Sverre Sandberg

List of Publications by Year in descending order

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		61984	79698
303	8,109	43	73
papers	citations	h-index	g-index
322	322	322	5332
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	Defining analytical performance specifications: Consensus Statement from the 1st Strategic Conference of the European Federation of Clinical Chemistry and Laboratory Medicine. Clinical Chemistry and Laboratory Medicine, 2015, 53, 833-5.	2.3	398
2	Current Issues in Measurement and Reporting of Urinary Albumin Excretion. Clinical Chemistry, 2009, 55, 24-38.	3.2	298
3	Preanalytical quality improvement: from dream to reality. Clinical Chemistry and Laboratory Medicine, 2011, 49, 1113-26.	2.3	256
4	The incidence of inherited porphyrias in Europe. Journal of Inherited Metabolic Disease, 2013, 36, 849-857.	3.6	220
5	Preanalytical quality improvement: in quality we trust. Clinical Chemistry and Laboratory Medicine, 2013, 51, 229-241.	2.3	162
6	The Biological Variation Data Critical Appraisal Checklist: A Standard for Evaluating Studies on Biological Variation. Clinical Chemistry, 2018, 64, 501-514.	3.2	152
7	From biomarkers to medical tests: The changing landscape of test evaluation. Clinica Chimica Acta, 2014, 427, 49-57.	1.1	148
8	Criteria for assigning laboratory measurands to models for analytical performance specifications defined in the 1st EFLM Strategic Conference. Clinical Chemistry and Laboratory Medicine, 2017, 55, 189-194.	2.3	130
9	A checklist for critical appraisal of studies of biological variation. Clinical Chemistry and Laboratory Medicine, 2015, 53, 879-85.	2.3	120
10	Instruments for Self-Monitoring of Blood Glucose: Comparisons of Testing Quality Achieved by Patients and a Technician. Clinical Chemistry, 2002, 48, 994-1003.	3.2	118
11	Confidence Intervals and Power Calculations for Within-Person Biological Variation: Effect of Analytical Imprecision, Number of Replicates, Number of Samples, and Number of Individuals. Clinical Chemistry, 2012, 58, 1306-1313.	3.2	118
12	EXPLORE: A Prospective, Multinational, Natural History Study of Patients with Acute Hepatic Porphyria with Recurrent Attacks. Hepatology, 2020, 71, 1546-1558.	7.3	103
13	Porphyrin-induced photodamage at the cellular and the subcellular level as related to the solubility of the porphyrin. Clinica Chimica Acta, 1981, 109, 193-201.	1.1	98
14	Biological Variation: The Effect of Different Distributions on Estimated Within-Person Variation and Reference Change Values. Clinical Chemistry, 2016, 62, 725-736.	3.2	94
15	Recurrent attacks of acute hepatic porphyria: major role of the chronic inflammatory response in the liver. Journal of Internal Medicine, 2018, 284, 78-91.	6.0	88
16	Cobalamin Deficiency in General Practice. Assessment of the Diagnostic Utility and Cost-Benefit Analysis of Methylmalonic Acid Determination in Relation to Current Diagnostic Strategies. Clinical Chemistry, 1999, 45, 189-198.	3.2	87
17	A multicentre study of reference intervals for haemoglobin, basic blood cell counts and erythrocyte indices in the adult population of the Nordic countries. Scandinavian Journal of Clinical and Laboratory Investigation, 2004, 64, 385-398.	1.2	79
18	Weekly and 90-Minute Biological Variations in Cardiac Troponin T and Cardiac Troponin I in Hemodialysis Patients and Healthy Controls. Clinical Chemistry, 2014, 60, 838-847.	3.2	77

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19	Sample collections from healthy volunteers for biological variation estimates' update: a new project undertaken by the Working Group on Biological Variation established by the European Federation of Clinical Chemistry and Laboratory Medicine. Clinical Chemistry and Laboratory Medicine, 2016, 54, 1599-1608.	2.3	76
20	The EuBIVAS: Within- and Between-Subject Biological Variation Data for Electrolytes, Lipids, Urea, Uric Acid, Total Protein, Total Bilirubin, Direct Bilirubin, and Glucose. Clinical Chemistry, 2018, 64, 1380-1393.	3.2	75
