

II 研究成果の刊行物リスト

書籍(日本語)

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
阿部 純也, 西小森 隆太, 平家 俊男	Aicardi-Goutieres症候群(解説/特集)	宮坂伸之	別冊日本臨床 免疫症候群(第2版)III	日本臨床社	大阪	2016	544-550
西小森 隆太, 西村 玄, 平家 俊男	Spondyloenchondrolyplasia with immune dysregulation (SPENCDI)	宮坂伸之	別冊日本臨床 免疫症候群(第2版)III	日本臨床社	大阪	2016	551-553
田中 孝之, 西小森 隆太, 平家 俊男	メバロン酸キナーゼ欠損症(高IgD症候群)	宮坂伸之	別冊日本臨床 免疫症候群(第2版)III	日本臨床社	大阪	2016	779-78
井澤 和司, 西小森 隆太, 平家 俊男	PLCG2異常症	宮坂伸之	別冊日本臨床 免疫症候群(第2版)III	日本臨床社	大阪	2016	783-786
井澤 和司, 河合 朋樹, 西小森 隆太, 平家 俊男	SAVI	宮坂伸之	別冊日本臨床 免疫症候群(第2版)III	日本臨床社	大阪	2016	831-833
河合 朋樹, 平家 俊男	免疫不全を伴う無汗性外胚葉形成異常症(EDA-ID) NEMO異常症とNFKBIA異常症	宮坂伸之	別冊日本臨床 免疫症候群(第2版)III	日本臨牀社	大阪	2016	304-310
河合 朋樹, 平家 俊男	LUBAC異常症(HOIL-1欠損症とHOIP欠損症)	宮坂伸之	別冊日本臨床 免疫症候群(第2版)III	日本臨床社	大阪	2016	311-316
大西秀典	MCM4欠損症、重症ウイルス感染症易感染疾患	宮坂伸之	別冊日本臨床 免疫症候群(第2版)III	日本臨床社	大阪	2016	216-218
大西秀典	単純ヘルペス脳炎易感染疾患	宮坂伸之	別冊日本臨床 免疫症候群(第2版)III	日本臨床社	大阪	2016	687-708
金澤伸雄	中條 - 西村症候群	宮坂伸之	別冊日本臨床 免疫症候群(第2版)III	日本臨牀社	大阪	2016	825-830
岡野翼, 今井耕輔, 金兼弘和	γc欠損症(X連鎖重症複合免疫不全症)	宮坂伸之	別冊日本臨床 免疫症候群(第2版)III	日本臨牀社	大阪	2016	16-19
山下基, 今井耕輔, 金兼弘和	JAK3欠損症	宮坂伸之	別冊日本臨床 免疫症候群(第2版)III	日本臨牀社	大阪	2016	20-24
大川哲平, 今井耕輔	CD3 欠損症	宮坂伸之	別冊日本臨床 免疫症候群(第2版)III	日本臨牀社	大阪	2016	87-89
岡本圭祐, 今井耕輔, 金兼弘和	Omenn症候群	宮坂伸之	別冊日本臨床 免疫症候群(第2版)III	日本臨牀社	大阪	2016	175-178
岡本圭祐, 今井耕輔	DOCK2欠損症 /Coronin-1A欠損症/CD3 欠損症	宮坂伸之	別冊日本臨床 免疫症候群(第2版)III	日本臨牀社	大阪	2016	76-79/43-47/39-42
田中桂輔, 今井耕輔	CD3 epsilon欠損症/CD45欠損症/CD3 欠損症	宮坂伸之	別冊日本臨床 免疫症候群(第2版)III	日本臨牀社	大阪	2016	36-38/29-32/33-35
金澤伸雄	Blau症候群	杉山幸比古	呼吸器科医のためのサルコイドーシス診療ガイド	南江堂	東京	2016	118-122
金澤伸雄	結節性紅斑	山口徹、北原光男	今日の治療指針2016年版	医学書院	東京	2016	1242-1243
右田清志	全身性疾患に伴う関節炎 3 悪性腫瘍、血液疾患 4 アミロイドーシス	日本リウマチ財団、日本リウマチ学会	リウマチ病学テキスト 改訂第2版	診断と治療社	東京	2016	365-373

学会誌・雑誌等における論文掲載（英語）

発表者名	論文タイトル名	発表雑誌	巻号	ページ	出版年
Shunpei Y, Imagawa T, Nishikomori R, Takada H, Abrams K, Lheritier K, Heike T, Hara T.	Long-term safety and efficacy of canakinumab in cryopyrin-associated periodic syndrome: results from an open-label, phase III pivotal study in Japanese patients.	Clin Exp Rheumatol.	印刷中	印刷中	2017
Eroglu FK, Kasapcopur O, Beşbaş N, Ozaltın F, Bilginer Y, Barut K, Mensa-Vilaro A, Nakagawa K, Heike T, Nishikomori R, Aróstegui J, Ozan S.	Genetic and clinical features of cryopyrin-associated periodic syndromes in Turkish children.	Clin Exp Rheumatol.	102(6)	115-120	2016
Kawasaki Y, Oda H, Ito J, Niwa A, Tanaka T, Hijikata A, Seki R, Nagashashi A, Osawa M, Asaka I, Watanabe A, Nishimata S, Shirai T, Kawashima H, Ohara O, Nakahata T, Nishikomori R, Heike T, Saito MK.	Identification of a High-Frequency Somatic NLRC4 Mutation as a Cause of Autoinflammation by Pluripotent Cell-Based Phenotype Dissection.	Arthritis Rheumatol.	印刷中	印刷中	2017
Imamura S, Narita S, Nishikomori R, Tsuruta H, Numakura K, Maeno A, Saito M, Inoue T, Tsuchiya N, Nanjo H, Heike T, Satoh S, Habuchi T.	Secondary bladder amyloidosis with familial Mediterranean fever in a living donor kidney transplant recipient: a case report.	BMC Res Notes.	9(1)	473	2016
