

研究成果の刊行に関する一覧表

<書籍>

著者氏名	論文タイトル名	書籍全体の編集者名	書籍名	出版社名	出版地	出版年	ページ
村田満	1 特発性血小板減少性紫斑病 [指定難病63]	水澤英洋, 五十嵐隆, 北川泰久, 高橋和久, 弓倉整	指定難病ペディア	医学書出版	東京	2019	
羽藤高明	血小板の同種抗原	面川進	日本輸血・細胞治療学会認定医制度指定カリキュラム	日本輸血・細胞治療学会	東京	2019	91-93
羽藤高明	輸血・血液型検査	矢富裕、他	今日の臨床検査2019-2020	南江堂	東京	2019	110-117
羽藤高明	ヒト白血球抗原 (HLA) 検査	矢富裕、他	今日の臨床検査2019-2020	南江堂	東京	2019	118-123
羽藤高明	先天性および後天性血管障害による出血	中尾眞二、松村到、神田善伸	血液疾患 最新の治療 2020-2022	南江堂	東京	2019	231-234
立忝良崇、加藤規利、丸山彰一	aHUS		プロフェッショナル腎臓内科学	中外医学社		In press	
森下英理子	遺伝性血栓性素因	中尾眞二、松村到、神田善伸	血液疾患最新の治療2020-2022	南江堂	東京	2019	258-261
森下英理子	先天性血栓傾向	南学正臣	改訂第9版 内科学書—血液造血管器疾患、神経疾患	中山書店	東京	2019	245-247
森下英理子	遺伝性血栓性素因は静脈血栓塞栓症 (VTE) リスクにどの程度注意したらよいか? 欠乏症の種類によってリスクは異なるのか?	日本静脈学会	新臨床静脈学会	メジカルビュー社	東京	2019	429-431
小林隆夫	肺血栓塞栓症の予防と治療指針	岡元和文	救急・集中治療最新ガイドライン2020-'21	総合医学社	東京	2020	335-340

小林隆夫	産科・婦人科領域のDIC	丸山征郎	ファーマナビゲーターDIC編改訂版	メディカルレビュー社	東京	2019	156-165
小林隆夫	産褥期の静脈血栓塞栓症	猿田享男, 北村惣一郎	専門家による私の治療 2019-20年度版	日本医事新報社	東京	2019	1369-1371
小林隆夫	血液疾患合併妊娠	永井良三, 綾部琢哉, 大須賀穰	産婦人科研修ノート	診断と治療社	東京	2019	456-459
富山佳昭	特発性血小板減少性紫斑病	門脇孝, 小室一成, 宮地良樹監修	日常診療に活かす診療ガイドライン UP-TO-DATE 2018-2019	メディカルレビュー社	大阪	2018	pp465-469
富山佳昭	妊娠時の ITP 管理	朝倉英策編	臨床に直結する血栓止血学	中外医学社	東京	2018	pp163-167
羽藤高明	血液製剤の適正使用 血小板	前田平生, 大戸斉, 岡崎仁	輸血学 (改訂第4版)	中外医学社	東京	2018	857-865
羽藤高明	輸血・血液型検査	櫻林郁之介	今日の臨床検査 2019-2020	南江堂	東京	2019	110-117
羽藤高明	HLA 検査	櫻林郁之介	今日の臨床検査 2019-2020	南江堂	東京	2019	118-123
丸山慶子, 小亀浩市	PS Tokushima (K196E) 変異の検査	朝倉英策	臨床に直結する血栓止血学 改訂2版	中外医学社	東京	2018	85-87
川端千晶・池田洋一郎	非典型溶血性尿毒症症候群の遺伝子診断		血液内科	科学評論社	東京	2019	234-238
池田洋一郎	非典型溶血性尿毒症症候群 (aHUS)		日本血栓止血学会誌	日本血栓止血学会	東京	2019	164-167
吉田瑤子, 南学正臣	非典型溶血性尿毒症症候群 (aHUS) の病態と臨床検査		炎症と免疫	先端医学社	東京	2019	17-41
吉田瑤子, 南学正臣	フォンヴィレブランド因子		腎と透析	東京医学社	東京	2019	226-228
吉田瑤子, 南学正臣	補体関連非典型溶血性尿毒症症候群		腎と透析	東京医学社	東京	2019	603-608

長屋聡美、 森下英理子	ホモシステイン	「腎と透析」 編集委員会	『腎と透析 ベッドサイド 検査事典』 (編集)	東京医学 社	東京	2018	108- 110
森下英理子	血栓性素因の検 査の進め方	朝倉英策	『臨床に直結す る血栓止血学 改訂2版』	中外医学 社	東京	2018	30-35
森下英理子	出血性素因の検 査の進め方	朝倉英策	『臨床に直結す る血栓止血学 改訂2版』	中外医学 社	東京	2018	36-41
森下英理子	アンチトロンビ ン, プロテイン S, プロテイン C.	朝倉英策	『臨床に直結す る血栓止血学 改訂2版』	中外医学 社	東京	2018	71-76
森下英理子	DOAC の AT・ PC・ PS 測定への影 響	朝倉英策	『臨床に直結す る血栓止血学 改訂2版』	中外医学 社	東京	2018	77-81
森下英理子	先天性 AT・ PC・PS 欠乏症 の遺伝子検査	朝倉英策	『臨床に直結す る血栓止血学 改訂2版』	中外医学 社	東京	2018	82-84
森下英理子	Lp(a), ホモシ ステイン	朝倉英策	『臨床に直結す る血栓止血学 改訂2版』	中外医学 社	東京	2018	151- 153
森下英理子	稀な先天性凝固 因子欠乏/異常 症	朝倉英策	『臨床に直結す る血栓止血学 改訂2版』	中外医学 社	東京	2018	185- 192
長屋聡美、 森下英理子	先天性第 X 因子 欠乏症の臨床と 遺伝子検査	朝倉英策	『臨床に直結す る血栓止血学 改訂2版』	中外医学 社	東京	2018	193- 195
森下英理子	アミロイドーシ ス	朝倉英策	『臨床に直結す る血栓止血学 改訂2版』	中外医学 社	東京	2018	272- 276
森下英理子	先天性アンチト ロンビン (AT)・プロテイン C(PC)・プロ テインS(PS) 欠乏症	朝倉英策	『臨床に直結す る血栓止血学 改訂2版』	中外医学 社	東京	2018	424- 431
小嶋哲人	プロテインC、 プロテインS	『腎と透 析』編集委 員会	腎と透析ベッ ドサイド検査 事典 腎と透析 2018年84巻 増刊号	東京医学 社	東京	2018	231- 233

