

雑誌（英文）

発表者氏名	論文タイトル名	発表誌名	巻号	ページ	出版年
Kido T, Hosaka S, Imagawa K, Fukushima H, Morio T, Nonoyama S, Takada H.	Initial manifestations in Patients with Inborn Errors of Immunity Based on Onset Age: a Study from a Nationwide Survey in Japan.	J Clin Immunol.		doi: 10.1007/s10875-023-01434-6.	2023
Inoue K, Miyamoto S, Tomomasa D, Adachi E, Azumi S, Horikoshi Y, Ishihara T, Osone S, Kawahara Y, Kudo K, Kato Z, Ohnishi H, Kashimada K, Imai K, Ohara O, van Zelm MC, Cowan MJ, Morio T, Kanegane H.	Clinical and Genetic Characterization of Patients with Artemis Deficiency in Japan.	J Clin Immunol.	43(3)	585-594	2022
Leiding JW, Vogel TP, Santarlas VGJ, Mhaskar R, Smith MR, Carisey A, Vargas-Hernandez A, Silva-Carmona M, Heeg M, Rensing-Ehl A, Neven B, Hadjadj J, Hambleton S, Leahy TR, Van Hagen M, Cunningham-Rundles C, Dutmer CM, Sharapova SO, Taskinen M, Chua I, Hague R, Kleemann C, Kostyuchenko L, Morio T, Thatayatikom A, Ozen A, Scherbina A, Bauer CS, Flanagan SE, Gambineri E, Giovannini-Chami L, Heimall J, Sullivan KE, Allenspach E, Romberg N, Deane SG, Prince BT, Rose M, Bohnsack J, Mousalem T, Jesudas R, Dos Santos Vilela MM, O'Sullivan M, Schmid JP, Průhová Š, Rees M, Su H, Bahna S, Baris S, Bartnikas LM, Chang Berger A, Briggs TA, Brothers S, Bundy V, Chan AY, Chandrasekaran S, Christiansen M, Cole T, Cook MC, Desai MM, Fischer U, Fulcher DA, Gallo S, Gauthier A, Gennery AR, Marques JG, Gottrand F, Grimbacher B, Grunebaum E, Haapaniemi E, Hämäläinen S, Heiskanen K, Heiskanen-Kosma T, Hoffman HM, Gonzalez-Granado LI, Guerreiro AL, Kainulainen L, Kumar A, Lawrence MG, Levin C, Martelius T, Neth O, Olbrich P, Palma A, Patel NC, Pozos T, Preece K, Lugo Reyes SO, Russell MA, Schejter Y, Seroogy C, Sinclair J, Skevofilax E, Suan D, Suegeorgz D, Szaboles P, Velasco H, Warnatz K, Walkovich K, Worth A; STAT3 GOF	Monogenic early-onset lymphoproliferation and autoimmunity: Natural history of STAT3 gain-of-function syndrome.	J Allergy Clin Immunol.	11	S0091-6749(22)01182-4	2022
Bousfiha A, Moundir A, Tangye SG, Picard C, Jeddane L, Al-Herz W, Rundles CC, Franco JL, Holland SM, Klein C, Morio T, Oksenhendler E, Puel A, Puck J, Seppänen MRJ, Somech R, Su HC, Sullivan KE, Torgerson TR, Meyts I.	The 2022 Update of IUIS Phenotypical Classification for Human Inborn Errors of Immunity.	J Clin Immunol.	42(7)	1508-1520	2022
Eto S, Nukui Y, Tsumura M, Nakagama Y, Kashimada K, Mizoguchi Y, Utsumi T, Taniguchi M, Sakura F, Noma K, Yoshida Y, Ohshima S, Nagashima S, Okamoto K, Endo A, Imai K, Kanegane H, Ohnishi H, Hirata S, Sugiyama E, Shime N, Ito M, Ohge H, Kido Y, Bastard P, Casanova JL, Tanaka J, Morio T, Okada S.	Neutralizing Type I Interferon Autoantibodies in Japanese Patients with Severe COVID-19.	J Clin Immunol.	42(7)	1360-1370	2022
