

II 研究成果の刊行物リスト

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
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)111	日本臨床社	大阪	2016	544-550
西小森 隆太, 西村 玄, 平家 俊男	Spondyloenchondrodysplasia with immune dysregulation (SPENCDI)(解説/特集)	宮坂伸之	別冊日本臨床免疫症候群(第2版)111	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)(解説/特集)	宮坂伸之	別冊日本臨床免疫症候群(第2版)111	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(解説/特集)	宮坂伸之	別冊日本臨床免疫症候群(第2版)111	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(解説/特集)	宮坂伸之	別冊日本臨床免疫症候群(第2版)111	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(解説・特集)	宮坂伸之	別冊日本臨床免疫症候群(第2版)111	日本臨床社	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)(解説/特集)	宮坂伸之	別冊日本臨床免疫症候群(第2版)111	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)111	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)111	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)111	日本臨床社	大阪	2016	825-830
岡野翼, 今井耕輔, 金兼弘和	xc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)111	日本臨床社	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)111	日本臨床社	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)111	日本臨床社	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)111	日本臨床社	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)111	日本臨床社	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)111	日本臨床社	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
西小森 隆太	自己炎症性疾患	「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	日本	2015	766-770
河合 朋樹 平家俊男	新しく報告された免疫不全症	「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	日本	2015	779 - 785
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	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
河合利尚	原発性免疫不全症候群	尾崎承一	難病事典	学研メディカル秀潤社	東京	2015	464-467
Nobuo Kanazawa	Rare hereditary autoinflammatory disorders.	Yan-Hua Lian	Dermatology Research	NOVA Science Publishers.	NY	2015	pp.3-18
金澤伸雄	中條 西村症候群		別冊日本臨床 新領域症候群シリーズ No.27神経症候群(第2版)	日本臨床社	東京	2014	pp.683-688
金澤伸雄	Blau症候群、サルコイドーシス診療Q&A集	杉山幸比古 監修、山口哲生 四十	厚生労働科学研究費補助金難治性疾患克服研究事業びまん性	鈴木印刷	宇都宮	2014	pp.114-116
原 寿郎	易感染性		小児血液腫瘍学テキスト	診断と治療社	東京	2015	印刷中
原 寿郎	先天性補体欠損症		小児血液腫瘍学テキスト	診断と治療社	東京	2015	印刷中

雑誌等における論文掲載（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)	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	1 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.	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, 黒濱大和, 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
河合 朋樹 平家俊男	自己炎症性疾患	アレルギー・免疫	9月号	1234-81	2015
井澤和司 西小森 隆太 平家俊男	クリオピリン関連周期熱症候群	リウマチ科	54	149-154	2015
古本 雅宏, 岡田 まゆみ, 柴 直子, 丸山 悠	PFAPA症候群100例の臨床像	日本小児科学会雑誌	119巻6号	985-990	2015
樋高 秀憲, 坂田 資尚, 上松 一永, 下田	潰瘍性大腸炎類似の区域性腸炎を伴った家族性地中海熱の	Gastroenterological Endoscopy	57巻4号	1203-1209	2015
右田清志	家族性地中海熱(FMF)	リウマチ科	54(2)	137-142	2015
金澤伸雄	なじみのない蕁麻疹様皮疹	MB Derma	228巻	1-8	2015
金澤伸雄	蕁麻疹の鑑別診断 - 自己炎症症候群	MB Derma	236巻	27-34	2015
金澤伸雄	中條 - 西村症候群	リウマチ科	54巻2号	178-184	2015
大西秀典、金子英雄.	IgGサブクラス欠損症およびIgA欠損症	小児内科47巻増刊号	47	718-721	2015
河合利尚	Blau症候群と若年発症サルコイドーシス	リウマチ科	54(2)	160-5	2015
谷内江昭宏	Autoimmune lymphoproliferative	日本臨床	73巻増刊号 6	53-58	2015

今井耕助	原発性免疫不全症 原因探索法・診断法の進歩	アレルギー・免疫	9月号	1196-1207	2015
今井耕助	iPS細胞を用いた難病研究－臨床病態解明と創薬に向けた	遺伝子医学MOOK	27号	152-157	2015
今井耕助	リンパ球解析 多パラメータ解析による免疫担当細胞亜群同定と機能解析	医学のあゆみ	252巻1号	48-54	2015
西小森隆太、中川権史、栗屋美絵、河合朋	自己炎症性疾患の新展開（総説）	臨床リウマチ	26巻2号	79-87	2014
金澤伸雄	中條-西村症候群	別冊日本臨床 新領域症候群シリーズ		683-688	2014
金澤伸雄	中條-西村症候群	分子リウマチ治療	7巻	25-29	2014
荻野篤彦、金澤伸雄、古江増隆	皮膚を編む 小児掌蹠丘疹性皮膚炎（砂かぶれ様皮膚炎）や自己炎症疾患の臨床と病態	Seminaria Dermatologie No. 227 マルホ皮膚科セミナー「ラジ		4-17	2014
金澤伸雄	中條-西村症候群：和歌山発・プロテアソーム不全による新しい自己炎症疾患	日本臨床皮膚科医学会近畿ブロック会報	29	4-5	2014
金澤伸雄	サルコイドーシス	別冊BIO Clinica	3(2)	80-85	2014
中村悠美、神戸直智	特集「最近のトピックス2014」最近話題の皮膚疾患	臨皮	68(5増)	10-14	2014
高田紗奈美、神戸直智	アレルギー用語解説シリーズ インフラマソーム	アレルギー	63	1142-1143	2014
高田紗奈美、神戸直智	Trend in Allergy インフラマソーム	皮膚アレルギーフロンティア	12	164-165	2014
中野倫代、神戸直智	特集「自己炎症症候群の診断と治療」 若年発症サルコイ	分子リウマチ治療	7	22-24	2014
江原瑞枝、神戸直智	総説 若年発症サルコイドーシス / Blau症候群	呼吸	33	3-9	2014
若林正一郎、神戸直智	内科疾患と皮疹 自己炎症症候群と皮疹	medicina	51	871-875	2014
原 寿郎	原発性免疫不全症～診断と治療の進歩～	MEDICAMENT NEWS	2188号	印刷中	2015
原 寿郎	易感染性	小児内科		印刷中	2015
石村匡崇、高田英俊、原 寿郎	限られたウイルスに易感染性を示す免疫不全症（単純ヘルペス脳炎、EBV、パピローマウイルス、細胞融解型感染形式をとるウイルス）	小児内科	46	1470-4	2014
戸田尚子、原 寿郎	先天性免疫不全症と低栄養	臨床栄養（別冊JCNセレクト9）	Feb	167-72	2014
森尾友宏	自然免疫と発熱	小児内科	46	324-7	2014
高島健浩、森尾友宏	原発性免疫不全症の分子的背景と免疫異常	リウマチ科	51	590-1	2014
中畑龍俊	iPS細胞からHTSに耐えうる疾患モデル評価系の構築	国際医薬品情報	通巻第1026号	25-27	2015
中畑龍俊	特集によせて（iPS細胞を用いた難病研究－臨床病態解明と創薬に向けた研究の最新知見）	遺伝子医学MOOK	27	23-26	2015
大西秀典、加藤善一郎	細胞内寄生菌に脆弱性を示す免疫不全症（MSMDなど）	小児内科	46	1492-7	2014
川口鎮司	膠原病における疾患関連マーカーと肺病変	呼吸器内科	26	220-223	2014
川口鎮司	注目される間質性肺炎の依存症 肺高血圧症	日本医師会雑誌	143	970	2014