21	A systematic review of data on biological variation for alanine aminotransferase, aspartate aminotransferase and Î ³ -glutamyl transferase. Clinical Chemistry and Laboratory Medicine, 2013, 51, 1997-2007.	2.3	74
22	The Index of Individuality Is Often a Misinterpreted Quantity Characteristic. Clinical Chemistry and Laboratory Medicine, 1999, 37, 655-61.	2.3	68
23	Between-Lot Variation in External Quality Assessment of Glucose: Clinical Importance and Effect on Participant Performance Evaluation. Clinical Chemistry, 2005, 51, 1632-1636.	3.2	68
24	Standardized Evaluation of Instruments for Self-Monitoring of Blood Glucose by Patients and a Technologist. Clinical Chemistry, 2004, 50, 1068-1071.	3.2	67
25	Influence of pH on porphyrin production in Propionibacterium acnes. Archives of Dermatological Research, 1984, 276, 396-400.	1.9	66
26	The EuBIVAS Project: Within- and Between-Subject Biological Variation Data for Serum Creatinine Using Enzymatic and Alkaline Picrate Methods and Implications for Monitoring. Clinical Chemistry, 2017, 63, 1527-1536.	3.2	66
27	The use of error and uncertainty methods in the medical laboratory. Clinical Chemistry and Laboratory Medicine, 2018, 56, 209-219.	2.3	66
28	Evidence-Based Guidelines in Laboratory Medicine: Principles and Methods. Clinical Chemistry, 2004, 50, 806-818.	3.2	65
29	Glycemic control and complications in patients with type 1 diabetesÂ-Âa registry-based longitudinal study of adolescents and young adults. Pediatric Diabetes, 2017, 18, 188-195.	2.9	62
30	Porphyrin-sensitized photodynamic damage of isolated rat liver mitochondria. Biochimica Et Biophysica Acta - Bioenergetics, 1980, 593, 187-195.	1.0	59
31	Hyperkinetic or Attention Deficit Disorder. British Journal of Psychiatry, 1996, 169, 10-17.	2.8	58
32	Patient-derived Quality Specifications for Instruments Used in Self-Monitoring of Blood Glucose. Clinical Chemistry, 2001, 47, 67-73.	3.2	56
33	Strategies to define performance specifications in laboratory medicine: 3 years on from the Milan Strategic Conference. Clinical Chemistry and Laboratory Medicine, 2017, 55, 1849-1856.	2.3	56
34	Mechanisms of photosensitivity in porphyric patients with special emphasis on erythropoietic protoporphyria. Journal of Photochemistry and Photobiology B: Biology, 1991, 10, 285-302.	3.8	53
35	Position paper of the EPMA and EFLM: a global vision of the consolidated promotion of an integrative medical approach to advance health care. EPMA Journal, 2013, 4, 12.	6.1	53
36	Estimation and Application of Biological Variation of Urinary δ-Aminolevulinic Acid and Porphobilinogen in Healthy Individuals and in Patients with Acute Intermittent Porphyria. Clinical Chemistry, 2006, 52, 650-656.	3.2	51

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37	Biological Variation Estimates Obtained from 91 Healthy Study Participants for 9 Enzymes in Serum. Clinical Chemistry, 2017, 63, 1141-1150.	3.2	51
38	Within-subject and between-subject biological variation estimates of 21 hematological parameters in 30 healthy subjects. Clinical Chemistry and Laboratory Medicine, 2018, 56, 1309-1318.	2.3	51
39	A comprehensive multiple matrix model representing the life cycle of the tick that transmits agent of lyme disease. Journal of Theoretical Biology, 1992, 157, 203-220.	1.7	50
40	Factors important for the measurement of chemiluminescence production by polymorphonuclear leukocytes. Journal of Immunological Methods, 1986, 88, 121-128.	1.4	49
41	Standardized Evaluation of Nine Instruments for Self-Monitoring of Blood Glucose. Diabetes Technology and Therapeutics, 2008, 10, 467-477.	4.4	49
42	Lead in Tissues of Deseased Lead Smelter Workers. Journal of Trace Elements in Medicine and Biology, 1995, 9, 136-143.	3.0	48
43	A new method for isolation of reticulocytes: positive selection of human reticulocytes by immunomagnetic separation. Blood, 1990, 76, 2397-2403.	1.4	45
44	Setting analytical performance specifications based on outcome studies – is it possible?. Clinical Chemistry and Laboratory Medicine, 2015, 53, 841-8.	2.3	45