小林隆夫	産科領域の DIC	朝倉栄策	臨床に直結する血栓止血学改訂 2 版	中外医学社	東京	2018	387-393
Satoh T and Kuwana M	T cell abnormalities	Ishida Y and Tomoyama Y	Autoimmune Thrombocytopenia	Springer	London, UK	2017	63-74
Satoh T and Kuwana M	Differential diagnosis: secondary ITP	Ishida Y and Tomoyama Y	Autoimmune Thrombocytopenia	Springer	London, UK	2017	97-106
Hato T, Kurata Y	Epidemiology.	Ishida Y, Tomiyama Y	Autoimmune Thrombocytopenia	Springer	Singapore	2017	41-49
Hato T	Transfusion	Ishida Y, Tomiyama Y	Autoimmune Thrombocytopenia	Springer	Singapore	2017	191-197
羽藤高明	輸血・血液型検査	櫻林郁之介	今日の臨床検査 2017-2018	南江堂	東京	2017	129-136
羽藤高明	HLA 検査	櫻林郁之介	今日の臨床検査 2017-2018	南江堂	東京	2017	137-142
Fujimura Y, Kokame K, Matsumoto M	Upshaw-Schulman syndrome	Nima Rezaei	Genetic Syndromes: A Comprehensive Reference Guide	Springer	In press	In press	In press
香美祥二	溶血性尿毒症症候群と非典型溶血性尿毒症症候群	矢崎義雄	内科学第 11 版	朝倉書店	東京	2017	1449-1451
小林隆夫	肺血栓塞栓症の予防と治療指針	岡元和文	救急・集中治療最新ガイドライン 2018-' 19	総合医学社	東京	2018	327-331
小林隆夫	産褥期の静脈血栓塞栓症	猿田享男, 北村惣一郎	私の治療 2017- 2018 年度版	日本医事新報社	東京	2017	1541-1543
小林隆夫	産科 DIC	日本産婦人科・新生児血液学会	産婦人科・新生児領域の血液疾患診療の手引き	メディカルレビュー社	東京	2017	97-108
小林隆夫	血栓塞栓症合併妊娠	日本産婦人科・新生児血液学会	産婦人科・新生児領域の血液疾患診療の手引き	メディカルレビュー社	東京	2017	41-52

<雑誌>

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Oka S, Wakui M, Fujimori Y, Kuroda Y, Nakamura S, Kondo Y, Nakagawa T, Katagiri H, Murata M	Application of clot-fibrinolysis waveform analysis to assessment of in vitro effects of direct oral anticoagulants on fibrinolysis.	Int J Lab Hematol.	In press		2020
Wakui M, Kawai K, Mizushima T, Nishime C, Serizawa A, Suemizu H, Asakura K, Yamauchi Y, Hayashida T, Suematsu M, Murata M	Fatty Acid β -Oxidation-dependent and -independent Responses and Tumor Aggressiveness Acquired Under Mild Hypoxia.	Anticancer Res.	39(1)	191-200	2019
Wakui M, Fujimori Y, Katagiri H, Nakamura S, Kondo Y, Kuroda Y, Nakagawa T, Shimizu N, Murata M	Assessment of in vitro effects of direct thrombin inhibitors and activated factor X inhibitors through clot waveform analysis.	J Clin Pathol	72(3)	244-250	2019
Uwamino Y, Kubota H, Sasaki T, Kosaka A, Furuhashi M, Uno S, Kudoh J, Murata M, Hasegawa N	Recovery of FRI-5 carbapenemase at a Japanese hospital where FRI-4 carbapenemase was discovered.	J Antimicrob Chemother.	74(11)	3390-3392	2019
Nakagawa T, Wakui M, Hayashida T, Nishime C, Murata M	Intensive optimization and evaluation of global DNA methylation quantification using LC-MS/MS.	Anal Bioanal Chem.	411(27)	7221-7231	2019
Uwamino Y, Nishimura T, Sato Y, Tamizu E, Uno S, Mori M, Fujiwara H, Kawabe H, Murata M, Hasegawa N	Showering is associated with Mycobacterium avium complex lung disease: An observational study in Japanese women	J Infect Chemother	26(3)	211-214	2019
Uwamino Y, Sakai A, Nishimura T, Noguchi M, Uno S, Fujiwara H, Mori M, Wakui M, Murata M, Hasegawa N.	Effect of refrigeration of blood samples in lithium-heparin tubes on QuantiFERON TB Gold Plus test result.	J Infect Chemother.	25(3)	312-314	2019