川口鎮司	混合性結合組織病	日本内科学会雑誌	103	2501-2506	2014
川口鎮司	膠原病に伴う神経障害	別冊日本臨床	27	572-576	2014
	強皮症	神経症候群（第2版）			
右田清志*，和泉泰衛，地内友香，川原知瑛子，川上 純．	家族性地中海熱．特集：自己炎症症候群の診断と治療．	分子リウマチ治療	7	7-12	2014
右田清志*，野中文陽，清水俊匡，江口勝美．	尿酸によるNLRP3インフラマソームの活性化機構．	Clinical Immunology & Allergology. 臨床免疫・アレルギー科．	62	338-343	2014
右田清志*，藤川敬太，川上 純．	IL-6と家族性地中海熱．	Rheumatic & Autoimmune Diseases	2	30-31	2014
右田清志*，川上 純，江口勝美．	自己炎症疾患の診断と治療．	日本内科学会雑誌	103	2594-2602	2014
金兼弘和	小児免疫不全症の現状と展望 造血不全を合併する原発性免疫不全症	日本小児血液・がん学会雑誌	51	510-514	2014
西田直徳，金兼弘和	ピンポイント小児医療 - 免疫不全症を疑うときの初期検査	小児内科	46	237-243	2014
金兼弘和	ガンマグロブリン補充療法 - 静注製剤と皮下注製剤	小児内科	46	1449-1453	2014
星野顕宏，金兼弘和	自己免疫リンパ増殖症候群	小児科	55	1633-1637	2014
金兼弘和	原発性免疫不全症に合併する自己炎症疾患～炎症性腸疾患をモデルとして～	日本小児科学会雑誌	118	1588-1594	2014
江口郁，野村 裕一，久保田 知洋，山遠剛，井之上 寿美，丸山 慎介，西川 拓朗，和田 昭宏，河野 嘉文，武井 修治．	川崎病の診断基準を満たした若年性特発性関節炎の1例 インターロイキン18値測定の有用性．	小児科臨床	67（7）	1173-1176	2014
武井修治	自然免疫と適応免疫のクロストーク-SLEにおける自然免疫の機能不全．臨床とウイルス	臨床とウイルス	42（3）	89-96	2014

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
Shumpei 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, Arostegui J, Ozen 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, Nagahashi 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 NLR4 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
Nakashimai 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, Sintés J, González-Roca E, Ruiz-Ortiz E, Heike T, Martínez-García JJ, Baroja-Mazo A, Cerutti A, Nishikomori R, Yagüe J, Pelegrín P, Delgado-Beltrán C, Arostegui 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 T, Kunishima S, Nakagawa K, Izawa K, Hiejima E, Kawai T, Yasumi T, Doi H, Katamura 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, Kusuhara K, Nakatsu Y, Tsuzuki T, Mitani K, Hara T.	BTK gene targeting by homologous recombination using a helper-dependent adenovirus/adenovirus-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	Mod Rheumatol.	14	1-5	2016
Ohnishi H, Kawamoto N, Seishima M, Ohara O, Fukao T.	A Japanese family case with juvenile onset Behçet'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 β -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, Kanegane 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, Hirokawa 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.	TheScientificWorldJournal.	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 (gAChR) 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	9E+06	2016
Oka S, Furukawa H, Shimada K, Sugii S, Hashimoto A, Komiya 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, Komiya 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, Shukuwa 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	4E+06	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, Umeda 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, Umeda 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, Camcioglu 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, Lilić D.	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
Gómez-Díaz L, August D, Stepensky 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, Kanegane 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, Screatton 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, Milota T, Faust SN, Williams AP, Hayman G, Kucuk ZY, Hague R, French P, Brooker R, Forsyth P, Herriot R, Cancrini 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, Pellier I, Fischer A, Touzot F, Casanova JL, Durandy A, Burns SO, Savic S, Kumararatne DS, Moshous 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, Mlunde 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
Yokoyama K, Ikeya M, Umeda K, Oda H, Nodomi S, Nasu A, Matsumoto Y, Izawa K, Horigome K, Kusaka T, Tanaka T, Saito MK, Yasumi T, Nishikomori R, Ohara O, Nakayama N, Nakahata T, Heike T, Toguchida J.	Enhanced chondrogenesis of induced pluripotent stem cells from patients with neonatal-onset multisystem inflammatory disease occurs via the caspase 1-independent cAMP/Protein Kinase A/CREB pathway.	Arthritis Rheumatol	67	302-314	2015
Hirano M, Seguchi J, Yamamura M, Narita A, Okanobu H, Nishikomori R, Heike T, Hosokawa M, Morizane Y, Shiraga F.	Successful resolution of stromal keratitis and uveitis using canakinumab in a patient with chronic infantile neurologic, cutaneous, and articular syndrome: a case study.	J Ophthalmic Inflamm Infect.		印刷中	2015
Harada Y, Fukiage K, Nishikomori R, Suzuki S, Futami T.	CINCA syndrome with surgical intervention for valgus deformity and flexion contracture of the knee joint: A case report.	Mod Rheumatol.		印刷中	2015
Kido J, Mizukami T, Ohara O, Takada H, Yanai M.	Idiopathic disseminated bacillus Calmette-Guerin infection in three infants.	Pediatr Int.	57	750-3	2015
Kusuda T, Nakashima Y, Murata K, Kanno S, Nishio H, Saito M, Tanaka T, Yamamura K, Sakai Y, Takada H, Miyamoto T, Mizuno Y, Ouchi K, Waki K, Hara T.	Kawasaki disease-specific molecules in the sera are linked to microbe-associated molecular patterns in the biofilms.	PLoS One.		印刷中	2014
Nozawa T, Mori M, Nishimura K, Sakurai N, Kikuchi M, Hara R, Yokota S.	Usefulness of two IFN- release assays for patients with rheumatic disease.	Pediatr Int.		印刷中	2015
Yokota S, Itoh Y, Morio T, Sumitomo N, Daimaru K, Minota S.	Macrophage Activation Syndrome in Patients with Systemic Juvenile Idiopathic Arthritis under Treatment with Tocilizumab.	J Rheumatol.	42	712-22	2015