Nakashima H, Miyake F, Ohki S, Hattori S, Matsubayashi T, Izawa K, Nishikomori R, Heike T, Honda Y, Shigematsu Y.	Febrile attacks triggered by milk allergy in an infant with mevalonate kinase deficiency.	Rheumatol Int.	36(10)	1477-8	2016
Iwasaki T, Kaneko N, Ito Y, Takeda H, Sawasaki T, Heike T, Migita K, Agematsu K, Kawakami A, Morikawa S, Mokuda S, Kurata M, Masumoto J.	Nod2-Nodosome in a Cell-Free System: Implications in Pathogenesis and Drug Discovery for Blau Syndrome and Early-Onset Sarcoidosis.	ScientificWorldJournal.	2016	2597376	2016
Mensa-Vilaro A, Teresa Bosque M, Magri G, Honda Y, Martínez-Banaclocha H, Casorran-Berges M, Sintes J, González-Roca E, Ruiz-Ortiz E, Heike T, Martínez-García JJ, Baroja-Mazo A, Cerutti A, Nishikomori R, Yagüe J, Pelegrín P, Delgado-Beltran C, Aróstegui JI.	Brief Report: Late-Onset Cryopyrin-Associated Periodic Syndrome Due to Myeloid-Restricted Somatic NLRP3 Mosaicism.	Arthritis Rheumatol.	68(12)	3035-3041	2016
Oda H, Sato I, Kunishima S, Nakagawa K, Izawa K, Hiejima E, Kawai T, Yasumi T, Doi H, Katanura K, Numabe H, Okamoto S, Nakase H, Hijikata A, Ohara O, Suzuki H, Morisaki H, Morisaki T, Nunoi H, Hattori S, Nishikomori R, Heike T.	Exon skipping causes atypical phenotypes associated with a loss-of-function mutation in FLNA by restoring its protein function.	Eur J Hum Genet.	24(3)	408-14	2016
Takada H, Ishimura M, Hara T. Bone Marrow Transplant.	Insufficient immune reconstitution after allogeneic cord blood transplantation without chemotherapy conditioning in patients with SCID caused by CD3 deficiency.	Bone Marrow Transplant	51(8)	1131-3	2016
Nanishi E, Hoshina T, Takada H, Ishimura M, Nishio H, Uehara T, Mizuno Y, Hasegawa S, Ohga S, Nagao M, Igarashi M, Yajima S, Kusumoto Y, Onishi N, Sasahara Y, Yasumi T, Heike T, Hara T; PID-Infection Study Group.	A nationwide survey of common viral infections in childhood among patients with primary immunodeficiency diseases.	J Infect	73(4)	358-68	2016
Takada H, Ishimura M, Takimoto T, Kohagura T, Yoshikawa H, Imaizumi M, Shichijyou K, Shimabukuro Y, Kise T, Hyakuna N, Ohara O, Nonoyama S, Hara T.	Invasive bacterial infection in patients with interleukin-1 receptor-associated kinase 4 deficiency: Case report.	Medicine (Baltimore).	95(4)	e2437	2016
Yamamoto H, Ishimura M, Ochiai M, Takada H, Kusuvara K, Nakatsu Y, Tsuzuki T, Mitani K, Hara T.	BTK gene targeting by homologous recombination using a helper-dependent adenovirus/adeno-associated virus hybrid vector.	Gene Ther	23(2)	205-13	2016
Ito N, Hataya H, Saida K, Amano Y, Hidaka Y, Motoyoshi Y, Ohta T, Yoshida Y, Terano C, Iwasa T, Kubota W, Takada H, Hara T, Fujimura Y, Ito S.	Efficacy and safety of eculizumab in childhood atypical hemolytic uremic syndrome in Japan.	Clin Exp Nephrol	20(2)	265-72	2016