Uwamino Y, Nishimura T, Sato Y, Tamizu E, Asakura T, Uno S, Mori M, Fujiwara H, Ishii M, Kawabe H, Murata M, Hasegawa N	Low serum estradiol levels are related to Mycobacterium avium complex lung disease: a cross-sectional study.	BMC Infect Dis.	19(1)	1055	2019
Wakui M, Fujimori Y, Nakamura S, Kondo Y, Kuroda Y, Oka S, Nakagawa T, Katagiri H, Murata M	Distinct features of bivalent direct thrombin inhibitors, hirudin and bivalirudin, revealed by clot waveform analysis and enzyme kinetics in coagulation assays.	J Clin Pathol.	72(12)	817-824	2019
Mori T, Kikuchi T, Kato J, Koda Y, Sakurai M, Kikumi O, Inose R, Murata M, Hasegawa N, Nakayama H, Yamazaki R, Okamoto S	Seasonal changes in indoor airborne fungal concentration in a hematology ward.	J Infect Chemother.	26(4)	363-366	2019
藤森祐多, 涌井昌俊, 村田満	【血栓塞栓症の防止対策-抗凝固療法の最近の動向】 Key words 静脈血栓症と臨床検査	カレントセラピー	37 巻 3 号	298	2019
柏木浩和、桑名正隆、羽藤高明、高蓋寿朗、藤村欣吾、倉田義之、村田満、富山佳昭	成人特発性血小板減少性紫斑病治療の参照ガイド 2019 改訂版	臨床血液	60(8)	877-896	2019
Kashiwagi H, Kuwana M, Hato T, Takafuta T, Fujimura K, Kurata Y, Murata M, Tomiyama Y	Reference guide for management of adult immune thrombocytopenia in Japan: 2019 Revision.	Int J Hematol.	111(3)	329-351	2020
Akuta K, Kashiwagi H, Yujiri T, Nishiura N, Morikawa Y, Kato H, Honda S, Kanakura Y, Tomiyama Y.	A unique phenotype of acquired Glanzmann thrombasthenia due to non-function-blocking anti- α IIb β 3 autoantibodies.	J Thromb Haemost.	17(1)	206-219	2019
Hato T, Shimada N, Kurata Y, Kuwana M, Fujimura K, Kashiwagi H, Takafuta T, Murata M, Tomiyama Y	Risk factors for skin, mucosal, and organ bleeding in adults with primary ITP: a nationwide study in Japan	Blood Advances	In press		2020

Ikeda Y, Yamanouchi J, Hato T, Yasukawa M, Takenaka K	Safe childbirth for a type 1 antithrombin-deficient woman with novel mutation in the SERPINC1 gene undergoing antithrombin concentrate therapy.	Blood Coagulation and Fibrinolysis	30(1)	47-51	2019
Joko K, Hato T	Acute liver failure and intractable gastric ulcer in plasma prekallikrein deficiency	松山赤十字病院医学雑誌	44(1)	21-27	2019
Fujimura Y, Lämmle B, Tanabe S, Sakai K, Kimura T, Kokame K, Miyata T, Takahashi Y, Taniguchi S, Matsumoto M	Patent ductus arteriosus generates neonatal hemolytic jaundice with thrombocytopenia in Upshaw-Schulman syndrome.	Blood Adv	3(21)	3191-3195	2019
van Dorland HA, Mansouri Taleghani M, Sakai K, Friedman KD, George JN, Hrachovinova I, Knöbl PN, von Krogh AS, Schneppenheim R, Aebi-Huber I, Bütikofer L, Largiadèr CR, Cermakova Z, Kokame K, Miyata T, Yagi H, Terrell DR, Vesely SK, Matsumoto M, Lämmle B, Fujimura Y, Kremer Hovinga JA.	The International Hereditary Thrombotic Thrombocytopenic Purpura Registry: Key findings at enrolment until 2017.	Haematologica.	104(10)	2107-2115	2019
Sakai K, Wada H, Nakatsuka Y, Kubo M, Hayakawa M, Matsumoto M.	Characteristics behaviors of coagulation and fibrinolysis markers in acquired thrombotic thrombocytopenic purpura.	J Intensive Care Med	In press		

Sakai K, Kuwana M, Tanaka H, Hosomichi K, Hasegawa A, Uyama H, Nishio K, Omae T, Hishizawa M, Matsui M, Iwato K, Okamoto A, Okuhiro K, Yamashita Y, Itoh M, Kumekawa H, Takezako N, Kawano N, Matsukawa T, Sano H, Oshiro K, Hayashi K, Ueda Y, Mushino T, Ogawa Y, Yamada Y, Murata M, Matsumoto M.	HLA loci predisposing to immune TTP in Japanese: potential role of the shared ADAMTS13 peptide bound to different HLA-DR.	Blood	In press		
Kato H, et al	Safety and effectiveness of eculizumab for adult patients with atypical hemolytic uremic syndrome in Japan: interim analysis of post-marketing surveillance.	Clin Exp Nephrol	23	65-75	2019
Ito S, et al.	Safety and effectiveness of eculizumab for pediatric patients with atypical hemolytic-uremic syndrome in Japan: interim analysis of post-marketing surveillance.	Clin Exp Nephrol	23	112-121	2019
宮川義隆	免疫性血小板減少症と血栓性血小板減少性紫斑病に対するリツキシマブ	臨床血液	60	480-487	2019
Nakajima-Doi S, Seguchi O, Shintani Y, Fujita T, Fukushima S, Matsumoto Y, Eura Y, Kokame K, Miyata S, Matsuda S, Mochizuki H, Iwasaki K, Kimura Y, Toda K, Kumai Y, Kuroda K, Watanabe T, Yanase M, Kobayashi J, Fukushima N	Experience of the use of octreotide for refractory gastrointestinal bleeding in a patient with Jarvik2000 left ventricular assist device	J Artif Organs	22 (4)	334-337	2019