Yokota S, Itoh Y, Morio T, Origasa H, Sumitomo N, Tomobe M, Tanaka K, Minota S.	Tocilizumab in systemic juvenile idiopathic arthritis in a real-world clinical setting: results from 1 year of postmarketing surveillance follow-up of 417 patients in Japan.	Ann Rheum Dis.		印刷中	2015
Yokota S, Kikuchi M, Nozawa T, Kanetaka T, Sato T, Yamazaki K, Sakurai N, Hara R, Mori M	Pathogenesis of systemic inflammatory diseases in childhood: "Lessons from clinical trials of anti-cytokine monoclonal antibodies for Kawasaki disease, systemic onset juvenile idiopathic arthritis, and cryopyrin-associated periodic fever syndrome". Yokota S, Kikuchi M, Nozawa T, Kanetaka T, Sato T, Yamazaki K, Sakurai N, Hara R, Mori M	Mod Rheumatol	25	1-10	2015
Ishikura K, Yoshikawa N, Nakazato H, Sasaki S, Nakanishi K, Matsuyama T, Ito S, Hamasaki Y, Yata N, Ando T, Iijima K, Honda M.	Japanese Study Group of Renal Disease in Children. Morbidity in children with frequently relapsing nephrosis: 10-year follow-up of a randomized controlled trial.	Pediatr Nephrol.	30	459-68	2015
Kobayashi I, Mori M, Yamaguchi K, Ito S, Iwata N, Masunaga K, Shimojo N, Ariga T, Okada K, Takei S.	Rheumatology Association of Japan recommendation for vaccination in pediatric rheumatic diseases.	Pediatric Mod Rheumatol.	25	335-43	2015
Takahashi T, Fujisawa T, Kimura M, Ohnishi H, Seishima M.	Familial Mediterranean fever variant with repeated atypical skin eruptions.	J Dermatol	42	903-905	2015
Takahashi T, Fujimoto N, Yamaguchi A, Hayashi H, Migita K, Ida H, Tanaka T.	Familial Mediterranean fever with onset in the 70s showing various neutrophilic dermatosis.	J Eur Acad Dermatol Venereol.		印刷中	2015
Yasunami M, Nakamura H, Agematsu K, Nakamura A, Yazaki M, Kishida D, Yachie A, Toma T, Masumoto J, Ida H, Koga T, Kawakami A, Eguchi K, Furukawa H, Nakamura T, Nakamura M, Migita K.	Identification of disease-promoting HLA class I and protective class II modifiers in Japanese patients with Familial Mediterranean Fever.	PLoS One.	10	e0125938	2015
Yamazaki T, Shigemura T, Kobayashi N, Honda K, Yazaki M, Masumoto J, Migita K, 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		印刷中	2015
Migita K, Izumi Y, Fujikawa K, Agematsu K, Masumoto J, Jiuchi Y, Kozuru H, Nonaka F, Shimizu T, Nakamura T, Iwanaga N, Furukawa H, Yasunami M, Kawakami A, Eguchi K	Dysregulated mature IL-1beta production in familial Mediterranean fever.	Rheumatology		印刷中	2015
Yamagami K, Nakamura T, Nakamura R, Hanioka Y, Seki K, Chiba H, Kobayashi K, Agematsu K.	Familial Mediterranean fever with P369S/R408Q exon3 variant in pyrin presenting as symptoms of PFAPA.	Mod Rheumatol		印刷中	2015
Kawamura S, Agematsu K, Kawamura D, Kawamura G, Suzuki K, Minami M.	A Case Report of Familial Mediterranean Fever Diagnosed Following the Total Knee Arthroplasty.	HSS Journal	11	278-280	2015
Takahara T, Shimizu M, Nakagishi Y, Kinjo N, Yachie A	Serum IL-18 as a potential specific marker for differentiating systemic juvenile idiopathic arthritis from incomplete Kawasaki disease.	Rheumatol Int	35	81-84	2015
Shimizu M, Nakagishi Y, Inoue N, Mizuta M, Ko G, Saikawa Y, Kubota T, Yamasaki Y, Takei S, Yachie A.	Interleukin-18 for predicting the development of macrophage activation syndrome in systemic juvenile idiopathic arthritis.	Clin Immunol.	160	277-81	2015

Kawai T, Arai K, Harayama S, Nakazawa Y, Goto F, Maekawa T, Tamura E, Uchiyama T, Onodera M.	Severe and Rapid Progression in Very Early-Onset Chronic Granulomatous Disease-Associated Colitis.	J Clin Immunol.	35	583-8	2015
Nerome Y, Akaike H, Nonaka Y, Takezaki T, Kubota T, Yamato T, Yamasaki Y, Imanaka H, Kawano Y, Takei S.	The safety and effectiveness of HBV vaccination in patients with juvenile idiopathic arthritis controlled by treatment.	Mod Rheumatol		印刷中	2015
Yamasaki Y, Takei S, Imanaka H, Nerome Y, Kubota T, Nonaka Y, Akaike H, Takezaki T, Kawano Y.	Prediction of long-term remission of oligo/polyarticular juvenile idiopathic arthritis with S100A12 and vasucular endothelial growth factor.	Mod Rheumatol		印刷中	2015
Kubota T, Imanaka H, Takei S, Yamatou T, Nerome Y, Yamasaki Y, Nonaka Y, Akaike H, Takezaki T, Kawano Y.	Disease activity score in 28 joints at 3 months after the initiation of biologic agent can be a predictive target for switching to the second biologic agents in patients with polyarticular juvenile.	Mod Rheumatol		印刷中	2015
Kobayashi I, Mori M, Yamaguchi KI, Ito S, Iwata N, Masunaga K, Shimojo N, Ariga T, Okada K, Takei S.	Pediatric Rheumatology Association of Japan recommendation for vaccination in pediatric rheumatic diseases.	Mod Rheumatol	25	335-343	2015
Umeda M, Aramaki T, Fujikawa K, Iwamoto N, Ichinose K, Terada K, Takeo G, Yonemitsu N, Ueki Y, Migita K, Kawakami A.	Tocilizumab is effective in a familial Mediterranean fever patient complicated with histologically proven recurrent fasciitis and myositis.	Int J Rheum Dis		印刷中	2015
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		印刷中	2015
Nonaka F, Migita K, Jiuchi Y, Shimizu T, Umeda M, Iwamoto N, Izumi Y, Mizokami A, Nakashima M, Ueki Y, Yasunami M, Kawakami A, Eguchi K.	Increased prevalence of MEFV exon 10 variants in Japanese patients with adult-onset Still's disease.	Clin Exp Immunol	179	392-7	2015
Jo T, Horio K, Migita K.	Sweet's syndrome in patients with MDS and MEFV mutations.	N Engl J Med	372	686-8	2015
Migita K, Hisanaga S, Izumi Y, Kawahara C, Shigemitsu Y, Iwanaga N, Araki T, Kamata M, Izumi M, Kumagai K, Kawakami A.	Protracted arthritis in a Japanese patient with familial Mediterranean fever.	Mod Rheumatol	30	1-5	2015
Miyamae T, Tanaka E, Kishi T, Matsuyama T, Igarashi T, Fujikawa S, Taniguchi A, Momohara S, Yamanaka H	Long-term outcome of 114 adult JIA patients in a non-pediatric rheumatology institute in Japan.	Mod Rheumatol	25	62-66	2015
Kaji M, Kishi T, Miyamae T, Nagata S, Yamanaka H, and Fujikawa S.	Efficacy of Adalimumab in a Girl with Refractory Intestinal Behcet's Disease.	Case reports in rheumatology		716138	2015
Yanagi Y, Mizuochi T, Takaki Y, Eda K, Mitsuyama K, Ishimura M, Takada H, Shouval DS, Griffith AE, Snapper SB, Yamashita Y, Yamamoto K.	Novel exonic mutation inducing aberrant splicing in the IL10RA gene and resulting in infantile-onset inflammatory bowel disease: a case report	BMC Gastroenterology		in press	
Shimizu M, Hamaguchi Y, Ishikawa S, Ueno K, Yachie A.	Successful treatment with tocilizumab of a psoriasiform skin lesion induced by etanercept in a patient with juvenile idiopathic arthritis.	Mod Rheumatol	25 (6)	972-973	2015
Fujimaru T, Ito S, Masuda H, Oana S, Kamei K, Ishiguro A, Kato H, Abe J	Decreased levels of inflammatory cytokines in immunoglobulin-resistant Kawasaki disease after plasma exchange.	Cytokine	70	156-60	2014