Teranishi H, Ishimura M, Koga Y, Eguchi K, Sonoda M, Kobayashi T, Shiraishi S, Nakashima K, Ikegami K, Aman M, Yamamoto H, Takada H, Ohga S.	Activated phosphoinositide 3-kinase syndrome presenting with gut-associated T-cell lymphoproliferative disease.	Rinsho Ketsueki	58(1)	20-25	2017

Kagawa R, Fujiki R, Tsumura M, Sakata S, Nishimura S, Itan Y, Kong XF, Kato Z, Ohnishi H, Hirata O, Saito S, Ikeda M, El Baghdadi J, Bousfiha A, Fujiwara K, Oleastro M, Yancoski J, Perez L, Danielian S, Ailal F, Takada H, Hara T, Puel A, Boisson-Dupuis S, Bustamante J, Casanova JL, Ohara O, Okada S, Kobayashi M.	Alanine-scanning mutagenesis of human signal transducer and activator of transcription 1 to estimate loss- or gain-of-function variants.	J Allergy Clin Immunol	S0091-6749(16)3	1281-7	2016
Hori M, Yasumi T, Shimodera S, Shibata H, Hiejima E, Oda H, Izawa K, Kawai T, Ishimura M, Nakano N, Shirakawa R, Nishikomori R, Takada H, Morita S, Horiuchi H, Ohara O, Ishii E, Heike T.	A CD57 ⁺ CTL Degranulation Assay Effectively Identifies Familial Hemophagocytic Lymphohistiocytosis Type 3 Patients.	J Clin Immunol	37(1)	92-99.	2017
Ueki M, Yamada M, Ito K, Tozawa Y, Morino S, Horikoshi Y, Takada H, Abdrabou SS, Takezaki S, Kobayashi I, Ariga T.	A heterozygous dominant-negative mutation in the coiled-coil domain of STAT1 is the cause of autosomal-dominant Mendelian susceptibility to mycobacterial diseases.	Clin Immunol	14;174	24-31	2016
Wakamiya T, Hokosaki T, Tsujimoto SI, Kadota K, Nakano Y, Watanabe S, Iwamoto M, Yanagimachi M, Ito S.	Effect of VKORC1, CYP2C9, CYP4F2, and GGCX Gene Polymorphisms on Warfarin Dose in Japanese Pediatric Patients.	Mol Diagn Ther.	20(4)	393-400	2016
Nozawa T, Nishimura K, Ohara A, Hara R, Ito S	Primary varicella infection in children with systemic juvenile idiopathic arthritis under tocilizumab therapy	Mod Rheumatol.	14	1-5	2016
Ohnishi H, Kawamoto N, Seishima M, Ohara O, Fukao T.	A Japanese family case with juvenile onset Behcet's disease caused by TNFAIP3 mutation.	Allergol Int.	66	146-148	2017
Mizutani Y, Okano T, Takahashi T, Ohnishi H, Ohara O, Sano A, Seishima M.	Pyoderma Gangrenosum, Acne and Suppurative Hidradenitis Syndrome Treated with Granulocyte and Monocyte Adsorption Apheresis.	Acta Derm Venereol.		in press	
Ueno HM, Kato T, Ohnishi H, Kawamoto N, Kato Z, Kaneko H, Kondo N, Nakano T.	T-cell epitope-containing hypoallergenic -lactoglobulin for oral immunotherapy in milk allergy.	Pediatr Allergy Immunol.	27	818-824	2016
Tsujita Y, Mitsui-Sekinaka K, Imai K, Yeh TW, Mitsuiki N, Asano T, Ohnishi H, Kato Z, Sekinaka Y, Zaha K, Kato T, Okano T, Takashima T, Kobayashi K, Kimura M, Kunitsu T, Maruo Y, Kanegae H, Takagi M, Yoshida K, Okuno Y, Muramatsu H, Shiraishi Y, Chiba K, Tanaka H, Miyano S, Kojima S, Ogawa S, Ohara O, Okada S, Kobayashi M, Morio T, Nonoyama S.	PTEN mutation can cause Activated PI3 Kinase Delta Syndrome (APDS)-like immunodeficiency.	J Allergy Clin Immunol.	138	1672-1680	2016