Sugawara Y, Kato H, Yoshida Y, Fujisawa M, Kokame K, Miyata T, Akioka Y, Miura K, Hattori M, Nangaku M	Novel CFHR2-CFHR1 hybrid in C3 glomerulopathy identified by genomic structural variation analysis	Kidney Int Rep	4 (12)	1759-1762	2019
Usui M, Ozawa T, Kim Y, Mashiko T, Matsuzono K, Maruyama K, Kokame K, Usui R, Koide R, Fujimoto S	Cerebral venous sinus thrombosis associated with protein S deficiency during pregnancy: a case report	J Obstet Gynaecol	40 (1)	135-136	2020
Miyoshi T, Maruyama K, Oku H, Asahara S, Hanada H, Neki R, Yoshimatsu J, Kokame K, Miyata T	Predictive value of protein S-specific activity and ELISA testing in patients with the protein S K196E mutation	Thromb Res	185	1-4	2020
Akuta K, Kiyomizu K, Kashiwagi H, Kunishima S, Nishiura N, Banno F, Kokame K, Kato H, Kanakura Y, Miyata T, Tomiyama Y	Knock-in mice bearing constitutively active α IIb(R990W) mutation develop macrothrombocytopenia with severe platelet dysfunction	J Thromb Haemost	18 (2)	497-509	2020
秋山正志, 小亀浩市	ADAMTS13の構造変化と機能発現	Thromb Med	9 (3)	189-198	2019
宮田敏行, 小亀浩市	TMAの遺伝子診断：TTPとaHUS	日本血栓止血誌	31 (1)	17-27	2020
Esumi S, Morishita E, Yasuda M, Nakajima K, Imashuku S.	Portal cavernoma cholangiopathy due to extrahepatic portal vein and supra-mesenteric vein thromboses associated with congenital protein C deficiency in a young adult	Arch Clin Med Case Rep	3(6)	436-441	2019
Nomoto H, Takami A, Espinoza JL, Onizuka M, Kashiwase K, Morishima Y, Fukuda T, Koderu Y, Doki N, Miyamura k, Mori T, akao S, Morishita E.	Recipient ADAMTS13 single-nucleotide polymorphism predicts relapse after unrelated bone marrow transplantation of hematologic malignancy.	Int J Mol Sci	20(1)	214	2019
Setaka T, Hirano K, Moriya K, Morita S, Shinakai T, Morishita E, Ichida T.	Portal vein thrombosis in a patient with hereditary antithrombin deficiency.	Intern Med.	58(12)	1733-1737	2019

Yamada S, Arahita M, Morishita E, Ichinose A, Asakura H.	The first reported case of acquired haemophilia A in which bleeding episodes were successfully treated via administration of a single-dose mixture of activated factor VIIa/X.	Haemophilia	25(5)	e350-e352	2019
Ueda k, Morishita E, Shiraki H, Matsuoka S, Imashuku S.	Aortic mural thrombus associated with congenital protein C deficiency in an elderly patient.	J Atheroscler Thromb	27(1)	100-103	2020
Yamashita M, Komaki T, Tashiro K, Inada Y, Iwata A, Morishita E, Miura SI.	Hereditary antithrombin deficiency presenting with deep venous thrombosis during the second pregnancy: A case report.	Intern Med.	59(2)	235-239	2020
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Horio T, Morishita E, Mizuno S, Uchino K, Hanamura I, Espinoza JL, Morishima Y, Kodera Y, Onizuka M, Kashiwase K, Fukuda T, Doki N, Miyamura K, Mori T, Nakao S, Takami A.	Donor heme oxygenase-1 promoter gene polymorphism predicts survival after unrelated bone marrow transplantation for high-risk patients.	Cancers (Basel)	12(2)	424	2020
Watanabe S, Matsumoto S, Nakahara I, Ishii A, Hatano T, Mori M, Morishita E, Nagata I.	A case of ischemic stroke with congenital protein C deficiency and carotid web successfully treated by anticoagulant and carotid stenting.	Front Neurol	18	10.3389/fneur.2020.00099.	2020

Yamada S, Okumura H, Morishita E, Asakura H.	Complete hemostasis achieved by factor XIII concentrate administration in a patient with bleeding after teeth extraction as a complication of aplastic anemia and chronic disseminated intravascular coagulation.	Blood Coagul Fibrinolysis	9	doi: c.000000 00000009 02.	2020
Togashi T, Meguro-Horike M, Nagaya S, Sugihara S, Ichinohe T, Araiso Y, Yamaguchi K, Mori K, Imai Y, Kuzasa K, Horike SI, Asakura H, Watanabe A, Morishita E.	Molecular genetic analysis of inherited protein C deficiency caused by the novel large deletion across two exons of PROC.	Thromb Res	188	115-118	2020
小林隆夫	肺血栓塞栓症. 特集 妊産婦死亡をどう防ぐかI	産婦人科の実際	68(12)	1455-1463	2019
小林隆夫	連載「DIC診療の新たな展開」第6回 産科DICにおける遺伝子組換えトロンボモジュリン製剤の有用性	Thromb Med	9(2)	63-68	2019
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Ochiai M, Kurata H, Inoue H, Ichiyama M, Fujiyoshi J, Watabe S, Hiroma T, Nakamura T, Ohga S.	Transcutaneous blood gas monitoring among neonatal intensive care units in Japan.	Pediatr Int	62(2)	169-174	2020
Taira R, Inoue H, Sawano T, Fujiyoshi J, Ichimiya Y, Torio M, Sanefuji M, Ochiai M, Sakai Y, Ohga S.	Management of apnea in infants with trisomy 18.	Dev Med Child Neurol			2019
Sonoda M, Ishimura M, Eguchi K, Shiraishi A, Kanno S, Kaku N, Inoue H, Motomura Y, Ochiai M, Sakai Y, Nakayama M, Ohara O, Ohga S.	Prognostic factors for survival of herpes simplex virus-associated hemophagocytic lymphohistiocytosis.	Int J Hematol	111(1)	131-136	2020