Marciano BE, Huang CY, Joshi G, Rezaei N, Carvalho BC, Allwood Z, Ilkinciogullari A, Reda SM, Gennery A, Thon V, Espinosa-Rosales F, Al-Herz W, Porras O, Shcherbina A, Szaflarska A, Kiliç S, Franco JL, Gómez Raccio AC, Roxo P Jr, Esteves I, Galal N, Grumach AS, Al-Tamemi S, Yildiran A, Orellana JC, Yamada M, Morio T, Liberatore D, Ohtsuka Y, Lau YL, Nishikomori R, Torres-Lozano C, Mazzucchelli JT, Vilela MM, Tavares FS, Cunha L, Pinto JA, Espinosa-Padilla SE, Hernandez-Nieto L, Elfeky RA, Ariga T, Toshio H, Dogu F, Cipe F, Formankova R, Nuñez-Nuñez ME, Bezrodnik L, Marques JG, Pereira MI, Listello V, Slatter MA, Nademi Z, Kowalczyk D, Fleisher TA, Davies G, Neven B, Rosenzweig SD	BCG vaccination in patients with severe combined immunodeficiency: complications, risks, and vaccination policies.	J Allergy Clin Immunol.	133	1134-41	2014
Shirasaki Y, Yamagishi M, Suzuki N, Izawa K, Nakahara A, Mizuno J, Shoji S, Heike T, Harada Y, Nishikomori R, Ohara O.	Real-time single-cell imaging of protein secretion.	Sci Rep.	4	4736	2014
Oda H, Nakagawa K, Abe J, Awaya T, Funabiki M, Hijikata A, Nishikomori R, Funatsuka M, Ohshima Y, Sugawara Y, Yasumi T, Kato H, Shirai T, Ohara O, Fujita T, Heike T.	Aicardi-Goutières syndrome is caused by IFIH1 mutations.	Am J Hum Genet.	95	121-5	2014
Wada T, Yasumi T, Toma T, Hori M, Maeda S, Umeda K, Heike T, Adachi S, Usami I, Yachie A.	Munc13-4 deficiency with CD5 downregulation on activated CD8+ T cells.	Pediatr Int.	56	605-8	2014
Yokoyama K, Ikeya M, Umeda K, Oda H, Nodomi S, Nasu A, Matsumoto Y, Izawa K, Horigome K, Kusaka T, Tanaka T, Saito MK, Yasumi T, Nishikomori R, Ohara O, Nakayama N, Nakahata T, Heike T, Toguchida J.	Enhanced chondrogenesis of iPS cells from neonatal-onset multisystem inflammatory disease occurs via the caspase-1-independent cAMP/PKA/CREB pathway.	Arthritis Rheumatol.	In press	In press	In press
Nobuo Kanazawa, Kayo Kunimoto, Norihisa Ishii, Yasuji Inamo, Fukumi Furukawa	Is CANDLE the best nomenclature?	Br J Dermatol.	171	659-60	2014
Kanazawa N, Tchernev G, Wollina U	Autoimmunity versus autoinflammation - friend or foe?	Wien Med Wochenschr	164	274-7	2014
Kanazawa N	Hereditary disorders presenting with urticaria.	Immunol Allergy Clin N Am	34	169-179	2014
Ikeda K, Kambe N, Takei S, Nakano T, Inoue Y, Tomiita M, Oyake N, Satoh T, Yamatou T, Kubota T, Okafuji I, Kanazawa N, Nishikomori R, Shimojo N, Matsue H, Nakajima H.	Ultrasonographic assessment reveals detailed distribution of synovial inflammation in Blau syndrome.	Arthritis Res Ther.	16	89	2014
T. Liu, Y. Yamaguchi, Y. Shirasaki, K. Shikada, M. Yamagishi, K. Hoshino, T. Kaisho, K. Takemoto, T. Suzuki, E. Kuranaga, O. Ohara, M. Miura	Real-time single cell analysis provides direct evidence that digital activation of caspase-1 couples macrophage cell death and IL-1 β secretion	Cell Reports	8	974-82	2014
Saito Y, Kagami S, Sanayama Y, Ikeda K, Suto A, Kashiwakuma D, Furuta S, Iwamoto I, Nonaka K, Ohara O, Nakajima H.	AT-rich-interactive domain-containing protein 5A functions as a negative regulator of retinoic acid receptor-related orphan nuclear receptor γ -induced Th17 cell differentiation.	Arthritis Rheumatol.	66	1185-94	2014