Sotoma S, Iimura J, Igarashi R, Hirosewa KM, Ohnishi H, Mizukami S, Kikuchi K, Fujiwara TK, Shirakawa M, Tochio H.	Selective Labeling of Proteins on Living Cell Membranes Using Fluorescent Nanodiamond Probes.	Nanomaterials	6	56	2016
Yamazaki T, Shigemura T, Kobayashi N, Honda K, Yazaki M, Masumoto J, Migita K, and Agematsu K.	IL-18 serum concentration is markedly elevated in typical familial Mediterranean fever with M694I mutation and can distinguish it from atypical type.	Mod Rheumatol.	26(2)	315-7	2016
Yamagami K, Nakamura T, Nakamura R, Hanioka Y, Seki K, Chiba H, Kobayashi K, and Agematsu K.	Familial Mediterranean fever with P369S/R408Q exon3 variant in pyrin presenting as symptoms of PFAPA.	Mod Rheumatol.	27(2)	356-359	2016
Shigemura T, Kaneko N, Kobayashi N, Kobayashi K, Takeuchi Y, Nakano N, Masumoto J, and Agematsu K.	Novel heterozygous C243Y A20/TNFAIP3 gene mutation is responsible for chronic inflammation in autosomal-dominant Behcet's disease.	RMD open.	2(1)	e000223	2016
Migita K, Izumi Y, Jiuchi Y, Iwanaga N, Kawahara C, Agematsu K, Yachie A, Masumoto J, Fujikawa K, Yamasaki S, et al.	Familial Mediterranean fever is no longer a rare disease in Japan.	Arthritis research & therapy.	30(18)	175	2016
Koga T, Migita K, Sato S, Umeda M, Nonaka F, Kawashiri SY, Iwamoto N, Ichinose K, Tamai M, Nakamura H, et al.	Multiple Serum Cytokine Profiling to Identify Combinational Diagnostic Biomarkers in Attacks of Familial Mediterranean Fever.	Medicine (Baltimore)	95(16)	e3449	2016
Iwasaki T, Kaneko N, Ito Y, Takeda H, Sawasaki T, Heike T, Migita K, Agematsu K, Kawakami A, Morikawa S, et al.	Nod2-Nodosome in a Cell-Free System: Implications in Pathogenesis and Drug Discovery for Blau Syndrome and Early-Onset Sarcoidosis.	TheScientificWorldJourn al.	2016	2597376	2016
Hokibara S, Kobayashi N, Kobayashi K, Shigemura T, Nagumo H, Takizawa M, Yamazaki T, and Agematsu K.	Markedly elevated CD64 expression on neutrophils and monocytes as a biomarker for diagnosis and therapy assessment in Kawasaki disease.	Inflammation research	65(7)	579-85	2016
Nishiguchi M, Furukawa F, Kanazawa N	Leprosy versus sarcoidosis: different diagnosis and review of misdiagnosed cases.	J Dermatol Clin Res	4	1087	2016

Kanazawa N, Tchernev G, Chokoeva AA, Maximov GK, Wollina U, Lotti T, Patterson JW, Guarneri C, Tana C, Furukawa F	Interstitial granulomatous dermatitis demonstrating small, discrete skin-colored papules.	J Biol Regul Homeost Agents	30	49-52	2016
Kamio Y, Kanazawa N, Mine Y, Utani A	Intractable leg ulcers in Blau syndrome.	J Dermatol	43	1096-1097	2016
Harada J, Nakajima T, Kanazawa N	A case of Blau syndrome with NOD2 E383K mutation.	Pediatr Dermatol	33	e385-e387	2016