Matsushita Y, Sakai Y, Torio M, Inoue H, Ochiai M, Yasuoka K, Kurata H, Fujiyoshi J, Ichiyama M, Taguchi T, Kato K, Ohga S; Neonatal Research Network of Japan (NRNJ).	Association of perinatal factors of epilepsy in very low birth weight infants, using a nationwide database in Japan.	J Perinatol	39(11)	1472-1479	2019
Ochiai M, Nagata H, Tanaka K, Ihara K, Ohga S.	Critical association of Pallister-Hall syndrome and congenital heart disease.	Pediatr Int.	61(8)	827-828	2019
Ishimura M, Ohga S.	[Cancer predisposition in inherited bone marrow failure syndromes and primary immunodeficiency diseases].	Rinsho Ketsueki	60(6)	702-707	2019
Ishimura M, Eguchi K, Shiraishi A, Sonoda M, Azuma Y, Yamamoto H, Imadome KI, Ohga S.	Systemic Epstein-Barr Virus-Positive T/NK Lymphoproliferative Diseases With SH2D1A/XIAP Hypomorphic.	Front Pediatr	7	183	2019
Yasuoka K, Inoue H, Egami N, Ochiai M, Tanaka K, Sawano T, Kurata H, Ichiyama M, Fujiyoshi J, Matsushita Y, Sakai Y, Ohga S	Neonatal Research Network of Japan. Late-Onset Circulatory Collapse and Risk of Cerebral Palsy in Extremely Preterm Infants.	J Pediatr	212	117-123	2019
Ichimiya Y, Sonoda M, Ishimura M, Kanno S, Ohga S	Hemorrhagic Pneumonia as the First Manifestation of Anhidrotic Ectodermal Dysplasia with Immunodeficiency.	J Clin Immunol	39(3)	264-266	2019
Ogiwara K, Nogami K, Mizumachi K, Nakagawa T, Noda N, Ohga S, Shima M.	Hemostatic assessment of combined anticoagulant therapy using warfarin and prothrombin complex concentrates in a case of severe protein C deficiency.	Int J Hematol.	109(6)	650-656	2019
Taguchi T, Yanagi Y, Yoshimaru K, Zhang XY, Matsuura T, Nakayama K, Kobayashi E, Yamaza H, Nonaka K, Ohga S, Yamaza T.	Regenerative medicine using stem cells from human exfoliated deciduous teeth (SHED): a promising new treatment in pediatric surgery.	Surg Today	49(4)	316-322	2019

Fujiyoshi J, Yamaza H, Sonoda S, Yuniartha R, Ihara K, Nonaka K, Taguchi T, Ohga S, Yamaza T.	Therapeutic potential of hepatocyte-like-cells converted from stem cells from human exfoliated deciduous teeth in fulminant Wilson's disease.	Sci Rep	9(1)	1535	2019
Kurata H, Ochiai M, Inoue H, Ichiyama M, Yasuoka K, Fujiyoshi J, Matsushita Y, Honjo S, Sakai Y, Ohga S; Neonatal Research Network of Japan.	A nationwide survey on tracheostomy for very-low-birth-weight infants in Japan.	Pediatr Pulmonol	54(1)	53-60	2019
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Ichiyama M, Inoue H, Ochiai M, Ishimura M, Shiraishi A, Fujiyoshi J, Yamashita H, Sato K, Matsumoto S, Hotta T, Uchiumi T, Kang D, Ohga S.	Diagnostic challenge of the newborn patients with heritable protein C deficiency.	J Perinatol	39(2)	212-219	2019
Sonoda Y, Yamamura K, Ishii K, Ohkubo K, Ihara K, Sakai Y, Ohga S.	A Child with Prostaglandin I(2)-associated Thyrotoxicosis: Case Report.	J Clin Res Pediatr Endocrinol	11(2)	207-210	2019
Kinjo T, Inoue H, Kusuda T, Fujiyoshi J, Ochiai M, Takahata Y, Honjo S, Koga Y, Hara T, Ohga S.	Chemokine levels predict progressive liver disease in Down syndrome patients with transient abnormal myelopoiesis.	Pediatr Neonatol	60(4)	382-388	2019
Nagao A, Suzuki N, Takedani H, Yamasaki N, Chikasawa Y, Sawada A, Kanematsu T, Nojima M, Higasa S, Amano K, Fukutake K, Fujii T, Matsushita T, Suzuki T.	Ischaemic events are rare, and the prevalence of hypertension is not high in Japanese adults with haemophilia: First multicentre study in Asia.	Haemophilia	25(4)	e223-e230	2019

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Suzuki A, Suzuki N, Kanematsu T, Shinohara S, Arai N, Kikuchi R, Matsushita T.	Performance evaluation of Revohem TM FVIII chromogenic and Revohem TM FIX chromogenic in the CS-5100 autoanalyser.	Int J Lab Hematol	41(5)	664-670	2019
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Yaish H, Matsushita T, Belhani M, Jiménez-Yuste V, Kavakli K, Korsholm L, Matytsina I, Philipp C, Reichwald K, Wu R	Safety and efficacy of turoctocog alfa in the prevention and treatment of bleeds in previously untreated paediatric patients with severe haemophilia A: Results from the guardian 4 multinational clinical trial.	Haemophilia	26(1)	64-72	2020
Ogawa M, Suzuki N, Takahashi N, Tamura S, Suzuki A, Suzuki S, Hattori Y, Kakihara M, Kanematsu T, Kojima T, Katsumi A, Hayakawa F, Kojima T, Ishiguro N, Kiyoi H, Matsushita T	Higher FVIII:C measured by chromogenic substrate assay than by one-stage assay is associated with silent hemophilic arthropathy	Thromb Res	Epub ahead of print		2020
松下正, 長谷川雄一, 玉井佳子, 宮田茂樹, 安村敏, 山本晃士, 松本雅則, 日本輸血・細胞治療学会「ガイドライン委員会」	科学的根拠に基づいた新鮮凍結血漿(FFP)の使用ガイドライン[改訂第2版]	日本細胞治療学会誌	65巻3号	525-537	2019