Obata Y, Furusawa Y, Endo TA, Sharif J, Takahashi D, Atarashi K, Nakayama M, Onawa S, Fujimura Y, Takahashi M, Ikawa T, Otsubo T, Kawamura YI, Dohi T, Tajima S, Masumoto H, Ohara O, Honda K, Hori S, Ohno H, Koseki H, Hase K.	The epigenetic regulator Uhrf1 facilitates the proliferation and maturation of colonic regulatory T cells	Nature Immunology	15	571-9	2014
Takimoto T, Takada H, Ishimura M, Kirino M, Hata K, Ohara O, Morio T, Hara T	Wiskott-Aldrich syndrome in a girl caused by heterozygous WASP mutation and extremely skewed X-chromosome inactivation: A novel association with maternal uniparental isodisomy 6	Neonatology	In press	In press	2014
Kanegane H, Imai K, Yamada M, Takada H, Ariga T, Bexon M, Rojavin M, Hu W, Kobayashi M, Lawo JP, Nonoyama S, Hara T, Miyawaki T	Efficacy and safety of IgPro20, a subcutaneous immunoglobulin, in Japanese patients with primary immunodeficiency diseases	J Clin Immunol	34	204-11	2014
Yamamura K, Takada H, Uike K, Nakashima Y, Hirata Y, Nagata H, Takimoto T, Ishimura M, Morihana E, Ohga S, Hara T	Early progression of atherosclerosis in children with chronic infantile neurological cutaneous and articular syndrome	Rheumatology (Oxford)	53	1783-7	2014
Koga Y, Takada H, Suminoe A, Ohga S, Hara T	Successful treatment of non-Hodgkin's lymphoma using R-CHOP in a patient with Wiskott-Aldrich syndrome followed by a reduced-intensity stem cell transplant	Pediatr Transplant.	18	E208-11	2014
Kanno S, Nishio H, Tanaka T, Motomura Y, Murata K, Ihara K, Onimaru M, Yamasaki S, Kono H, Sueishi K, Hara T	Activation of an Innate Immune Receptor, Nod1, Accelerates Atherogenesis in Apoe ^{-/-} Mice	J Immunol.	194	773-80	2015
Fukuta M, Nakai Y, Kirino K, Nakagawa M, Sekiguchi K, Nagata S, Matsumoto Y, Yamamoto T, Umeda K, Heike T, Okumura N, Koizumi N, Sato T, Nakahata T, Saito M, Otsuka T, Kinoshita S, Ueno M, Ikeya M, Toguchida J.	Derivation of Mesenchymal Stromal Cells from Pluripotent Stem Cells through a Neural Crest Lineage using Small Molecule Compounds with Defined Media.	Plos one	9	e112291	2014
Yokoyama K, Ikeya M, Umeda K, Oda H, Nodomi S, Nasu A, Matsumoto Y, Izawa K, Horigome K, Kusaka T, Tanaka T, Saito MK, Yasumi T, Nishikomori R, Ohara O, Nakayama N, Nakahata T, Heike T, Toguchida J.	Enhanced chondrogenesis of iPS cells from neonatal-onset multisystem inflammatory disease occurs via the caspase-1-independent cAMP/PKA/CREB pathway.	Arthritis Rheum.	In press	In press	In press
Shimizu M, Hamaguchi Y, Ishikawa S, Ueno K, Yachie A.	Successful treatment with tocilizumab of a psoriasiform skin lesion induced by etanercept in a patient with juvenile idiopathic arthritis.	Modern Rheumatol.	In press	In press	In press
Shimizu M, Nakagishi Y, Yoshida A, Yachie A.	Serum interleukin 18 as a diagnostic remission criterion in systemic juvenile idiopathic arthritis.	J Rheumatol.	41	2328-30	2014
Shimizu M, Ueno K, Ishikawa S, Tokuhisa Y, Inoue N, Yachie A.	Treatment of refractory polyarticular juvenile idiopathic arthritis with tacrolimus.	Rheumatology	53	2120-2	2014
Kovtunovich G, Ghosh MC, Ollivierre W, Weitzel RP, Eckhaus MA, Tisdale JF, Yachie A, Rouault TA.	Wild-type macrophages reverse disease in heme oxygenase 1-deficient mice.	Blood	124	1522-30	2014
Takahara T, Shimizu M, Nakagishi Y, Kinjo N, Yachie A.	Serum IL-18 as a potential specific marker for differentiating systemic juvenile idiopathic arthritis from incomplete Kawasaki disease.	Rheumatology Int.	35	81-4	2015
Kubokawa I, Yachie A, Hayakawa A, Hirase S, Yamamoto N, Mori T, Yanai T, Takeshima Y, Kyo E, Kageyama G, Nagai H, Uehara K, Kojima M, Iijima K.	The first report of adolescent TAFRO syndrome, a unique clinicopathologic variant of multicentric Castleman's disease.	BMC Pediatr.	35	81-4	2015

Migita K, Agematsu K, Yazaki M, Nonaka F, Nakamura A, Toma T, Kishida D, Uehara R, Nakamura Y, Jiuchi Y, Masumoto J, Furukawa H, Ida H, Terai C, Nakashima Y, Kawakami A, Nakamura T, Eguchi K, Yasunami M, Yachie A.	Familial Mediterranean fever: genotype-phenotype correlations in Japanese patients.	Medicine	93	158-64	2014
Nakagishi Y, Shimizu M, Kasai K, Miyoshi M, Yachie A.	Successful therapy of macrophage activation syndrome with dexamethasone palmitate.	Modern Rheumatol.	In press	In press	In press
Takahara T, Shimizu M, Nakagishi Y, Kinjo N, Yachie A.	Serum IL-18 as a potential specific marker for differentiating systemic juvenile idiopathic arthritis from incomplete Kawasaki disease.	Rheumatology Int.	35	81-4	2015
Mitsui-Sekinaka K, Imai K, Sato H, Tomizawa D, Kajiwara M, Nagasawa M, Morio T, Nonoyama S.	Clinical features and hematopoietic stem cell transplantations for CD40 ligand deficiency in Japan.	J Allergy Clin Immunol.	in press		2015
Kato T, Crestani E, Kamae C, Honma K, Yokosuka T, Ikegawa T, Nishida N, Kanegane H, Wada T, Yachie A, Ohara O, Morio T, Notarangelo L.D, Imai K, Nonoyama S.	RAG1 deficiency may present clinically as selective IgA deficiency.	J Clin Immunol.	In press	In press	In press
Oshima K, Imai K, Albert M.H, Bittner T.C, Strauss G, Filipovich A.H, Morio T, Kapoor N, Dalal J, Schultz K.R, Casper J.T, Notarangelo L.D, Ochs H.D, Nonoyama S.	Hematopoietic Stem Cell Transplantation for X-Linked Thrombocytopenia With Mutations in the WAS gene.	J Clin Immunol.	In press	In press	In press
Horiuchi K, Imai K, Mitsui-Sekinaka K, Yeh ZW, Ochs HD, Durandy A, Nonoyama S.	Analysis of somatic hypermutation in the IgM switch region in human B cells.	J Allergy Clin Immunol.	134	411-419	2014
Nakatani K, Imai K, Shigeno M, Sato H, Tezuka M, Okawa T, Mitsui N, Isoda T, Tomizawa D, Takagi M, Nagasawa M, Kajiwara M, Yamamoto M, Arai A, Miura O, Kamae C, Nakagawa N, Honma K, Nonoyama S, Mizutani S, Morio T.	Cord blood transplantation is associated with rapid B cell neogenesis compared with bone marrow transplantation.	Bone Marrow Transplant.	49	1155-61	2014
Al-Herz W, Bousfiha A, Casanova JL, Chatila T, Conley ME, Cunningham-Rundles C, Etzioni A, Franco JL, Boly Gaspar H, Holland SM, Klein C, Nonoyama S, Ochs HD, Oksenhandler E, Picard C, Puck JM, Sullivan KE, Tang ML.	Primary immunodeficiency diseases: an update on the classification from the International Union of Immunological Societies Expert Committee for Primary Immunodeficiency.	Front Immunol.	5	1-33	2014
Tsutsumi N, Kimura T, Arita K, Ariyoshi M, Ohnishi H, Yamamoto T, Zuo X, Maenaka K, Park EY, Kondo N, Shirakawa M, Tochio H, Kato Z.	The structural basis for receptor recognition of human interleukin-18.	Nat Commun.			2014
Kimura T, Tsutsumi N, Arita K, Ariyoshi M, Ohnishi H, Kondo N, Shirakawa M, Kato Z, Tochio H.	Purification, crystallization and preliminary X-ray crystallographic analysis of human IL-18 and its extracellular complexes.	Acta Crystallogr F Struct Biol Commun.	70	1351-6	2014
Funato M, Uemura O, Ushijima K, Ohnishi H, Orii K, Kato Z, Yamakawa S, Nagai T, Ohara O, Kaneko H, Kondo N.	A complement factor B mutation in a large kindred with atypical hemolytic uremic syndrome.	J Clin Immunol.	34	691-5	2014
Kubota K, Ohnishi H, Teramoto T, Norio Kawamoto, Kasahara K, Ohara O, Kondo N.	Clinical and genetic characterization of Japanese sporadic cases of periodic fever, aphthous stomatitis, pharyngitis and adenitis syndrome from a single Medical Center in Japan.	J Clin Immunol.	34	584-93	2014