Sakai T, Izumi M, Kumagai K, Kidera K, Yamaguchi T, Asahara T, Kozuru H, Jiuchi Y, Mawatari M, Osaki M, Motokawa S, Migita K.	Effects of a Foot Pump on the Incidence of Deep Vein Thrombosis After Total Knee Arthroplasty in Patients Given Edoxaban: A Randomized Controlled Study.	Medicine (Baltimore).	95(1)	e2247	2016
Bito S, Migita K, Nakamura M, Shinohara K, Sato T, Tonai T, Shimizu M, Shibata Y, Kishi K, Kubota C, Nakahara S, Mori T, Ikeda K, Ota S, Minamizaki T, Yamada S, Shiota N, Kamei M, Motokawa S.	Mechanical prophylaxis is a heparin-independent risk for anti-platelet factor 4/heparin antibody formation after orthopedic surgery.	Blood.	127(8)	1036-43	2016
Maeda Y, Migita K, Higuchi O, Mukaino A, Furukawa H, Komori A, Nakamura M, Hashimoto S, Nagaoka S, Abiru S, Yatsuhashi H, Matsuo H, Kawakami A, Yasunami M, Nakane S.	Association between Anti-Ganglionic Nicotinic Acetylcholine Receptor (GACnR) Antibodies and HLA-DRB1 Alleles in the Japanese Population.	PLoS One.	11(1)	e0146048	2016
Hirayama K, Iwanaga N, Izumi Y, Yoshimura S, Kurohama K, Yamashita M, Takahata T, Oku R, Ito M, Kawakami A, Migita K.	A Case of Relapsing Polychondritis Initiating with Unexplained Fever.	Case Rep Med.	2016	9462489	2016
Oka S, Furukawa H, Shimada K, Sugii S, Hashimoto A, Komiyama A, Fukui N, Suda A, Tsunoda S, Ito S, Katayama M, Nakamura T, Saisho K, Sano H, Migita K, Nagaoka S, Tsuchiya N, Tohma S.	Association of human leukocyte antigen alleles with chronic lung diseases in rheumatoid arthritis.	Rheumatology (Oxford).	55(7)	1301-7	2016
Koga T, Migita K, Sato S, Umeda M, Nonaka F, Kawashiri SY, Iwamoto N, Ichinose K, Tamai M, Nakamura H, Origuchi T, Ueki Y, Masumoto J, Agematsu K, Yachie A, Yoshiura K, Eguchi K, Kawakami A.	Multiple Serum Cytokine Profiling to Identify Combinational Diagnostic Biomarkers in Attacks of Familial Mediterranean Fever.	Medicine (Baltimore).	95(16)	e3449	2016
Furukawa H, Oka S, Kawasaki A, Shimada K, Sugii S, Matsushita T, Hashimoto A, Komiyama A, Fukui N, Kobayashi K, Osada A, Ihata A, Kondo Y, Nagai T, Setoguchi K, Okamoto A, Okamoto A, Chiba N, Suematsu E, Kono H, Katayama M, Hirohata S, Sumida T, Migita K, Hasegawa M, Fujimoto M, Sato S, Nagaoka S, Takehara K, Tohma S, Tsuchiya N.	Human Leukocyte Antigen and Systemic Sclerosis in Japanese: The Sign of the Four Independent Protective Alleles, DRB1*13:02, DRB1*14:06, DQB1*03:01, and DPB1*02:01.	PLoS One.	11(4)	e0154255	2016
Mori S, Hidaka M, Kawakita T, Hidaka T, Tsuda H, Yoshitama T, Migita K, Ueki Y.	Factors Associated with Myelosuppression Related to Low-Dose Methotrexate Therapy for Inflammatory Rheumatic Diseases.	PLoS One.	11(4)	e0154744	2016
Ashida M, Koike Y, Kuwatsuka S, Ichinose K, Migita K, Sano S, Utani A.	Psoriasis-like lesions in a patient with familial Mediterranean fever.	J Dermatol.	43(3)	314-7	2016
Izumi Y, Nakaoka K, Kamata M, Iwanaga N, Imadachi S, Kurohama H, Ito M, Migita K.	Steroid-resistant protein-losing gastroenteropathy complicated with Sjögren's syndrome successfully treated with mizoribine.	Mod Rheumatol.		1-5	2016