Sugawara M, Goto Y, Yamazaki T, Teramoto T, Oikawa S, Shimada K, Uchiyama S, Ando K, Ishizuka N, Murata M, Yokoyama K, Uemura Y, Ikeda Y	Low-Dose Aspirin for Primary Prevention of Cardiovascular Events in Elderly Japanese Patients with Atherosclerotic Risk Factors: Subanalysis of a Randomized Clinical Trial (JPPP-70).	Am J Cardiovasc Drugs.		doi: 10.1007/s40256-018-0313-0.	2018
Tozawa K, Ono-Uruga Y, Yazawa M, Mori T, Murata M, Okamoto S, Ikeda Y, Matsubara Y	Unique megakaryocytes and platelets from novel human adipose-derived mesenchymal stem cell line.	Blood.		doi: https://doi.org/10.1182/blood-2018-04-842641	2018
Nakayama H, Kikuchi T, Abe R, Tozawa K, Watanuki S, Shimizu T, Mitsuhashi T, Murata M, Okamoto S, Mori T	Evans syndrome complicated with multicentric Castleman disease successfully treated with tocilizumab.	Rinsho Ketsueki.	59 (8)	997-1001. doi: 10.11406/rinketsu.59.997.	2018
Wakui M, Fujimori Y, Katagiri H, Nakamura S, Kondo Y, Kuroda Y, Nakagawa T, Shimizu N, Murata M	Assessment of in vitro effects of direct thrombin inhibitors and activated factor X inhibitors through clot waveform analysis.	J Clin Pathol.	72 (3)	244-250. doi: 10.1136/jclinpath-2018-205517	2019
村田満	偽性血小板減少症	ドクターサロン	62	19 (259)-22 (262)	2018
Akuta K, Kashiwagi H, Yujiri T, Nishiura N, Morikawa Y, Kato H, Honda S, Kanakura Y, Tomiyama Y	A unique phenotype of acquired Glanzmann thrombasthenia due to non-function-blocking anti- α IIb β 3 autoantibodies.	J Thromb Haemost	17	206-219	2019
富山佳昭	ITPに対する治療の進展-TPO受容体作動薬、リツキシマブの臨床効果	血液内科	77	54-60	2018
富山佳昭	抗血小板抗体の検出とその臨床的意義	日本輸血細胞治療学会誌	64	681-687	2018
Kuwana M	Personalized medicine in connective tissue disease: historical and future perspectives.	Pers. Med. Univers	7 (1)	1-6	2018

Oku K, Atsumi T, Akiyama Y, Amano H, Azuma N, Bohgaki T, Horita T, Hosoya T, Ichinose K, Kato M, Katsumata Y, Kawaguchi Y, Kawakami A, Koga T, Kohsaka H, Kondo Y, Kubo K, Kuwana M, et al.	Evaluation of the alternative classification criteria of systemic lupus erythematosus established by Systemic Lupus International Collaborating Clinics (SLICC)	Mod. Rheumatol	28(4)	642-648	2018
Rauch J, Salem D, Subang R, Kuwana M, and Levine JS	β 2-glycoprotein I-reactive T cells in autoimmune disease	Front. Immunol	9(12)	2836	2018
Salem D, Subang R, Kuwana M, Levine JS, and Rauch J	T cells from induced and spontaneous models of SLE recognize a common T cell epitope on β 2-glycoprotein I	Cell. Mol. Immunol			Epub ahead of print
桑名正隆	後天性無巨核球形血小板減少症	血液内科	77(1)	66-70	2018
桑名正隆	特発性血小板減少症紫斑病 (ITP) の診断	日本血栓止血学会誌	29(6)	625-629	2018
Ikeda Y, Yamanouchi J, Hato T, Yasukawa M, Takenaka K	Safe childbirth for a type I antithrombin-deficient woman with novel mutation in the SERPINC1 gene undergoing antithrombin concentrate therapy.	<u>Blood</u> <u>Coagul</u> <u>Fibrinolysis</u>	30(1)	47-51	2019
Casey N, Fujiwara H, Azuma T, Murakami Y, Yoshimitsu M, Masamoto I, Nawa Y, Yamanouchi J, Narumi H, Yakushijin Y, Hato T, Yasukawa M	An unusual, CD4 and CD8 dual-positive, CD25 negative, tumor cell phenotype in a patient with adult T-cell leukemia/lymphoma.	Leuk Lymphoma	59(11)	2740-2742	2018
Ikeda Y, Yamanouchi J, Kumon Y, Yasukawa M, Hato T	Association of platelet response to cilostazol with clinical outcome and CYP genotype in patients with cerebral infarction.	Thromb Res	172	14-20	2018
Yamanouchi J, Tokumoto D, Ikeda Y, Maruta M, Kaneko M, Hato T, Yasukawa M	Development of an FVIII Inhibitor in a Mild Hemophilia Patient with a Phe595Cys Mutation	Internal Med	57(21)	3179-3182	2018