Mizuno Y, Kato G, Shu E, Ohnishi H, Fukao T, Ohara O, Fukumoto H, Katano H, Seishima M.	Merkel Cell Polyomavirus-positive Merkel Cell Carcinoma in a Patient with Epidermodysplasia Verruciformis.	Acta Derm Venereol.	95	98-9	2015
Sugiura T, Kawaguchi Y, Goto K, Hayashi Y, Gono T, Furuya T, Nishino I, Yamanaka H	Positive association between <i>C8orf13-BLK</i> polymorphisms and polymyositis/dermatomyositis in the Japanese population	PLoS One	9	e90019	2014
Ichida H, Kawaguchi Y, Sugiura T, Takagi K, Katsumata Y, Gono T, Ota Y, Kataoka S, Kawasumi H, Yamanaka H	Clinical Manifestations of Adult-Onset Still's Disease Presenting with Erosive Arthritis: Association with Low Levels of Ferritin and IL-18	Arthritis Care Res	66	642-646	2014
Gono T, Kaneko H, Kawaguchi Y, Hanaoka M, Kataoka S, Kuwana M, Takagi K, Katsumata Y, Ota Y, Kawasumi H, Yamanaka H	Cytokine profiles in polymyositis and dermatomyositis complicated with rapidly progressive or chronic interstitial lung disease	Rheumatology	53	2196-2203	2014
Nonaka F, Migita K*, Jiuchi Y, Shimizu T, Umeda M, Iwamoto N, Fujikawa K, Izumi Y, Mizokami A, Nakashima M, Ueki Y, Yasunami M, Kawakami A, Eguchi K	Increased prevalence of MEFV exon 10 variants in Japanese patients with adult onset Still's disease.	Clin Exp Immunol.	179	392-7	2015
Sugiyama R, Agematsu K, Migita K, Nakayama J, Mokuda S, Ogura F, Haraikawa K, Okumura C, Suehiro S, Morikawa S, Ito Y, Masumoto J	Defect of suppression of inflammasome-independent interleukin-8 secretion from SW982 synovial sarcoma cells by familial Mediterranean fever-derived pyrin mutations.	Mol Biol Rep	41	545-53	2014
Fujikawa K, Migita K*, Shigemitsu Y, Umeda M, Nonaka F, Tamai M, Nakamura H, Mizokami A, Tsukada T, Origuchi T, Yonemitsu N, Yasunami M, Kawakami A, Eguchi K.	MEFV gene polymorphisms and TNFRSF1A mutation in patients with inflammatory myopathy with abundant macrophages.	Clin Exp Immunol	178	224-8	2014
Migita K*, Izumi Y, Fujikawa K, Agematsu K, Masumoto J, Jiuchi Y, Kozuru H, Nonaka F, Shimizu T, Nakamura T, Iwanaga N, Furukawa H, Yasunami M, Kawakami A, Eguchi K.	Dysregulated mature IL-1 production in familial Mediterranean fever.	Rheumatology (Oxford)	[Epub ahead of print]		2014
Migita K*, Izumi Y, Jiuchi Y, Kozuru H1, Kawahara C, Nakamura M, Nakamura T, Agematsu K, Masumoto J, Yasunami M, Kawakami A, Eguchi K.	Serum amyloid A induces NLRP-3-mediated IL-1 secretion in neutrophils.	PLoS One	9	e96703	2014
Yamazaki T*, Shigemura T, Kobayashi N, Honda K, Yazaki M, Masumoto J, Migita K, 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	22	1-3	2014
Jo T*, Horio K, Migita K.	Sweet Syndrome in Myelodysplastic Syndrome Patients with MEFV Gene Mutations.	New Engl J Med.	327	7	2015
Migita K*, Abiru S, Sasaki O, Miyashita T, Izumi Y, Nishino A, Jiuchi Y, Kawakami A, Yasunami M.	Coexistence of familial Mediterranean fever and rheumatoid arthritis.	Mod Rheumatol	24	212-6	2014
Nonaka F, Migita K*, Haramura T, Sumiyoshi R, Kawakami A, Eguchi K.	Colchicine-responsive protracted gouty arthritis with systemic inflammatory reactions.	Mod Rheumatol	24	540-3	2014
Nonaka F, Migita K*, Iwasaki K, Shimizu T, Kawakami A, Yasunami M, Eguchi K.	Overlap Syndrome between Familial Mediterranean Fever and Tumor Necrosis Factor Receptor-Associated Periodic Syndrome in a Lupus Patient.	Tohoku J Exp Med	233	73-77	2014
Komatsu S*, Honma M, Igawa S, Tsuji H, Ishida-Yamamoto A, Migita K, Ida H, Iizuka H.	Cutaneous necrotizing vasculitis as a manifestation of familial Mediterranean fever.	J Dermatol	41	827-9	2014
Izumi Y*, Miyashita T, Migita K.	Safety of tacrolimus treatment during pregnancy and lactation in systemic lupus erythematosus: a report of two patients.	Tohoku J Exp Med	234	51-6	2014