Tsurukawa S, Iwanaga N, Izumi Y, Shirakawa A, Kawahara C, Shikuwa T, Inamoto M, Kawakami A, Migita K.	Herpes Zoster Meningitis Complicating Combined Tocilizumab and Cyclosporine Therapy for Adult-Onset Still's Disease.	Case Rep Rheumatol.	2016	4232657	2016
Koga T, Fukushima C, Umeda M, Migita K, Kawakami A.	Familial Mediterranean fever complicated with refractory asthma: Successful treatment with colchicine.	Mod Rheumatol.		1-2	2016
Umeda M, Migita K, Ueki Y, Nonaka F, Aramaki T, Terada K, Koga T, Ichinose K, Eguchi K, Kawakami A.	<Letter to the Editor> A Japanese familial Mediterranean fever patient with a rare G632S MEFV mutation in exon 10.	Mod Rheumatol.		1-2	2016
Maeda Y, Nakane S, Higuchi O, Nakamura H, Komori A, Migita K, Mukaino A, Umeda M, Ichinose K, Tamai M, Kawashiri SY, Sakai W, Yatsuhashi H, Kawakami A, Matsuo H.	Ganglionic acetylcholine receptor autoantibodies in patients with autoimmune diseases including primary biliary cirrhosis.	Mod Rheumatol.	30	42740	2016

Higuchi T, Oka S, Furukawa H, Nakamura M, Komori A, Abiru S, Nagaoka S, Hashimoto S, Naganuma A, Naeshiro N, Yoshizawa K, Shimada M, Nishimura H, Tomizawa M, Kikuchi M, Makita F, Yamashita H, Ario K, Yatsuhashi H, Tohma S, Kawasaki A, Ohira H, Tsuchiya N and Migita K.	Association of a single nucleotide polymorphism upstream of ICOS with Japanese autoimmune hepatitis type 1.	Journal of Human Genetics.	1-4	2016	
Fukui S, Ichinose K, Tsuji S, Ueda M, Nishino A, Nakashima Y, Suzuki T, Horai Y, Koga T, Kawashiri SY, Iwamoto N, Hirai Y, Tamai M, Nakamura H, Sato S, Aramaki T, Iwanaga N, Izumi Y, Origuchi T, Migita K, Ueki Y, Kawakami A.	Hypocholesterolemia predicts relapses in patients with Takayasu arteritis.	Mod Rheumatol.	26(3)	415-420	2016
Takahashi T, Fujimoto N, Yamaguchi A, Hayashi H, Migita K, Ida H, Tanaka T.	<Letter to the Editor> Familial Mediterranean fever with onset in the 70s showing various neutrophilic dermatosis.	J Eur Acad Dermatol Venereol.	30(11)	e129-e131	2016
Izumi M, Sakai T, Shirakawa A, Kozuru H, Jiuchi Y, Izumi Y, Asahara T, Kumagai K, Mawatari M, Osaki M, Motokawa S, Migita K.	Reduced induction of anti-PF4/heparin antibody in RA patients after total knee arthroplasty.	Arthritis Res Ther.	18	191.	2016
Fukui S, Iwamoto N, Shimizu T, Ueda M, Nishino A, Koga T, Kawashiri SY, Ichinose K, Hirai Y, Tamai M, Nakamura H, Aramaki T, Iwanaga N, Izumi Y, Origuchi T, Migita K, Ueki Y, Sato S, Kawakami A.	Fewer subsequent relapses and lower levels of IL-17 in Takayasu arteritis developed after the age of 40 years.	Arthritis Res Ther.	18(1)	293	2016
Koga T, Migita K, Kawakami A.	Biologic therapy in familial Mediterranean fever. (レビュー)	Mod Rheumatol.	26(5)	637-41	2016