Tangye SG, Al-Herz W, Bousfiha A, Cunningham-Rundles C, Franco JL, Holland SM, Klein C, Morio T, Oksenhendler E, Picard C, Puel A, Puck J, Seppänen MRJ, Somech R, Su HC, Sullivan KE, Torgerson TR, Meyts I.	Human Inborn Errors of Immunity: 2022 Update on the Classification from the International Union of Immunological Societies Expert Committee.	J Clin Immunol.	42(7)	1473-1507	2022

Manry J, Bastard P, Gervais A, Le Voyer T, Rosain J, Philippot Q, Michailidis E, Hoffmann HH, Eto S, Garcia-Prat M, Bizien L, Parra-Martínez A, Yang R, Haljasmägi L, Migaud M, Särekannu K, Maslovskaja J, de Prost N, Tandjaoui-Lambotte Y, Luyt CE, Amador-Borrero B, Gaudet A, Poissy J, Morel P, Richard P, Cognasse F, Troya J, Trouillet-Assant S, Belot A, Saker K, Garçon P, Rivière JG, Lagier JC, Gentile S, Rosen LB, Shaw E, Morio T, Tanaka J, Dalmau D, Tharaux PL, Sene D, Stepanian A, Mégarbane B, Triantafyllia V, Fekkar A, Heath JR, Franco JL, Anaya JM, Solé-Violán J, Imberti L, Biondi A, Bonfanti P, Castagnoli R, Delmonte OM, Zhang Y, Snow AL, Holland SM, Biggs CM, Moncada-Vélez M, Arias AA, Lorenzo L, Boucherit S, Anglicheau D, Planas AM, Haerynck F, Duvlis S, Ozcelik T, Keles S, Bousfiha AA, El Bakkouri J, Ramirez-Santana C, Paul S, Pan-Hammarström Q, Hammarström L, Dupont A, Kurolap A, Metz CN, Aiuti A, Casari G, Lampasona V, Ciceri F, Barreiros LA, Dominguez-Garrido E, Vidigal M, Zatz M, van de Beek D, Sahanic S, Tancevski I, Stepanovskyy Y, Boyarchuk O, Nukui Y, Tsumura M, Vidaur L, Tangye SG, Burrel S, Duffy D, Quintana-Murci L, Klocperk A, Kann NY, Shcherbina A, Lau YL, Leung D, Coulongeat M, Marlet J, Koning R, Reves LF.	The risk of COVID-19 death is much greater and age dependent with type I IFN autoantibodies.	Proc Natl Acad Sci U S A.	119(21)	e2200413119	2022
Yamashita M, Morio T.	AIOLOS Variants Causing Immunodeficiency in Human and Mice.	Front Immunol.	13	866582	2022
Abdrabou SSMA, Toita N, Ichihara S, Tozawa Y, Takahashi M, Fujiwara S, Ashida T, Ohara O, Ariga T, Manabe A, Konno M, Yamada M.	Absent XIAP expression in T cell blasts and causal XIAP mutations including non-coding deletion.	Pediatr. Int.	64(1)	e14892	2022
Sbihi Z, Tanita K, Bachelet C, Bole C, Jabot-Hanin F, Tores F, Le Loch M, Khodr R, Hoshino A, Lenoir C, Oleastro M, Villa M, Sposrito L, Prieto E, Danielian S, Brunet E, Picard C, Taga T, Abdrabou SSMA, Isoda T, Yamada M, Palma A, Kanegane H, Latour S.	Identification of germline non-coding deletions in XIAP gene causing XIAP deficiency reveals a key promoter sequence.	J Clin Immunol.	42(3)	559-571	2022
Uchiyama T, Takahashi S, Nakabayashi K, Okamura K, Edasawa K, Yamada M, Watanabe N, Mochizuki E, Yasuda T, Miura A, Kato M, Tomizawa D, Otsu M, Ariga T, Onodera M.	Nonconditioned ADA-SCID gene therapy reveals ADA requirement in the hematopoietic system and clonal dominance of vector-marked clones.	Mol Ther Methods Clin.	23	424-433	2021