Yokota K*, Fukuda M, Migita K, Tanaka E, Okamoto T, Kimura K.	Three patients with familial Mediterranean fever: a possible underdiagnosed entity in Japan.	Intern Med	53	2013-6	2014
Fujikawa K*, Migita K, Nagasato A, Tsukada T, Kawakami A, Eguchi K.	Mediterranean fever (MEFV) variant P369S/R408Q in a patient with entero-Beçet's disease who successfully responded to treatment with colchicine.	Intern Med	53	2381-4	2014
Nakamura T*, Migita K, Ando Y, Takaoka H, Suzushima H, Shiraishi N.	Amyloid A amyloidosis in a Japanese patient with familial Mediterranean fever associated with homozygosity for the pyrin variant M694I/M694L.	Mod Rheumatol	24	349-52	2014
Fujikawa K*, Migita K, Tsukada T, Kawakami A, Eguchi K.	Protracted febrile myalgia syndrome in a Japanese patient with fasciitis detected on MRI.	Intern Med	53	2817-9	2014
Izumi Y, Takeshita H, Moriwaki Y, Hisatomi K, Masakazu M, Yamashita N, Kawahara C, Shigemitsu Y, Iwanaga N, Kawakami A, Kurohama H, Niino D, Ito M, Migita K*.	Multicentric Castleman disease mimicking IgG4-related disease: a case report.	Mod Rheumatol	22	1-4	2014
Hoshino A, Okuno Y, Migita M, Ban H, Yang X, Kiyokawa N, Adachi Y, Kojima S, Ohara O, Kanegane H.	X-Linked Agammaglobulinemia Associated with B-Precursor Acute Lymphoblastic Leukemia	J Clin Immunol	35	108-111	2015
Mitsuiki N, Yang X, Bartol SJ, Grosserichter-Wagener C, Kosaka Y, Takada H, Imai K, Kanegane H, Mizutani S, van der Burg M, van Zelm MC, Ohara O, Morio T.	Mutations in Bruton's tyrosine kinase impair IgA responses.	Int J Hematol	101	305-313	2015
Mori N, Makino T, Mizawa M, Kagoyama K, Kanegane H, Sakaguchi H, Miyazono T, Kojima S, Shimizu T.	Late presentation of X-linked dyskeratosis congenita with a missense mutation in codon 350 of the dyskerin protein.	Eur J Dermatol	25	75-76	2015
Hoshino A, Nomura K, Hamashima T, Isobe T, Seki M, Hiwatari M, Yoshida K, Shiraishi Y, Chiba K, Tanaka H, Miyano S, Ogawa S, Takita J, Kanegane H.	Aggressive transformation of anaplastic large cell lymphoma with increased number of ALK-translocated chromosomes.	Int J Hematol	101	198-202	2015
Hoshino A, Nomura K, Noguchi K, Kanegane H.	Relapsed leukemia without peripheral blood abnormalities and clinical symptoms detected on MRI.	Pediatr Int	56	798	2014
Aguilar C, Lenoir C, Lambert N, Bègue B, Brousse N, Canioni D, Berrebi D, Roy M, Gérard S, Chapel H, Schwerd T, Siproudhis L, Schàppi M, Al-Ahmari A, Mori M, Yamaide A, Galicier L, Neven B, Routes J, Uhlig HH, Koletzko S, Patel S, Kanegane H, Picard C, Fischer A, Bensussan NC, Ruemmele F, Hugot JP, Latour S	Characterization of Crohn disease in X-linked inhibitor of apoptosis-deficient male patients and female symptomatic carriers	J Allergy Clin Immunol	134	1131-1141.e9	2014
Igarashi A, Kanegane H, Kobayashi M, Miyawaki T, Tsutani K	Cost-Minimization Analysis of IgPro20, a Subcutaneous Immunoglobulin, in Japanese Patients With Primary Immunodeficiency	Clin Ther	36	1616-1624	2014
Koura U, Sakaki-Nakatsubo H, Otsubo K, Nomura K, Oshima K, Ohara O, Wada T, Yachie A, Imai K, Morio T, Miyawaki T, Kanegane H.	Successful treatment of systemic cytomegalovirus infection in severe combined immunodeficiency using allogeneic bone marrow transplantation followed by adoptive immunotherapy.	J Investig Allergol Clin Immunol	24	200-202	2014

Yabal M, Müller N, Adler H, Knies N, Groß CJ, Damgaard RB, Kanegane H, Ringelhan M, Kaufmann T, Heikenwälder M, Strasser A, Groß O, Ruland J, Peschel C, Gyrd-Hansen M, Jost PJ	XIAP Restricts TNF- and RIP3-Dependent Cell Death and Inflammasome Activation	Cell Rep	7	1796-1808	2014
Yamagami K, Miyashita T, Nakamura T, Shirano M, Nakamura T, Kameda K, Nishijima M, Imanishi M, Yang X, Kanegane H	Campylobacter fetus bacteremia with purulent pleurisy in a young adult with primary hypogammaglobulinemia.	Intern Med	53	1221-1225	2014
van Zelm MC, Bartol SJ, Driessen GJ, Mascart F, Reisli I, Franco JL, Wolska-Kusnierz B, Kanegane H, Boon L, van Dongen JJ, van der Burg M	Human CD19 and CD40L deficiencies impair antibody selection and differentially affect somatic hypermutation	J Allergy Clin Immunol	134	135-144.e7	2014
Hoshino A, Imai K, Ohshima Y, Yasutomi M, Kasai M, Terai M, Ishigaki K, Morio T, Miyawaki T, Kanegane H	Pneumothorax in patients with severe combined immunodeficiency	Pediatr Int	56	510-514	2014
Yasumura J, Wago M, Okada S, Nishikomori R, Takei S, Kobayashi M.	A 2-year-old Japanese girl with TNF receptor-associated periodic syndrome: A case report of the youngest diagnosed proband in Japan.	Mod Rheumatol	Epub ahead of print.		2014
Davi S, Minoia F, Pistorio A, Horne A, Consolaro A, Rosina S, Bovis F, Cimaz R, Gamir ML, Ilowite NT, Kone-Paut I, Feitosa de Oliveira SK, McCurdy D, Silva CA, Sztajn bok F, Tsitsami E, Unsal E, Weiss JE, Wulfraat N, Abinun M, Aggarwal A, Apaz MT, Astigarraga I, Corona F, Cuttica R, D'Angelo G, Eisenstein EM, Hashad S, Lepore L, Mulaosmanovic V, Nielsen S, Prahalad S, Rigante D, Stanevicha V, Sterba G, Susic G, Takei S, Trauzeddel R, Zletni M, Ruperto N, Martini A, Cron RQ, Ravelli A	Performance of current guidelines for diagnosis of macrophage activation syndrome complicating systemic juvenile idiopathic arthritis.	Arthritis Rheumatol	66 (10)	2871-2880	2014
Kobayashi I, Mori M, Yamaguchi K, Ito S, Iwata N, Masunaga K, Shimajo N, Ariga T, Okada K, Takei S.	Pediatric Rheumatology Association of Japan recommendation for vaccination in pediatric rheumatic diseases.	Mod Rheumatol	Epub ahead of print.		2014

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)	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	新薬と臨床	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	1 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.	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.	別冊日本臨床 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, 黒濱大和, 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
河合 朋樹 平家俊男	自己炎症性疾患	アレルギー・免疫	9月号	1234-81	2015
井澤和司 西小森 隆太 平家俊男	クリオピリン関連周期熱症候群	リウマチ科	54	149-154	2015
古本 雅宏. 岡田 まゆみ. 柴 直子. 丸山 悠太. 重村 倫成. 小林 法元. 小池 健一. 伯耆原 祥. 神田 仁. 本多 貴実子. 梅田 陽. 上松 一永	PFAPA症候群100例の臨床像	日本小児科学会雑誌	119巻6号	985-990	2015
樋高 秀憲, 坂田 資尚, 上松 一永, 下田 良, 藤本 一眞, 岩切 龍一	潰瘍性大腸炎類似の区域性腸炎を伴った家族性地中海熱の1例	Gastroenterological Endoscopy	57巻4号	1203-1209	2015
右田清志	家族性地中海熱(FMF)	リウマチ科	54(2)	137-142	2015
金澤伸雄	なじみのない蕁麻疹様皮疹	MB Derma	228巻	1-8	2015
金澤伸雄	蕁麻疹の鑑別診断 - 自己炎症症候群	MB Derma	236巻	27-34	2015