Toubiana J, Okada S, Hiller J, Oleastro M, Lagos Gomez M, Aldave Becerra JC, Ouachée-Chardin M, Fouyssac F, Girisha KM, Etzioni A, Van Montfrans J, Camcioğlu Y, Kerns LA, Belohradsky B, Blanche S, Bousfiha A, Rodriguez-Gallego C, Meyts I, Kisand K, Reichenbach J, Renner ED, Rosenzweig S, Grimbacher B, van de Veerdonk FL, Traidl-Hoffmann C, Picard C, Marodi L, Morio T, Kobayashi M, Lilic D, Milner JD, Holland S, Casanova JL, Puel A	Heterozygous STAT1 gain-of-function mutations underlie an unexpectedly broad clinical phenotype: an international survey of 274 patients from 167 kindreds.	Blood	127	3154-3164	2016
Gámez-Díaz L, August D, Stepenksy P, Revel-Vilk S, Seidel MG, Mituiki N, Morio T, Worth AJ, Blessing J, Van de Veerdonk F, Feuchtinger T, Kanariou M, Schmitt-GA Jung S, Seneviratne S, Burns S, Belohradsky BH, Rezaei N, Bakhtiar S, Speckmann C, Jordan M, Grimbacher B.	The extended phenotype of LPS-responsive beige-like anchor protein (LRBA) deficiency.	J. Allergy Clin. Immuno	137	223-23	2016
Hoshino A, Okada S, Yoshida K, Nishida N, Okuno Y, Ueno H, Yamashita M, Okano T, Tsumura M, Nishimura S, Sakata S, Kobayashi M, Nakamura H, Kamizono J, Mitsui-Sekinaka K9, Ichimura T10, Ohga S10, Nakazawa Y11, Takagi M12, Imai K12, Shiraishi Y, Chiba K, Tanaka H, Miyano S, Ogawa S, Kojima S, Nonoyama S, Morio T, Kanegae H	Abnormal hematopoiesis and autoimmunity in human subjects with germline IKZF1 mutations	J Allergy Clin Immunol	in press	2016	
Coulter TI, Chandra A, Bacon CM, Babar J, Curtis J, Sreaton N, Goodlad JR, Farmer G, Steele CL, Leahy TR, Doffinger R, Baxendale H, Bernatoniene J, Edgar JD, Longhurst HJ, Ehl S, Speckmann C, Grimbacher B, Sediva A, Miliota T, Faust SN, Williams AP, Hayman G, Kucuk ZY, Hague R, French P, Brooker R, Forsyth P, Herriot R, Cancrin C, Palma P, Ariganello P, Conlon N, Feighery C, Gavin PJ, Jones A, Imai K, Ibrahim MA, Markelj G, Abinun M, Rieux-Laucat F, Latour S, Peltier I, Fischer A, Touzot F, Casanova JL, Durandy A, Burns SO, Savic S, Kumararatne DS, Moskous D, Kracker S, Vanhaesebroeck B, Okkenhaug K, Picard C, Nejentsev S, Condliffe AM, Cant AJ	Clinical spectrum and features of activated phosphoinositide 3-kinase syndrome: A large patient cohort study.	J Allergy Clin Immunol	in press	2016	

Takagi M, Ogata S, Ueno H, Yoshida K, Yeh T, Hoshino A, Piao J, Yamashita M, Nanya M, Okano T, Kajiwara M, Kanegane H, Muramatsu H, Okuno Y, Shiraishi Y, Chiba K, Tanaka H, Bando Y, Kato M, Hayashi Y, Miyano S, Imai K, Ogawa S, Kojima S, Morio T	Haploinsufficiency of TNFAIP3 (A20) by germline mutation is involved in autoimmune lymphoproliferative syndrome	J Allergy Clin Immunol		in press	2016
Hayakawa S, Okada S, Tsumura M, Sakata S, Ueno Y, Imai K, Morio T, Ohara O, Chayama K, Kobayashi M	A Patient with CTLA-4 Haploinsufficiency Presenting Gastric Cancer	J Clin Immunol	36(1)	28-32	2016