45	Personalized Reference Intervals in Laboratory Medicine: A New Model Based on Within-Subject Biological Variation. Clinical Chemistry, 2021, 67, 374-384.	3.2	45
46	Decreased concentration of hemoglobin, accumulation of lipid oxidation products and unchanged skeletal muscle in Atlantic salmon (Salmo salar) fed low dietary vitamin E. Fish Physiology and Biochemistry, 1994, 12, 421-429.	2.3	44
47	Is Placenta a Good Indicator of Cadmium and Lead Exposure?. Archives of Environmental Health, 1996, 51, 389-394.	0.4	44
48	Within-subject biological variation of glucose and HbA1c in healthy persons and in type 1 diabetes patients. Clinical Chemistry and Laboratory Medicine, 2011, 49, 1501-7.	2.3	44
49	Influence of Index of Individuality on False Positives in Repeated Sampling from Healthy Individuals. Clinical Chemistry and Laboratory Medicine, 2001, 39, 160-5.	2.3	43
50	Self-Monitoring of Blood Glucose in Type 1 Diabetes Patients with Insufficient Metabolic Control: Focused Self-Monitoring of Blood Glucose Intervention Can Lower Glycated Hemoglobin A1C. Journal of Diabetes Science and Technology, 2009, 3, 83-88.	2.2	42
51	Diagnosing Diabetes Mellitus: Performance of Hemoglobin A1c Point-of-Care Instruments in General Practice Offices. Clinical Chemistry, 2013, 59, 1790-1801.	3.2	42
52	Comparison of transmission rates of HIV-1 and HIV-2 in a cohort of prostitutes in Senegal. Bulletin of Mathematical Biology, 1993, 55, 731-743.	1.9	41
53	Quality of Care for Patients With Type 2 Diabetes in Primary Care in Norway Is Improving: Results of cross-sectional surveys of 33 general practices in 1995 and 2005. Diabetes Care, 2009, 32, 81-83.	8.6	41
54	Nursing home patients with diabetes: Prevalence, drug treatment and glycemic control. Diabetes Research and Clinical Practice, 2014, 105, 102-109.	2.8	41

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55	Improving quality in the preanalytical phase through innovation, on behalf of the European Federation for Clinical Chemistry and Laboratory Medicine (EFLM) Working Group for Preanalytical Phase (WG-PRE). Clinical Chemistry and Laboratory Medicine, 2017, 55, 489-500.	2.3	41
56	PORPHYRIN-INDUCED PHOTODAMAGE TO ISOLATED HUMAN NEUTROPHILS. Photochemistry and Photobiology, 1981, 34, 471-475.	2.5	41
57	Postanalytical External Quality Assessment of Urine Albumin in Primary Health Care: An International Survey. Clinical Chemistry, 2008, 54, 1630-1636.	3.2	40
58	How to achieve harmonisation of laboratory testing —The complete picture. Clinica Chimica Acta, 2014, 432, 8-14.	1.1	40
59	Systematic review and meta-analysis of within-subject and between-subject biological variation estimates of 20 haematological parameters. Clinical Chemistry and Laboratory Medicine, 2019, 58, 25-32.	2.3	40
60	The European Biological Variation Study (EuBIVAS): a summary report. Clinical Chemistry and Laboratory Medicine, 2022, 60, 505-517.	2.3	40
61	Influence of tetracyclines on human polymorphonuclear leukocyte function. Antimicrobial Agents and Chemotherapy, 1984, 25, 354-357.	3.2	39
62	Familial and Sporadic Porphyria Cutanea Tarda: Characterization and Diagnostic Strategies. Clinical Chemistry, 2009, 55, 795-803.	3.2	39
63	Biomarker development targeting unmet clinical needs. Clinica Chimica Acta, 2016, 460, 211-219.	1.1	39
64	European Biological Variation Study (EuBIVAS): Within- and Between-Subject Biological Variation Data for 15 Frequently Measured Proteins. Clinical Chemistry, 2019, 65, 1031-1041.	3.2	39
65	Effect of Ambient Temperature on Analytical Performance of Self-Monitoring Blood Glucose Systems. Diabetes Technology and Therapeutics, 2011, 13, 883-892.	4.4	38
66	European Specialist Porphyria Laboratories: Diagnostic Strategies, Analytical Quality, Clinical Interpretation, and Reporting As Assessed by an External Quality Assurance Program. Clinical Chemistry, 2011, 57, 1514-1523.	3.2	38
67	Biological variation estimates for prostate specific antigen from the European Biological Variation Study; consequences for diagnosis and monitoring of prostate cancer. Clinica Chimica Acta, 2018, 486, 185-191.	1.1	37