金澤伸雄	中條 - 西村症候群	リウマチ科	54巻2号	178-184	2015
大西秀典、金子英雄	IgGサブクラス欠損症およびIgA欠損症	小児内科47巻増刊号	47	718-721	2015
河合利尚	Blau症候群と若年発症サルコイドーシス	リウマチ科	54(2)	160-5	2015
谷内江昭宏	Autoimmune lymphoproliferative	日本臨床	73巻増刊号 6	53-58	2015
今井耕助	原発性免疫不全症 原因探索法・診断法の進歩	アレルギー・免疫	9月号	1196-1207	2015
今井耕助	ips細胞を用いた難病研究—臨床病態解明と創薬に向けた	遺伝子医学MOOK	27号	152-157	2015
今井耕助	リンパ球解析 多パラメータ解析による免疫担当細胞亜群同定と機能解析	医学のあゆみ	252巻1号	48-54	2015
西小森隆太、中川権史、粟屋美絵、河合朋	自己炎症性疾患の新展開（総説）	臨床リウマチ	26巻2号	79-87	2014
金澤伸雄	中條-西村症候群	別冊日本臨床 新領域症候群シリーズ		683-688	2014
金澤伸雄	中條-西村症候群	分子リウマチ治療	7巻	25-29	2014
荻野篤彦、金澤伸雄、古江増隆	皮膚を編む 小児掌蹠丘疹性皮膚炎（砂かぶれ様皮膚炎）や自己炎症疾患の臨床と病態	Seminaria Dermatologie No. 227 マルホ皮膚科セミナー「ラジ		4-17	2014
金澤伸雄	中條 - 西村症候群：和歌山発・プロテアソーム不全による新しい自己炎症疾患	日本臨床皮膚科医会近畿ブロック会報	29	4-5	2014
金澤伸雄	サルコイドーシス	別冊BIO Clinica	3(2)	80-85	2014
中村悠美、神戸直智	特集「最近のトピックス2014」最近話題の皮膚疾患	臨皮	68(5増)	10-14	2014
高田紗奈美、神戸直智	アレルギー用語解説シリーズ インフラマソーム	アレルギー	63	1142-1143	2014
高田紗奈美、神戸直智	Trend in Allergy インフラマソーム	皮膚アレルギーフロンティア	12	164-165	2014
中野倫代、神戸直智	特集「自己炎症症候群の診断と治療」 若年発症サルコイ	分子リウマチ治療	7	22-24	2014
江原瑞枝、神戸直智	総説 若年発症サルコイドーシス / Blau症候群	呼吸	33	3-9	2014
若林正一郎、神戸直智	内科疾患と皮疹 自己炎症症候群と皮疹	medicina	51	871-875	2014
原 寿郎	原発性免疫不全症～診断と治療の進歩～	MEDICAMENT NEWS	2188号	印刷中	2015
原 寿郎	易感染性	小児内科		印刷中	2015
石村匡崇、高田英俊、原 寿郎	限られたウイルスに易感染性を示す免疫不全症（単純ヘルペス脳炎、EBV、パピローマウイルス、細胞融解型感染形式をとるウイルス）	小児内科	46	1470-4	2014
戸田尚子、原 寿郎	先天性免疫不全症と低栄養	臨床栄養（別冊JCNセレクト9）	Feb	167-72	2014
森尾友宏	自然免疫と発熱	小児内科	46	324-7	2014
高島健浩、森尾友宏	原発性免疫不全症の分子的背景と免疫異常	リウマチ科	51	590-1	2014

中畑龍俊	iPS細胞からHTSに耐えうる疾患モデル評価系の構築	国際医薬品情報	通巻第1026号	25-27	2015
中畑龍俊	特集によせて(iPS細胞を用いた難病研究-臨床病態解明と創薬に向けた研究の最新知見)	遺伝子医学MOOK	27	23-26	2015
大西秀典、加藤善一郎	細胞内寄生菌に脆弱性を示す免疫不全症(MSMDなど)	小児内科	46	1492-7	2014
川口鎮司	膠原病における疾患関連マーカーと肺病変	呼吸器内科	26	220-223	2014
川口鎮司	注目される間質性肺炎の依存症 肺高血圧症	日本医師会雑誌	143	970	2014
川口鎮司	混合性結合組織病	日本内科学会雑誌	103	2501-2506	2014
川口鎮司	膠原病に伴う神経障害	別冊日本臨床	27	572-576	2014
	強皮症	神経症候群(第2版)			
右田清志*, 和泉泰衛, 地内友香, 川原知瑛子, 川上 純	家族性地中海熱. 特集: 自己炎症症候群の診断と治療.	分子リウマチ治療	7	7-12	2014
右田清志*, 野中文陽, 清水俊匡, 江口勝美	尿酸によるNLRP3インフラマソームの活性化機構.	Clinical Immunology & Allergology. 臨床免疫・アレルギー科.	62	338-343	2014
右田清志*, 藤川敬太, 川上 純	IL-6と家族性地中海熱.	Rheumatic & Autoimmune Diseases	2	30-31	2014
右田清志*, 川上 純, 江口勝美	自己炎症疾患の診断と治療.	日本内科学会雑誌	103	2594-2602	2014
金兼弘和	小児免疫不全症の現状と展望 造血不全を合併する原発性免疫不全症	日本小児血液・がん学会雑誌	51	510-514	2014
西田直徳, 金兼弘和	ピンポイント小児医療-免疫不全症を疑うときの初期検査	小児内科	46	237-243	2014
金兼弘和	ガンマグロブリン補充療法- 静注製剤と皮下注製剤	小児内科	46	1449-1453	2014
星野顕宏, 金兼弘和	自己免疫リンパ増殖症候群	小児科	55	1633-1637	2014
金兼弘和	原発性免疫不全症に合併する自己炎症疾患~炎症性腸疾患をモデルとして~	日本小児科学会雑誌	118	1588-1594	2014
江口郁, 野村 裕一, 久保田 知洋, 山遠剛, 井之上 寿美, 丸山 慎介, 西川 拓朗, 和田 昭宏, 河野 嘉文, 武井 修治	川崎病の診断基準を満たした若年性特発性関節炎の1例 インターロイキン18値測定の有用性.	小児科臨床	67(7)	1173-1176	2014
武井修治	自然免疫と適応免疫のクロストーク-SLEにおける自然免疫の機能不全. 臨床とウイルス	臨床とウイルス	42(3)	89-96	2014