Miyamoto S, Umeda K, Kurata M, Yanagimachi M, Iguchi A, Sasahara Y, Okada K, Koike T, Tanoshima R, Ishimura M, Yamada M, Sato M, Takahashi Y, Kajiwara M, Kawaguchi H, Inoue M, Hashii Y, Yabe H, Kato K, Atsuta Y, Imai K, Morio T.	Hematopoietic Cell Transplantation for Inborn Errors of Immunity Other than Severe Combined Immunodeficiency in Japan: Retrospective Analysis for 1985-2016.	J Clin Immunol.	42	529-545	2022
Ueki M, Sakamoto K, Nishioka N, Ohata H, Nobuta T, Takezaki S, Manabe A, Yamada M.	Rheumatologic manifestations with elevated levels of IL-6, IL-17A, and IL-23 in a patient with scurvy.	Mod Rheumatol Case Rep.	9	rxac059.	2022
Kobayashi I, Shimomura M, Ueki M, Takezaki S, Okura Y, Nawate M, Yamada M, Takahashi M, Ariga T.	Development of Graves' disease during drug-free remission of juvenile dermatomyositis.	Mod Rheumatol Case Rep.	6	55-58	2022
Tanita K, Kawamura Y, Miura H, Mitsuiki N, Tomoda T, Inoue K, Iguchi A, Yamada M, Yoshida T, Muramatsu H, Tada N, Matsui T, Kato M, Eguchi K, Ishimura M, Ohga S, Imai K, Morio T, Yoshikawa T, Kanegane H.	Case Report: Rotavirus vaccination and severe combined immunodeficiency in Japan.	Front Immunol.	13	786375	
Muramoto Y, Nihira H, Shiokawa M, Izawa K, Hiejima E, Seno H; Japan Pediatric Inflammatory Bowel Disease Working Group	Anti-integrin $\alpha v\beta 6$ antibody as a diagnostic marker for pediatric patients with ulcerative colitis.	Gastroenterol.	163	1094-1097	2022
Nakano T, Sasahara Y, Kikuchi A, Moriya K, Niizuma H, Nihori T, Shirota M, Funayama R, Nakayama K, Aoki Y, Kure S.	Novel POLE mutations identified in patients with IMAGE-I syndrome cause aberrant subcellular localization and protein degradation in the nucleus.	J Med Genet.	59(1)	1116-1122	2022
Sasahara Y, Wada T, Morio T.	Impairment of cytokine production following immunological synapse formation in patients with Wiskott-Aldrich syndrome and leukocyte adhesion deficiency type 1.	Clin Immunol.	242	109098	2022
Nishitani-Isa M, Mukai K, Honda Y, Nihira H, Tanaka T, Shibata H, Kodama K, Hiejima E, Izawa K, Kawasaki Y, Osawa M, Katata Y, Onodera S, Watanabe T, Uchida T, Kure S, Takita J, Ohara O, Saito KM, Nishikomori R, Taguchi T, Sasahara Y, Yasumi T.	Trapping of CDC42 C-terminal variants in the Golgi drives pyrin inflammasome hyperactivation.	J Exp Med.	219(6)	e20211889	2022

Yamamoto M, Sato M, Onishi Y, Sasahara Y, Sano H, Masuko M, Nakamae H, Matsuoka K, Ara T, Washio K, Onizuka M, Watanabe K, Takahashi Y, Hirakawa T, Nishio M, Sakashita C, Kobayashi T, Sawada A, Ichinohe T, Fukuda T, Hashii Y, Atsuta Y, Arai A.	Registry data analysis of hematopoietic stem cell transplantation on systemic chronic active EBV infection patients in Japan.	Am J Hematol,	97(6)	780-790	2022