68	The susceptibility to nephrotoxicity of streptozotocin-induced diabetic rats subchronically exposed to cadmium chloride in drinking water. Toxicology, 1999, 142, 69-75.	4.2	36
69	How to conduct External Quality Assessment Schemes for the pre-analytical phase?. Biochemia Medica, 2014, 24, 114-122.	2.7	36
70	Type 2 diabetes in general practice in Norway 2005–2014: moderate improvements in risk factor control but still major gaps in complication screening. BMJ Open Diabetes Research and Care, 2017, 5, e000459.	2.8	35
71	Models for Combining Random and Systematic Errors. Assumptions and Consequences for different Models. Clinical Chemistry and Laboratory Medicine, 2001, 39, 589-95.	2.3	34
72	Biological variation – reliable data is essential. Clinical Chemistry and Laboratory Medicine, 2015, 53, 153-4.	2.3	34

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73	Dysregulation of homocysteine homeostasis in acute intermittent porphyria patients receiving heme arginate or givosiran. Journal of Inherited Metabolic Disease, 2021, 44, 961-971.	3.6	34
74	Postanalytical External Quality Assessment of Blood Glucose and Hemoglobin A1c: An International Survey. Clinical Chemistry, 2005, 51, 1145-1153.	3.2	33
75	Harmonization initiatives in the generation, reporting and application of biological variation data. Clinical Chemistry and Laboratory Medicine, 2018, 56, 1629-1636.	2.3	33
76	Light-induced protoporphyrin release from erythrocytes in erythropoietic protoporphyria Journal of Clinical Investigation, 1982, 70, 693-698.	8.2	33
77	Phototoxicity of protoporphyrin as related to its subcellular localization in mice livers after short-term feeding with griseofulvin. Biochemical Journal, 1981, 198, 67-74.	3.7	32
78	Clinical assessment of haemoglobin values by general practitioners related to analytical and biological variation. Scandinavian Journal of Clinical and Laboratory Investigation, 1991, 51, 453-459.	1.2	32
79	Acute hepatic porphyria and cancer risk: a nationwide cohort study. Journal of Internal Medicine, 2017, 282, 229-240.	6.0	32
80	Systematic review of the biological variation data for diabetes related analytes. Clinica Chimica Acta, 2019, 488, 61-67.	1.1	32
81	estimates of I2-isomerized C-terminal telopeptide of type I collagen (I2-CTX), N-terminal propeptide of type I collagen (PINP), osteocalcin, intact fibroblast growth factor 23 and uncarboxylated-unphosphorylated matrix-Gla protein—a cooperation between the EFLM Working Group on Biological Variation and the International Osteoporosis Foundation-International	3.1	31
82	PHOTODYNAMIC RELEASE OF PROTOPORPHYRIN FROM INTACT ERYTHROCYTES IN ERYTHROPOIETIC PROTOPORPHYRIA: THE EFFECT OF SMALL REPETITIVE LIGHT DOSES. Photochemistry and Photobiology, 1985, 41, 535-541.	2.5	30
83	Results and Feasibility of an External Quality Assessment Scheme for Self-Monitoring of Blood Glucose. Clinical Chemistry, 2006, 52, 1311-1317.	3.2	29
84	Feasibility of using self-reported patient data in a national diabetes register. BMC Health Services Research, 2015, 15, 553.	2.2	29
85	ERYTHROPOIETIC PROTOPORPHYRIA: PHOTODYNAMIC TRANSFER OF PROTOPORPHYRIN FROM INTACT ERYTHROCYTES TO OTHER CELLS. Photochemistry and Photobiology, 1990, 51, 573-577.	2.5	28
86	Protoporphyrin-induced photodamage to mitochondria and lysosomes from rat liver. Clinica Chimica Acta, 1981, 111, 55-60.	1.1	27
87	Implications of probability analysis for interpreting results of leukocyte esterase and nitrite test strips Clinical Chemistry, 1989, 35, 1663-1668.	3.2	27
88	Automated counting of white and red blood cells in the cerebrospinal fluid. International Journal of Laboratory Hematology, 2000, 22, 203-210.	0.2	27
89	Excess risk of adverse pregnancy outcomes in women with porphyria: a populationâ€based cohort study. Journal of Inherited Metabolic Disease, 2011, 34, 217-223.	3.6	27
90	The Importance of Reagent Lot Registration in External Quality Assurance/Proficiency Testing Schemes. Clinical Chemistry, 2016, 62, 708-715.	3.2	27