Kawano N, Yokota-Ikeda N, Kawano S, Kuriyama T, Yamashita K, Ono N, Ueda N, Ochiai H, Ishikawa F, Kikuchi I, Shimoda K, Matsumoto M.	Clinical effect of rituximab as early administration for refractory thrombotic thrombocytopenic purpura associated with connective tissue diseases.	Modern Rheumatology Case Reports	2	59-67	2018
Yamashita M, Matsumoto M, Hayakawa M, Sakai K, Fujimura Y, Ogata N.	Intravitreal injection of aflibercept, an anti-VEGF antagonist, down-regulates plasma von Willebrand factor in patients with age-related macular degeneration.	Sci Rep	24	1491	2018
Itami H, Hara S, Matsumoto M, Imamura S, Kanai R, Nishiyama K, Ishimura M, Ohga S, Yoshida M, Tanaka R, Ogawa Y, Asada Y, Sekita-Hatakeyama Y, Hatakeyama K, Ohbayashi C.	Complement activation associated with ADAMTS13 deficiency may contribute to the characteristic glomerular manifestations in Upshaw-Schulman syndrome.	Thromb Res.	1 (170)	148-155	2018
Wada H, Matsumoto T, Suzuki K, Imai H, Katayama N, Iba T, Matsumoto M.	Differences and similarities between disseminated intravascular coagulation and thrombotic microangiopathy.	Thromb J.	16	14	2018
van Dorland HA, Mansouri Taleghani M, Sakai K, Friedman KD, George JN, Hrachovinova I, Knöbl PN, von Krogh AS, Schneppenheim R, Aebi-Huber I, Bütikofer L, Largiadèr CR, Cermakova Z, Kokame K, Miyata T, Yagi H, Terrell DR, Vesely SK, Matsumoto M, Lämmle B, Fujimura Y, Kremer Hovinga JA; Hereditary TTP Registry.	The International Hereditary Thrombotic Thrombocytopenic Purpura Registry: Key findings at enrolment until 2017.	Haematologica		doi:10.3324/haematol.2019.216796.	2019

Hayakawa M, Kato S, Matsui T, Sakai K, Fujimura Y, Matsumoto M.	Blood group antigen A on von Willebrand factor is more protective against ADAMTS13 cleavage than antigens B and H.	J Thromb Haemost	In press	In press	2019
Horiuchi H, Doman T, Kokame K, Saiki Y, Matsumoto M.	Acquired von Willebrand Syndrome Associated with Cardiovascular Diseases.	J Thromb	26(4)	303-314	2019
宮川義隆	特発性血小板減少性紫斑病 (ITP) 合併妊娠への対応	血栓止血誌	29	634-636	2018
Kato H, et al.	Safety and effectiveness of eculizumab for adult patients with atypical hemolytic uremic syndrome in Japan: interim analysis of post-marketing surveillance	Clin Exp Nephrol	23	65-76	2019
Ito S, et al.	Safety and effectiveness of eculizumab for pediatric patients with atypical hemolytic-uremic syndrome in Japan: interim analysis of post-marketing surveillance	Clin Exp Nephrol	23	112-121	2019
Fujisawa M, Kato H, Yoshida Y, Usui T, Takata M, Fujimoto M, Wada H, Uchida Y, Kokame K, Matsumoto M, Fujimura Y, Miyata T, Nangaku M	Clinical characteristics and genetic backgrounds of Japanese patients with atypical hemolytic uremic syndrome	Clin Exp Nephrol	22 (5)	1088-1099	2018
Maruyama K, Akiyama M, Miyata T, Kokame K	Protein S K196E mutation reduces its cofactor activity for APC but not for TFPI	Res Pract Thromb Haemost	2 (4)	751-756	2018
Tsuda M, Shiratsuchi M, Nakashima Y, Ikeda M, Muta H, Narazaki T, Masuda T, Kimura D, Takamatsu A, Matsumoto M, Fujimura Y, Kokame K, Matsushima T, Ogawa Y	Upshaw-Schulman syndrome diagnosed during pregnancy complicated by reversible cerebral vasoconstriction syndrome	Transfus Apher Sci	57 (6)	790-792	2018

Miyoshi T, Oku H, Asahara S, Okamoto A, Kokame K, Nakai M, Nishimura K, Otsuka F, Higashiyama A, Yoshimatsu J, Miyata T	Effects of low-dose combined oral contraceptives and protein S K196E mutation on anticoagulation factors: a prospective observational study	Int J Hematol	In press	In press	In press
Horiuchi H, Doman T, Kokame K, Saiki Y, Matsumoto M	Acquired von Willebrand Syndrome Associated with Cardiovascular Diseases.	J Atheroscler Thromb	26 (4)	303-314	2019
小亀浩市	ADAMTS13	日本血栓止血学会誌	29 (6)	586-588	2018
Ito S, Hidaka Y, Inoue N, Kaname S, Kato H, Matsumoto M, Miyakawa Y, Mizuno M, Okada H, Shimono A, Matsuda T, Maruyama S, Fujimura Y, Nangaku M, Kagami S.	Safety and effectiveness of eculizumab for pediatric patients with atypical hemolytic-uremic syndrome in Japan: interim analysis of post-marketing surveillance.	Clin Exp Nephrol.	23	112-121	2019
Kato H, Miyakawa Y, Hidaka Y, Inoue N, Ito S, Kagami S, Kaname S, Matsumoto M, Mizuno M, Matsuda T, Shimono A, Maruyama S, Fujimura Y, Nangaku M, Okada H.	Safety and effectiveness of eculizumab for adult patients with atypical hemolytic-uremic syndrome in Japan: interim analysis of post-marketing surveillance.	Clin Exp Nephrol.	23	65-75	2019
Ikeda Y, Yoshida Y, Sugawara Y, Nangaku M.	Geographic Diversity in Atypical Hemolytic Uremic Syndrome (aHUS): The Genetic Background of aHUS Cohort in Japan.	Rare Dis Res Treat.	3(3)	1-3	2018
Yoshida Y, Kato H, Ikeda Y, Nangaku M.	Pathogenesis of Atypical Hemolytic Uremic Syndrome.	J Atheroscler Thromb.	26	99-110	2019
Yamamura T, Nozu K, Ueda H, Fujimaru R, Hisatomi R, Yoshida Y, Kato H, Nangaku M, Miyata T, Sawai T, Minamikawa S, Kaito H, Matsuo M, Iijima K.	Functional splicing analysis in an infantile case of atypical hemolytic uremic syndrome caused by digenic mutations in C3 and MCP genes.	J Hum Genet.	63	755-759	2018
Kato H, Nangaku M, Okada H, Kagami S.	Controversies of the classification of TMA and the terminology of aHUS.	Clin Exp Nephrol.	22	979-980	2018