Sasahara Y, Uchida T, Suzuki T, Abukawa D.	Primary immunodeficiencies associated with inflammatory bowel disease in Southeast and East Asia.	Front Immunol,	12	786538	2022
Miyamoto S, Umeda K, Kurata M, Yanagimachi M, Iguchi A, Sasahara Y, Okada K, Koike T, Tanoshima R, Ishimura M, Yamada M, Sato M, Takahashi Y, Kajiwara M, Kawaguchi H, Inoue M, Hashii Y, Yabe H, Kato K, Atsuta Y, Imai K, Morio T.	Hematopoietic cell transplantation for inborn errors of immunity other than severe combined immunodeficiency in Japan: retrospective analysis for 1985-2016.	J Clin Immunol.	42(3)	529-545	2022
Aoki M, Izawa K, Tanaka T, Honda Y, Shiba T, Maeda Y, Miyamoto T, Okamoto K, Nishitani-Isa M, Nihira H, Imai K, Takita J, Nishikomori R, Hiejima E, Yasumi T.	Case Report: A Pediatric Case of Familial Mediterranean Fever Concurrent With Autoimmune Hepatitis.	Front Immunol.	13	917398	2022
Hojo K, Furuta T, Komaki S, Yoshikane Y, Kikuchi J, Nakamura H, Ide M, Shima S, Hiyoshi Y, Araki J, Tanaka S, Ozono S, Yoshida A, Nobusawa S, Morioka M, Nishikomori R.	Systemic inflammation caused by an intracranial mesenchymal tumor with a EWSR1::CREM fusion presenting associated with IL-6/STAT3 signaling	Neuropathology.	doi: 10.1111/neup.12877.		2022
Kozycki CT, Kodati S, Huryn L, Wang H, Warner BM, Jani P, Hammoud D, Abu-Asab MS, Jittayasothorn Y, Mattapallil MJ, Tsai WL, Ullah E, Zhou P, Tian X, Soldatos A, Moutsopoulos N, Kao-Hsieh M, Heller T, Cowen EW, Lee CR, Toro C, Kalsi S, Khavandgar Z, Baer A, Beach M, Long Priel D, Nehrebecky M, Rosenzweig S, Romeo T, Deutlich N, Brenchley L, Pelayo E, Zein W, Sen N, Yang AH, Farlev G, Sweetser DA	Gain-of-function mutations in ALPK1 cause an NF- κ B-mediated autoinflammatory disease: functional assessment, clinical phenotyping, and disease course of patients with ROSAH syndrome.	Ann Rheum Dis.	81(10)	1453-1464	2022
Matsubayashi T, Yamamoto M, Takayama S, Otsuki Y, Yamadori I, Honda Y, Izawa K, Nishikomori R, Oto T.	Allograft dysfunction after lung transplantation for COPA syndrome: A case report and literature review.	Mod Rheumatol Case Rep.	6(2)	314-8	2022
Matsuyuki K, Ide M, Houjou K, Shima S, Tanak S, Watanabe Y, Tomino H, Egashira T, Takayanagi T, Tashiro K, Okamura K, Suzuki T, Miyamoto T, Shibata H, Yasumi T, Nishikomori R.	Novel AP3B1 mutations in a Hermansky-Pudlak syndrome type2 with neonatal interstitial lung disease.	Pediatr Allergy Immunol.	33(2)	e13748	2022
Miyamoto T, Honda Y, Izawa K, Kanazawa N, Kadokami S, Ohnishi H, Fujimoto M, Kambe N, Kase N, Shiba T, Nakagishi Y, Akizuki S, Murakami K, Bamba M, Nishida Y, Inui A, Fujisawa T, Nishida D, Iwata N, Otsubo Y, Ishimori S, Nishikori M, Tanizawa K, Nakamura T, Ueda T, Ohwada Y, Tsuyusaki Y, Shimizu M, Ebato T, Iwao K, Kubo A, Kawai T, Matsubayashi T, Miyazaki T, Kanayama T, Nishitani-Isa M, Nihira H, Abe J, Tanaka T, Hiejima E, Okada S, Ohara O, Saito MK, Takita J, Nishikomori R, Yasumi T	Assessment of type I interferon signatures in undifferentiated inflammatory diseases: A Japanese multicenter experience.	Front Immunol.	13	905960	2022