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91	Biological variation data for lipid cardiovascular risk assessment biomarkers. A systematic review applying the biological variation data critical appraisal checklist (BIVAC). Clinica Chimica Acta, 2019, 495, 467-475.	1.1	27
92	Pharmacokinetics and Metabolism of the Antimalarial Piperaquine After Intravenous and Oral Single Doses to the Rat. Journal of Pharmaceutical Sciences, 2008, 97, 3400-3410.	3.3	25
93	Conformational stability and activity analysis of two hydroxymethylbilane synthase mutants, K132N and V215E, with different phenotypic association with acute intermittent porphyria. Bioscience Reports, 2013, 33, .	2.4	25
94	To report or not to report: a proposal on how to deal with altered test results in hemolytic samples. Clinical Chemistry and Laboratory Medicine, 2017, 55, 1109-1111.	2.3	25
95	American Liver Guidelines and Cutoffs for "Normal―ALT: A Potential for Overdiagnosis. Clinical Chemistry, 2017, 63, 1196-1198.	3.2	25
96	Acute Intermittent Porphyria: An Overview of Therapy Developments and Future Perspectives Focusing on Stabilisation of HMBS and Proteostasis Regulators. International Journal of Molecular Sciences, 2021, 22, 675.	4.1	25
97	The European Biological Variation Study (EuBIVAS): weekly biological variation of cardiac troponin I estimated by the use of two different high-sensitivity cardiac troponin I assays. Clinical Chemistry and Laboratory Medicine, 2020, 58, 1741-1747.	2.3	25
98	DOXYCYCLINE INDUCED PHOTODAMAGE TO HUMAN NEUTROPHILS AND TRYPTOPHAN. Photochemistry and Photobiology, 1984, 39, 43-48.	2.5	24
99	Interpretation of Hemoglobin A1c (HbA1c) Values among Diabetic Patients. Clinical Chemistry, 2001, 47, 1212-1217.	3.2	24
100	Do Guidelines for the Diagnosis and Monitoring of Diabetes Mellitus Fulfill the Criteria of Evidence-Based Guideline Development?. Clinical Chemistry, 2008, 54, 1872-1882.	3.2	24
101	Withinâ€subject biological variation of reticulocytes and reticulocyteâ€derived parameters. European Journal of Haematology, 1998, 61, 42-48.	2.2	24
102	Quality specifications derived from objective analyses based upon clinical needs. Scandinavian Journal of Clinical and Laboratory Investigation, 1999, 59, 531-534.	1.2	23
103	External Quality Assessment of Point-of-Care Methods: Model For Combined Assessment of Method Bias and Single-Participant Performance by the Use of Native Patient Samples and Noncommutable Control Materials. Clinical Chemistry, 2013, 59, 363-371.	3.2	23
104	Total error vs. measurement uncertainty: the match continues. Clinical Chemistry and Laboratory Medicine, 2016, 54, 195-6.	2.3	23
105	A pragmatic approach to sample acceptance and rejection. Clinical Biochemistry, 2017, 50, 579-581.	1.9	23
106	Setting clinical performance specifications to develop and evaluate biomarkers for clinical use. Annals of Clinical Biochemistry, 2019, 56, 527-535.	1.6	23
107	Behandlingen av type 1-diabetes i spesialisthelsetjenesten – data fra Norsk diabetesregister for voksne. Tidsskrift for Den Norske Laegeforening, 2013, 133, 2257-2261.	0.2	23
108	Proposal for the modification of the conventional model for establishing performance specifications. Clinical Chemistry and Laboratory Medicine, 2015, 53, 925-37.	2.3	22

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109	Porfyrisykdommer i Norge. Tidsskrift for Den Norske Laegeforening, 2014, 134, 831-835.	0.2	22
110	PORPHYRIN-INDUCED PHOTODAMAGE TO ISOLATED HUMAN NEUTROPHILS. Photochemistry and Photobiology, 1981, 34, 471-475.	2.5	21
111	Calcium, Magnesium, Albumin, and Total Protein Measurement in Serum as Assessed with 20 Fresh-Frozen Single-Donation Sera. Clinical Chemistry, 2012, 58, 1597-1599.	3.2	21
112	Effect of Participating in a Quality Improvement System over Time for Point-of-Care C-Reactive Protein, Glucose, and Hemoglobin Testing. Clinical Chemistry, 2016, 62, 1474-1481.	3.2	21
