦	先天性骨系統疾患、先天異常症候群。	中村利孝、松野丈夫	標準整形外科学	医学書院	東京	2017	307-316
鬼頭浩史	FGFR3グループ概説 軟骨無形成症 軟骨低形成症 濃化異骨症 弯曲骨異形成症概説		今日の整形外科治療指針 7版)	医学書院	東京	2016	231-232, 232-233, 233-234, 252-253, 249-250,

雑誌

(本研究と密接の関連のある論文のみを抜粋)

発表者氏名	論文タイトル名	発表誌名	巻号	ページ	出版年
Matsushita M, Kitoh H, Mishima K, Yamashita S, Haga N, Fujiwara S, Ozono K, Kubota T, Kitaoka T, Ishiguro N.	Physical, mental and social problems of adolescent and adult patients with achondroplasia.	Calcif Tissue Int	104(4):	364-372,	2019

Nakahara Y, Kitoh H, Nakashima Y, Toguchida J, Haga N.	Longitudinal study of the activities of daily living and quality of life in Japanese patients with fibrodysplasia ossificans progressiva.	Disabil Rehabil	41(6):	699-704,	2019
Yamamoto K, Kawawai M, Yamazaki M, Tachikawa K, Kubota T, Ozono K, Michigami T.	CREB activation in hypertrophic chondrocytes is involved in the skeletal overgrowth in epiphyseal chondrodysplasia Miura type caused by activating mutations of natriuretic peptide receptor B.	Hum Mol Genet	28(7):	1183-1198,	2019
Offiah AC, Vockley J, Munns CF, Murotsuki J;	Differential diagnosis of perinatal hypophosphatasia: radiologic perspectives.	Pediatr Radiol	49:	3-22	2019
Oda T, Matsushita M, Ono Y, Kitoh H, Sakai T.	A novel heterozygous mutation in the T-box protein 4 gene in an adult case of small patella syndrome.	J Orthop Case Rep	8(1):	85-88,	2018
Okura T, Matsushita M, Mishima K, Esaki R, Seki T, Ishiguro N, Kitoh H.	Activated FGFR3 prevents subchondral bone sclerosis during the development of osteoarthritis in transgenic mice with achondroplasia.	J Orthop Res	36(1):	300-308,	2018
Hasegawa S, Kitoh H, Matsushita M, Mishima K, Kadono I, Susukiura H, Kitamura A, Ishiguro N.	Chronic lateral epiphyseal separation of the proximal tibia causes late-onset tibial vara.	J Pediatr Orthop B	27(1):	31-34,	2018

Akiyama T, Kubota T, Ozono K, Michigami T, Kobayashi D, Takeyari S, Sugiyama Y, Nodamura M, Harada D, Namba N, Suzuki A, Utoyama M, Kitanaka S, Uematsu M, Mitani Y, Matsunami K, Takishima S, Ogawa E, Kobayashi K.	Pyridoxal 5'-phosphate and related metabolites in hypophosphatasia: Effects of enzyme replacement therapy.	Molecular Genetics and Metabolism ,	125(1- 2),	174-180 ,	2018
Kashii M, Kanao S, Kitaoka T, Makino T, Kaito T, Iwasaki M, Kubota T, Yamamoto T, Ozono K, Yoshikawa H.	Development of scoliosis in young children with osteogenesis imperfecta undergoing intravenous bisphosphonate therapy.	J Bone Miner Metab ,	37(3):	545-553.	2018
Harada A, Miyashita S, Nagai R, Makino S, Murotsuki J	Prenatal sonographic findings and prognosis of craniosynostosis diagnosed during the fetal and neonatal periods.	Congenit Anom (Kyoto)		doi: 10.1111/1/cga.12308	2018

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Takahashi Y, Sawawai H, Murotsuki J, Satoh S, Yamada T, Hatayakawa H, Kouzuma Y, Saseda M, Watanabe A, Miyazaki O, Nishimura G.	Parental serum alkaline phosphatase activity as an auxiliary tool for prenatal diagnosis of hypophosphatasia.	Prenat Diagn	37(5)	491-496.	2017

Miyazaki O, Sawai H, Yamada T, Murotsuki J, Nishimura G.	Follow-Up Study on Fetal CT Radiation Dose in Japan: Validating the Decrease in Radiation Dose.	Am J Roentgenol.	208(4)	862-867.	2017
室月 淳	出生前診断は周産期医療をどのようにかえたか - 骨系統疾患を例として	日本周産期・新生児医学会雑誌	53	452	2017
室月 淳	胎児骨系統疾患の遺伝診療	産科と婦人科	84:	29-34	2017
Kusano C, Takagi M, Hori N, Murotsuki J, Nishimura G, Hasegawa T	A novel mutation in the C-propeptide of COL2A1 causes atypical spondyloepiphyseal dysplasia congenita.	Hum Genome Var.	2(4)	17003	2017
Imai 4, Miyazaki O, Horiuchi T, Asano K, Nishimura G, Sago H, Nosaka S.	Ultra-Low-Dose Fetal CT with Model-Based Iterative Reconstruction: A Prospective Pilot Study	Am J Roentgenol.	208	1-8	2017;
Kajita S, Yamamoto T, Tsugawa N, Nakayama H, Kubota T, Michigami T, Ozono K.	Serum calcitriol levels in a patient with X-linked hypophosphatemia complicated by autosomal dominant polycystic kidney disease.	CEN Case Report	6:	29-35,	2017.
Ueyama K, Namba N, Kitaoka T, Yamamoto K, Fujiwara M, Ohata Y, Kubota T, Ozono	K. Endocrinological and phenotype evaluation in a patient with acrodysostosis.	Clin Pediatric Endocrinol.	26(3):	177-182,	2017

Kitaoka T , Tajima T , Nagasaki K , Kikuchi T, Yamamoto K , Michigami T , Okada S , Fujiwara I , Kokaji M , Mochizuki H , Ogata T , Tatebayashi K , Watanabe A , Yatsuga S , Kubota T , Ozono K.	Safety and efficacy of treatment with asfotase alfa in patients with hypophosphatasia: Results from a Japanese clinical trial.	Clin Endocrinol (Oxf),	87(1):	10-19,	2017.
日本整形外科学会小児整形外科学委員会、骨系統疾患国際分類和訳作業WG、小崎慶介、北野利夫、鬼頭浩史、中島康晴、北中幸子、室月淳、西村玄、芳賀信彦:	2015年版骨系統疾患国際分類の和訳.	日整会誌	91(7):	462-505,	2017
芳賀信彦	:骨系統疾患と装具。(治療効率をあげる運動器装具療法のコツ)	Monthly Book Orthopaedics	30(6):	69-73,	2017
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Nakahara Y, Kitoh H, Nakashima Y, Toguchida J, Haga N:	The longitudinal study of activities of daily living and quality of life in Japanese patients with fibrodysplasia ossificans progressiva.	Disabil Rehabil,	Nov 16:1-6 [Epub ahead of print]		2017

Matsushita M, Mishima K, Esaki R, Ishiguro N, Ohno K, Kitoh H.	Maternal administration of meclozine for the treatment of foramen magnum stenosis in transgenic mice with achondroplasia.	J Neurosurg Pediatr:	19(1)	91-95,	2017
Osawa Y, Matsushita M, Hasegawa S, Esaki R, Fujio M, Ohgasawara B, Ishiguro N, Ohno K, Kitoh H.	Activated FGFR3 promotes bone formation via accelerating endochondral ossification in mouse model of distraction osteogenesis.	Bone	105:	42-49,	2017
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