Nishitani-Isa M, Mukai K, Honda Y, Nihira H, Tanaka T, Shibata H, Kodama K, Hiejima E, Izawa K, Kawasaki Y, Osawa M, Kataoka Y, Onodera S, Watanabe T, Uchida T, Kure S, Takita J, Ohara O, Saito KM, Nishikomori R, Taguchi T, Sasahara Y, Yasumi T.	Trapping of CDC42 C-terminal variants in the Golgi drives pyrin inflammasome hyperactivation.	J Exp Med.	219(6)	e20211889	2022
Ohto T, Tayeh AA, Nishikomori R, Abe H, Hashimoto K, Baba S, Arias-Loza AP, Soda N, Satoh S, Matsuda M, Iizuka Y, Kondo T, Koseki H, Yan N, Higuchi T, Fujita T, Kato H.	Intracellular virus sensor MDA5 mutation develops autoimmune myocarditis and nephritis.	J Autoimmun.	127	102794	2022
Okada E, Morisada N, Horinouchi T, Fujii H, Tsuji T, Miura M, Katori H, Kitagawa M, Morozumi K, Toriyama T, Nakamura Y, Nishikomori R, Nagai S, Kondo A, Aoto Y, Ishiko S, Rossanti R, Sakakibara N, Nagano C, Yamamura T, Ishimori S, Usui J, Yamagata K, Iijima K, Imasawa T, Nozu K.	Detecting MUC1 Variants in Patients Clinicopathologically Diagnosed With Having Autosomal Dominant Tubulointerstitial Kidney Disease.	Kidney Int Rep.	7(4)	857-66	2022
Ono R, Tsumura M, Shima S, Matsuda Y, Gotoh K, Miyata Y, Yoto Y, Tomomasa D, Utsumi T, Ohnishi H, Kato Z, Ishiwada N, Ishikawa A, Wada T, Uhara H, Nishikomori R, Hasegawa D, Okada S, Kanegane H.	Novel STAT1 Variants in Japanese Patients with Isolated Mendelian Susceptibility to Mycobacterial Diseases.	J Clin Immunol.	43(2)	466-478	2023
Tanaka T, Shiba T, Honda Y, Izawa K, Yasumi T, Saito KM, Nishikomori R.	Induced Pluripotent Stem Cell-Derived Monocytes/Macrophages in Autoinflammatory Diseases.	Front Immunol.	13	870535	2022
Toyofuku E, Takeshita K, Ohnishi H, Kiriodshi Y, Masuoka H, Kadokami T, Nishikomori R, Nishimura K, Kobayashi C, Ebato T, Shigemura T, Inoue Y, Suda W, Hattori M, Morio T, Honda K, Kanegane H.	Dysregulation of the Intestinal Microbiome in Patients With Haploinsufficiency of A20.	Front Cell Infect Microbiol.	11	787667	2021

Yamada Y, Inui K, Okano T, Mandai K, Nishikomori R, Nakamura H, Tsuruta D.	Ultrasound and biopsy findings in arthritis with familial Mediterranean fever.	J Med Ultrason (2001). 49(1)	115-6	2022
Matsukawa Y, Isshiki K, Osumi T, Fujiyama S, Fukushima H, Uchiyama T, Yamada M, Deguchi T, Imadome KI, Matsumoto K, Tomizawa D, Takada H, Onodera M, Kato M.	Successful hematopoietic stem cell transplantation with reduced dose of busulfan for Omenn syndrome.	Blood Cell Ther. 5(3)	75-78	2022