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榑佑介, 石東光司, 上床武史, 森下英理子, 杉森宏	一過性脳虚血性発作を契機に診断し得た先天性プロテインC欠損症の1例	脳卒中	40(5)	377-381	2018
Kuma H, Matsuda R, Nakashima A, Motoyama K, Takazaki S, Hatae H, Jin X, Tsuda T, Tsuda H, and Hamasaki N.	Protein S-specific activity assay system is not affected by direct oral anticoagulants.	Thromb Res.	168	60-62	2018
梶山倫未、安武健一郎、森口里利子、宮崎瞳、阿部志磨子、増田隆、今井克己、岩本昌子、津田博子、大部正代、河手久弥、上野宏美、小野美咲、川崎遥香、能口健太、市川彩絵、鬼木愛子、前田翔子、中野修治	食物摂取頻度調査報告 (FFQ 中村) で推定された女子大学生のナトリウム、カリウム摂取量の妥当性：24時間尿中排泄量との比較	中村学園大学・中村学園大学短期大学部研究紀要	51	105-111	2019
Yoshida R, Seki S, Hasegawa J, Koyama T, Yamazaki K, Takagi A, Kojima T, Yoshimura M	Familial pulmonary thromboembolism with a prothrombin mutation and antithrombin resistance.	Cardiol Cases.	17(6)	197-199	2018
Sanda N, Suzuki N, Suzuki A, Kanematsu T, Kishimoto M, Hasuwa H, Takagi A, Kojima T, Matsushita T, Nakamura S	Vwf K1362A resulted in failure of protein synthesis in mice.	Int J Hematol.	107(4)	428-35	2018
Tamura S, Suga Y, Tanamura M, Murata-Kawakami M, Takagi Y, Hattori Y, Kakiyama M, Suzuki S, Takagi A, Kojima T	Optimisation of antithrombin resistance assay as a practical clinical laboratory test: development of prothrombin activator using factors Xa/Va and automation of assay.	Int J Lab Hematol.	40(3)	312-319	2018

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Morikawa M, Umazume T, Hosokawa- Miyanishi A, Watari H, Kobayashi T, Seki H, Saito S	Relationship between antithrombin activity and interval from diagnosis to delivery among pregnant women with early-onset pre-eclampsia	Int J Gynaecol Obstet	82(2)	555-560	2019
杉浦和子, 尾島俊之, 浦野哲盟, 小林隆夫	わが国における経口避妊薬に関連した血栓塞栓症の年齢別発症数とその予後	日産婦誌	71(3)	546	2019
小林隆夫	エビデンスからみるOC・EP配合剤の血栓症リスクとマネジメント	Female Hormone Digest	第2回	1-5	2018
小林隆夫, 杉浦和子	連載「周術期に留意すべき凝固異常」第4回 後天性血栓性素因～女性ホルモン剤～	Thromb Med	8(3)	206-210	2018
小林隆夫	月経困難症治療におけるルナベル配合錠ULDの適正使用ー血栓症リスクを考えるー	東京	194	i-vii	2018
市山正子, 石村匡崇, 落合正行, 大賀正一	遺伝性血栓症	周産期医学	48	1397-1400	2018
大賀正一, 市山正子, 落合正行, 石村匡崇, 堀田多恵子, 内海健, 後藤和人, 康東天	小児の遺伝性血栓性素因の特徴とその治療	日本検査血液学会雑誌	19		2018
落合正行, 石村匡崇, 市山正子, 石黒精, 大賀正一	特集 血栓・塞栓症ー診断・治療・予防の最新動向ー新生児・小児の血栓症	日本臨床	76		2018
石黒精, 大賀正一	小児科で遭遇する血栓性疾患	日本血栓止血学会誌			2018

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Azuma K, Osawa Y, Tabata S, Katsukawa F, Ishida H, Oguma Y, Kawai T, Itoh H, Okuda H, Oguchi S, Ohta A, Kikuchi H, Murata M, Matsumoto H	Decrease in regional body fat after long-term high-intensity interval training.	The Journal of Physical Fitness and Sports Medicine.	Vol. 6	103-110	2017
Fujio Y, Kojima K, Hashiguchi M, Wakui M, Murata M, Amagai M, Yamagami J	Validation of chemiluminescent enzyme immunoassay in detection of autoantibodies in pemphigus and pemphigoid.	J Dermatol Sci	85/ 3	208-215	2017
Toyofuku M, Morozumi M, Hida M, Satoh Y, Sakata H, Shiro H, Ubukata K, Murata M, Iwata, S.	Effects of Intrapartum Antibiotic Prophylaxis on Neonatal Acquisition of Group B Streptococci.	The Journal of Pediatrics.	190	169-173	2017
大平賢太郎、藤森祐多、片桐尚子、清水長子、三ツ橋雄之、涌井昌俊、村田満	Xa阻害薬のPT, APTTと抗Xa活性測定法によるモニタリングの有用性検討	日本検査血液学会雑誌	第58巻 第4号	271-281	2017
村田満	《特集》血栓止血の臨床化学	臨床化学	Vol. 46 No. 2	94	2017
Rubic-Schneider T, Kuwana M, Christen B, Assenmacher M, Hainzl O, Zimmermann F, Fischer R, Koppenburg V, Chibout SD, Wright TM, Seidl A, and Kammüller M	T cell assays confirm immunogenicity of tungsten-induced erythropoietin aggregates associated with pure red cell aplasia	Blood Adv	1(6)	367-379	2017
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