113	Is D-dimer used according to clinical algorithms in the diagnostic work-up of patients with suspicion of venous thromboembolism? A study in six European countries. Thrombosis Research, 2016, 142, 1-7.	1.7	21
114	Biological Variation of Cardiac Troponins in Health and Disease: A Systematic Review and Meta-analysis. Clinical Chemistry, 2021, 67, 256-264.	3.2	21
115	European Biological Variation Study (EuBIVAS): within- and between-subject biological variation estimates for serum thyroid biomarkers based on weekly samplings from 91 healthy participants. Clinical Chemistry and Laboratory Medicine, 2022, 60, 523-532.	2.3	21
116	Light-induced release of protoporphyrin, but not of zinc protoporphyrin, from erythrocytes in a patient with greatly elevated erythrocyte protoporphyrin. Blood, 1983, 62, 846-851.	1.4	20
117	Effect of tetracyclines and UV light on oxygen consumption by human leukocytes. Antimicrobial Agents and Chemotherapy, 1984, 26, 489-492.	3.2	20
118	Discrepancies in International Normalized Ratio Results between Instruments: A Model to Split the Variation into Subcomponents. Clinical Chemistry, 2010, 56, 1618-1626.	3.2	20
119	The erythrocyte sedimentation rate in general practice: Clinical assessment based on case histories. Scandinavian Journal of Clinical and Laboratory Investigation, 1994, 54, 291-300.	1.2	19
120	Providing Correct Estimates of Biological Variation—Not an Easy Task. The Example of S100-β Protein and Neuron-Specific Enolase. Clinical Chemistry, 2018, 64, 1537-1539.	3.2	19
121	Within- and between-subject biological variation data for tumor markers based on the European Biological Variation Study. Clinical Chemistry and Laboratory Medicine, 2022, 60, 543-552.	2.3	19
122	Practical guide for identifying unmet clinical needs for biomarkers. Electronic Journal of the International Federation of Clinical Chemistry and Laboratory Medicine, 2018, 29, 129-137.	0.7	19
123	External quality assessment of prothrombin time: The splitâ€sample model compared with external quality assessment with commercial control material. Scandinavian Journal of Clinical and Laboratory Investigation, 2006, 66, 337-350.	1.2	18
124	The Prevalence of Self-Monitoring of Blood Glucose and Costs of Glucometer Strips in a Nationwide Cohort. Diabetes Technology and Therapeutics, 2010, 12, 701-705.	4.4	18
125	External quality assessment of point-of-care International Normalized Ratio (INR) testing in Europe. Clinical Chemistry and Laboratory Medicine, 2012, 50, 81-8.	2.3	18
126	Selfâ€monitoring of blood glucose in patients with diabetes who do not use insulin—are guidelines evidenceâ€based?. Diabetic Medicine, 2012, 29, 1226-1236.	2.3	18

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127	Biological Variation of Hemoglobin A1c: Consequences for Diagnosing Diabetes Mellitus. Clinical Chemistry, 2014, 60, 1570-1572.	3.2	18
128	Analytical performance specifications based on how clinicians use laboratory tests. Experiences from a post-analytical external quality assessment programme. Clinical Chemistry and Laboratory Medicine, 2015, 53, 857-62.	2.3	18
129	Guidance for the design and reporting of studies evaluating the clinical performance of tests for present or past SARS-CoV-2 infection. BMJ, The, 2021, 372, n568.	6.0	18
130	Hematoporphyrin Derivative: Chemical Composition, Photochemical and Photosensitizing Properties. Advances in Experimental Medicine and Biology, 1983, 160, 165-179.	1.6	18
131	Lot-to-lot reagent verification: challenges and possible solutions. Clinical Chemistry and Laboratory Medicine, 2022, 60, 675-680.	2.3	18
132	Use and interpretation of HbA1ctesting in general practice. Implications for quality of care. Scandinavian Journal of Clinical and Laboratory Investigation, 2000, 60, 349-356.	1.2	17
133	Circadian variation in serum cortisol and circulating neutrophils are markers for circadian variation of bone marrow proliferation in cancer patients. European Journal of Haematology, 1993, 50, 206-212.	2.2	17