Yamashita M, Eguchi S, Tomomasa D, Kamiya T, Niizato D, Mitsuiki N, Isoda T, Funakoshi H, Mizuno Y, Okamoto K, Nguyen TM, Takada H, Takagi M, Imai K, Morio T, Kanegane H.	Case report: HLA-haploidentical hematopoietic cell transplant with posttransplant cyclophosphamide in a patient with leukocyte adhesion deficiency type I.	Front Immunol. 13	1020362	2022
Morita A, Imagawa K, Asayama K, Terakado T, Takahashi S, Yaita K, Tagawa M, Matsubara D, Takada H.	Immunological characteristics of severe acute hepatitis of unknown origin in a child post SARS-CoV-2 infection.	Clin Immunol. 245	109138	2022
Hasegawa M, Fukushima H, Suzuki R, Yamaki Y, Hosaka S, Inaba M, Nakao T, Kobayashi C, Yoshimi A, Tsuchida M, Koike K, Fukushima T, Takada H.	Effect of Germline MEFV Polymorphisms on the Prognosis of Japanese Children with Cancer:	A Regional Analysis. Oncology. 100(7)	376-383	2022
Morita A, Hosaka S, Imagawa K, Ishiodori T, Nozaki Y, Murakami T, Takada H.	Time course of peripheral immunophenotypes of multisystem inflammatory syndrome in children.	Clin Immunol. 236	108955	2022
Usami M, Ikawa Y, Sakai Y, Fujiki T, Wada T.	Refractory gastroduodenal ulcers: A rare complication with Bloom syndrome.	Clin Case Rep. 10(9)	10.1002/ccr 3.6141	2022
Miyazawa H, Wada T.	Immune-mediated inflammatory diseases with chronic excess of serum interleukin-18.	Front Immunol. 13	930141	2022
Banday AZ, Kaur A, Akagi T, Bhattacharai D, Muraoka M, Dev D, Das J, Sachdeva MUS, Karmakar I, Arora K, Kaur G, Pandiarajan V, Jindal AK, Wada T, Koeffler HP, Suri D, Ahluwalia J, Kanegane H, Bhatia P, Rawat A, Singh S.	A Novel CEBPE Variant Causes Severe Infections and Profound Neutropenia.	J Clin Immunol. 42(7)	1434-1450	2022
Tozaki N, Tawada C, Niwa H, Mizutani Y, Shu E, Kawase A, Miwa Y, Ohnishi H, Sasai H, Miyako K, Hosokawa J, Kato A, Kobayashi K, Miyazaki T, Shirakami Y, Shimizu M, Iwata H.	A case of VEXAS syndrome (vacuoles, E1 enzyme, X-linked, autoinflammatory, somatic) with decreased oxidative stress levels after oral prednisone and tocilizumab treatment.	Front Med (Lausanne). 9	1046820	2022
Niwano T, Hosoya T, Kadokami S	An adult case of suspected A20 haploinsufficiency mimicking polyarteritis nodosa.	Rheumatology (Oxford). 61(11)	e337-e340	2022
Tobai H, Endo M, Ishimura M, Moriya K, Yano J, Kanamori K, Sato N, Amanuma F, Maruyama H, Muramatsu H, Shibahara J, Narita M, Fumoto S, Peltier D, Ohga S.	Neonatal intestinal obstruction in Hoyeraal-Hreidarsson syndrome with novel RTEL1 variants.	Pediatr Blood Cancer. 13	e30250	2023
Yamamori A, Hamada M, Muramatsu H, Wakamatsu M, Hama A, Narita A, Tsumura Y, Yoshida T, Doi T, Terada K, Higa T, Yamamoto N, Miura H, Shiota M, Watanabe K, Yoshida N, Maemura R, Imaya M, Miwata S, Narita K, Kataoka S, Taniguchi R, Suzuki K, Kawashima N, Nishio N, Iwafuchi H, Ito M, Kojima S, Okuno Y, Takahashi Y.	Germline and somatic RUNX1 variants in a pediatric bone marrow failure cohort.	Am J Hematol. doi: 10.1002/ajh.26874		2023