Ono S, Okano T, Hoshino A, Yanagimachi M, Hamamoto K, Nakazawa Y, Imamura T, Onuma M, Niizuma H, Sasahara Y, Tsujimoto H, Wada T, Kunisaki R, Takagi M, Imai K, Morio T, Kanegane H	Hematopoietic Stem Cell Transplantation for XIAP Deficiency in Japan	J Clin Immunol	37(1)	85-91	2016
Yamamoto S, Yamaga T, Sakai Y, Ishida T, Nakasone S, Ohira M, Ota E, Mori R.	Association between physical performance and cardiovascular events in patients with coronary artery disease: protocol for a meta-analysis.	Syst Rev.	5	32	2016
Yamamoto S, Hotta K, Ota E, Matsunaga A, Mori R.	Exercise-based cardiac rehabilitation for people with ventricular assist devices.	Cochrane Database of Systematic Reviews.	6		2016
Shoda T, Ishitsuka K, Kobayashi T, Ota E, Mori R.	TNF blockers for the treatment of Kawasaki disease in children.	The Cochrane Library.	6		2016
Miyazaki C, Moreno RG, Ota E, Swa T, Oladapo OT, Mori R.	Tocolysis for inhibiting preterm birth in extremely preterm birth, multiple gestations and in growth-restricted fetuses: a systematic review and meta-analysis.	Reprod Health.		13	2016
Liao Y, Ota E, Cheng K, Mori R.	Alternative prophylactic therapies (acupuncture and/or moxibustion) for reducing blood loss in the third stage of labour.	Cochrane Database of Systematic Reviews.	6		2016
da Silva Lopes K, Ota E, Tanigaki S, Mori R.	Bed rest with and without hospitalisation in multiple pregnancy for improving outcomes.	Cochrane Database of Systematic Reviews.	1		2016
Balogun OO, da Silva Lopes K, Ota E, Takemoto Y, Rumbold A, Takegata M, Mori R.	Vitamin supplementation for preventing miscarriage	Cochrane Database Syst Re	5		2016
Amiya RM, Miunde LB, Ota E, Swa T, Oladapo OT, Mori R.	Antenatal Corticosteroids for Reducing Adverse Maternal and Child Outcomes in Special Populations of Women at Risk of Imminent Preterm Birth: A Systematic Review and Meta-Analysis.	Plos One.	11(2)		2016
Amari S, Shahrook S, Ota E, Mori R.	Branched-chain amino acid supplementation for improving nutrition in term and preterm neonates	Cochrane Database of Systematic Reviews.	7		2016
Abe Sarah K, Balogun Olukunmi O, Ota E, Takahashi K, Mori R.	Supplementation with multiple micronutrients for breastfeeding women for improving outcomes for the mother and baby	Cochrane Database of Systematic Reviews.	2		2016

雑誌（日本語）

発表者名	論文タイトル名	発表雑誌	巻号	ページ	出版年
金澤伸雄、古川福実	差分解説 中條 - 西村症候群 (Nakajo-Nishimura syndrome)	日本医事新報	4813巻	52	2016
金澤伸雄	中條 - 西村症候群の病因と診断	新薬と臨牀	65巻	1212-1217	2016
金澤伸雄	自己炎症疾患	日本サルコイドーシス / 肉芽腫性疾患学会雑誌	36巻	21-26	2016
右田清志	I 全身性疾患に伴う関節炎 3 惠性腫瘍、血液疾患 4 アミロイドーシス.	リウマチ病学テキスト 改訂第2版.		365-373	2016
右田清志	家族性地中海熱の臨床.	別冊日本臨牀 免疫症候群(第2版)		775-778	2016
岩永 希, 原田康平, 辻 良香, 川原知瑛子, 黒瀧大和, 和泉泰衛, 吉田真一郎, 藤川敬太, 伊藤正博, 川上 純, 右田清志.	TAFRO症候群類似の臨床像を呈した原発性シェーグレン症候群の1例.	日本臨床免疫学会会誌.	39(5)	478-484	2016
山下 舞, 和泉泰衛, 森内由季, 辻 良香, 川原知瑛子, 岩永 希, 野中文陽, 右田清志, 川上 純.	好中球減少に対してガンマグロブリン大量療法が著効した双極性障害に合併した全身性エリテマトーデスの一例.	九州リウマチ.	36(2)	106-111	2016
右田清志	定期的に熱、家族性地中海熱とは?	読売新聞			2016
金兼弘和, 高島健浩, 今井耕輔	原発性免疫不全症における診断のすすめ方	モダンメディア別冊	62 (4)	130-137	2016