Maemura R, Wakamatsu M, Matsumoto K, Sakaguchi H, Yoshida N, Hama A, Yoshida T, Miwata S, Kitazawa H, Narita K, Kataoka S, Ichikawa D, Hamada M, Taniguchi R, Suzuki K, Kawashima N, Nishikawa E, Narita A, Okuno Y, Nishio N, Kato K, Kojima S, Morita K, Muramatsu H, Takahashi Y.	Clinical Impact of Melphalan Pharmacokinetics on Transplantation Outcomes in Children Undergoing Hematopoietic Stem Cell Transplantation.	Cell Transplant. 31	963689722114	2022
Wakamatsu M, Kojima D, Muramatsu H, Okuno Y, Kataoka S, Nakamura F, Sakai Y, Tsuge I, Ito T, Ueda K, Saito A, Morihana E, Ito Y, Ohashi N, Tanaka M, Tanaka T, Kojima S, Nakajima Y, Ito T, Takahashi Y.	TREC/KREC Newborn Screening followed by Next-Generation Sequencing for Severe Combined Immunodeficiency in Japan.	J Clin Immunol. 42(8)	1696-1707.	2022
Kobayashi A, Ohtaka R, Toki T, Hara J, Muramatsu H, Kanezaki R, Takahashi Y, Sato T, Kamio T, Kudo K, Sasaki S, Yoshida T, Utsugisawa T, Kanno H, Yoshida K, Nannya Y, Takahashi Y, Kojima S, Miyano S, Ogawa S, Terui K, Ito E.	Dyserythropoietic anaemia with an intronic GATA1 splicing mutation in patients suspected to have Diamond-Blackfan anaemia.	EJHaem. 3(1)	163-167.	2022
Miyagishima M, Hamada M, Hirayama Y, Muramatsu H, Tainaka T, Shirota C, Hinoki A, Imaizumi T, Nakatoh M, Kamei M, Nishikawa E, Kawashima N, Narita A, Nishio N, Kojima S, Takahashi Y.	Risk factors for unplanned removal of central venous catheters in hospitalized children with hematological and oncological disorders.	Int J Hematol. 116(2)	288-294	2022
Fujikawa H, Shimizu H, Nambu R, Takeuchi I, Matsui T, Sakamoto K, Gocho Y, Miyamoto T, Yasumi T, Yoshioka T, Arai K.	Monogenic inflammatory bowel disease with STXBP2 mutations is not resolved by hematopoietic stem cell transplantation but can be alleviated via immunosuppressive drug therapy.	Clin Immunol. 246	109203	2022

Tomomasa D, Hiejima E, Miyamoto T, Tanita K, Matsuoka M, Niizato D, Mitsuiki N, Isoda T, Yasumi T, van Zelm MC, Morio T, Kanegane H.	Recurrent tandem duplication of UNC13D in familial hemophagocytic lymphohistiocytosis type 3.	Clin Immunol.	242	109104	2022
Yamamoto S, Nakao S, Inoue H, Koga Y, Kojima-Ishii K, Semba Y, Maeda T, Akashi K, Ohga S.	A preterm newborn-onset juvenile myelomonocytic leukemia-like myeloproliferation with PTPN11 mutation.	Pediatr Blood Cancer.	70(2)	e29915	2023
Kimoto Y, Horiuchi T	The complement system and ANCA associated vasculitis in the era of anti-complement drugs.	Front. Immunol.	13	926044	2022
Kashiwado Y, Kimoto Y, Sawabe T, Irino K, Nakano S, Hiura J, Wang Q, Kawano S, Ayano M, Mitoma H, Ono N, Arinobu Y, Niiro H, Hotta T, Kang D, Akashi K, Ohshima S, Takeuchi T, Horiuchi T.	Antibody response to SARS-CoV-2 mRNA vaccines in patients with rheumatic diseases in Japan: Interim analysis of a multicenter cohort study	Mod Rheumatol.	33(2)	367-372	2022
Oku K, Kimoto Y, Horiuchi T, Yamamoto M, Kondo Y, Okamoto M, Atsumi T, Takeuchi T.	Risk factors for hospitalization or mortality for COVID-19 in patients with rheumatic diseases: Results of a nationwide JCR COVID-19 registry in Japan.	Mod Rheumatol.		roac104.	2022
Sugimori Y, Iwasaki Y, Takeshima Y, Okubo M, Kobayashi S, Hatano H, Yamada S, Nakano M, Yoshida R, Ota M, Tsuchida Y, Nagafuchi Y, Shimane K, Yoshida K, Kurosaka D, Sumitomo S, Shoda H, Yamamoto K, Okamura T, Fujio K.	Transcriptome Profiling of Immune Cell Types in Peripheral Blood Reveals Common and Specific Pathways Involved in the Pathogenesis of Myositis-Specific Antibody-Positive Inflammatory Myopathies	ACR Open Rheumatol.	5(2)	93-102	2023
Nakano M, Ota M, Takeshima Y, Iwasaki Y, Hatano H, Nagafuchi Y, Itamiya T, Maeda J, Yoshida R, Yamada S, Nishiwaki A, Takahashi H, Takahashi H, Akutsu Y, Kusuda T, Suetsugu H, Liu L, Kim K, Yin X, Bang SY, Cui Y, Lee HS, Shoda H, Zhang X, Bae SC, Terao C, Yamamoto K, Okamura T, Ishigaki K, Fujio K.	Distinct transcriptome architectures underlying lupus establishment and exacerbation.	Cell.	185(18)	3375-3389.e21	2022

雑誌（和文）

発表者氏名	論文タイトル名	発表誌名	巻号	ページ	出版年
山田雅文	易感染性 小児内科 小児疾患診療のための病態生理 2		53	668-673	2021
山田雅文	学術コーナー 新たに導入された新生児マススクリーニングの話題	原発性免疫不全症 北海道小児科医会	38	5-6	2022
仁平寛士, 井澤和司, 八角高裕, 西小森隆太.	【血管炎の診療update-診断・治療の新展開-】血管炎症候群の症候と診断 アデノシンデアミナーゼ2(ADA2)欠損症.	日本臨床	80(8)	1260-4	2022
仁平寛士, 井澤和司, 八角高裕, 西小森隆太.	【自己炎症性疾患の最前線】アデノシンデアミナーゼ2(ADA2)欠損症.	臨床免疫・アレルギー科	77(6)	705-10	2022
西小森隆太.	小児免疫関連異常症の診療.	久留米医学会雑誌	85(6-8)	117-25	2022
西小森隆太, 田中征治, 井手水紀, 井澤和司.	【自己炎症性疾患の治療最前線】自己炎症性疾患治療における生物学的製剤. 臨床免疫・アレルギー科. 2022; 78(5): 552-9.	臨床免疫・アレルギー科	78(5)	552-9	2022
西小森隆太, 田中征治, 井手水紀, 北城恵史郎.	【発熱と血液疾患】不明熱と自己炎症性疾患.	血液内科	85(5)	682-90	2022
西小森隆太, 田中征治, 八角高裕.	【サイトカインストームと小児疾患】サイトカインストームが関与する疾患 自己炎症性疾患に伴うサイトカインストーム.	小児科診療	85(4)	467-72	2022
穂坂翔, 高田英俊	補体と疾患 一補体受容体欠損症 日本臨床	日本臨床社	11(80)	1809-1812	2022
宮澤 英恵, 和田 泰三.	【先天性免疫異常症の進歩】血球貪食をきたす先天性免疫異常症(解説).	臨床免疫・アレルギー科	78(5)	504-511	2022
堀内孝彦	血管性浮腫 一アレルギー性と遺伝性 病態の差異と新しい治療法について	infoAllergy (日本アレルギー協会機関誌)	99	4	2022
柏戸佑介, 堀内孝彦	リウマチ科専攻医のための補体知識	リウマチ科	67(4)	482-490	2022
堀内孝彦	巻頭言：企画にあたって 新時代を迎えた補体学	日本臨牀	80(11)	1702-1705	2022

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
笹原洋二	免疫不全を伴う特徴的症候群	日本小児血液・がん学会 編集	小児血液・腫瘍 学テキスト 改訂 第2版	診断と治療社	東京	2022	422-426
高田英俊	リンパ球の発生と機能		ネルソン 小児科学	エルゼビア・ジャパン		印刷中	
高田英俊	先天性免疫不全症	日本血液学会編集	血液専門 医テキスト 第4 版 日本 血液学会 編集	南江堂	東京	印刷中	
高田英俊	易感染性	日本小児血液・がん学会 編集	小児血 液・腫瘍 学 (改定 第2版)	診断と治療社	東京	2022	37-39
高田英俊	予防接種	日本小児血液・がん学会 編集	小児血 液・腫瘍 学 (改定 第2版)	診断と治療社	東京	2022	263-265
高田英俊	自然免疫不全症・自己炎症性疾患	日本小児血液・がん学会 編集	小児血 液・腫瘍 学 (改定 第2版)	診断と治療社	東京	2022	435-438
大西秀典	自己炎症性症候群.	公益財団法人 日本リウマチ財団 教育研修委員会 一般社団法人 日本リママチ学会 生涯教育委員会 編	リウマチ 病学テキスト 改定 第3版	南江堂	東京	2022	428-432