134	Estimates of Within-Subject Biological Variation of Protein C, Antithrombin, Protein S Free, Protein S Activity, and Activated Protein C Resistance in Pregnant Women. Clinical Chemistry, 2017, 63, 898-907.	3.2	17
135	Harmonization activities of Noklus – a quality improvement organization for point-of-care laboratory examinations. Clinical Chemistry and Laboratory Medicine, 2018, 57, 106-114.	2.3	17
136	Analytical Performance Specifications for Lipoprotein(a), Apolipoprotein B-100, and Apolipoprotein A-I Using the Biological Variation Model in the EuBIVAS Population. Clinical Chemistry, 2020, 66, 727-736.	3.2	17
137	Exponential increase in age-specific prevalence of ventricular dysrhythmia among males. Journal of Chronic Diseases, 1982, 35, 743-750.	1.2	16
138	Phototoxicity of tetracyclines as related to singlet oxygen production and uptake by polymorphonuclear leukocytes. Biochemical Pharmacology, 1986, 35, 2883-2885.	4.4	16
139	Erythropoietic protoporphyria: A quantitative determination of erythrocyte protoporphyrin in individual cells by flow cytometry. Scandinavian Journal of Clinical and Laboratory Investigation, 1988, 48, 261-267.	1.2	16
140	A Model for Setting Analytical Quality Specifications and Design of Control for Measurements on the Ordinal Scale. Clinical Chemistry and Laboratory Medicine, 2000, 38, 545-51.	2.3	16
141	Postanalytical External Quality Assessment of Warfarin Monitoring in Primary Healthcare. Clinical Chemistry, 2006, 52, 1871-1878.	3.2	16
142	Glucose meters – fit for clinical purpose. Clinical Chemistry and Laboratory Medicine, 2013, 51, 943-52.	2.3	16
143	International Porphyria Molecular Diagnostic Collaborative: an evidence-based database of verified pathogenic and benign variants for the porphyrias. Genetics in Medicine, 2019, 21, 2605-2613.	2.4	16
144	Diagnosing Acute Porphyrias. Clinical Chemistry, 2004, 50, 803-805.	3.2	15

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145	A model for calculating the within-subject biological variation and likelihood ratios for analytes with a time-dependent change in concentrations; exemplified with the use of D-dimer in suspected venous thromboembolism in healthy pregnant women. Annals of Clinical Biochemistry, 2012, 49, 561-569.	1.6	15
146	Interpretation and management of INR results: A case history based survey in 13 countries. Thrombosis Research, 2012, 130, 309-315.	1.7	15
147	A Bayesian Approach to Biological Variation Analysis. Clinical Chemistry, 2019, 65, 995-1005.	3.2	15
148	Porphyria cutanea tarda increases risk of hepatocellular carcinoma and premature death: a nationwide cohort study. Orphanet Journal of Rare Diseases, 2019, 14, 77.	2.7	15
149	Critical appraisal and meta-analysis of biological variation estimates for kidney related analytes. Clinical Chemistry and Laboratory Medicine, 2022, 60, 469-478.	2.3	15
150	Biological variation estimates of thyroid related measurands– meta-analysis of BIVAC compliant studies. Clinical Chemistry and Laboratory Medicine, 2022, 60, 483-493.	2.3	15
151	Self-monitoring of blood glucose with a focus on analytical quality: an overview. Clinical Chemistry and Laboratory Medicine, 2010, 48, 963-972.	2.3	14
152	The European Biological Variation Study (EuBIVAS): Biological Variation Data for Coagulation Markers Estimated by a Bayesian Model. Clinical Chemistry, 2021, 67, 1259-1270.	3.2	14
153	Availability and analytical quality of hemoglobin A _{1c} point-of-care testing in general practitioners' offices are associated with better glycemic control in type 2 diabetes. Clinical Chemistry and Laboratory Medicine, 2020, 58, 1349-1356.	2.3	14
154	Effect of zinc on protoporphyrin induced photohaemolysis. Scandinavian Journal of Clinical and Laboratory Investigation, 1980, 40, 185-189.	1.2	13
155	â€~Likelihood-ratio' and â€~odds' applied to monitoring of patients as a supplement to â€~reference chan value' (RCV). Clinical Chemistry and Laboratory Medicine, 2008, 46, 157-64.	ge _{2.3}	13
156	Usefulness of factor V Leiden mutation testing in clinical practice. European Journal of Human Genetics, 2010, 18, 862-866.	2.8	13
157	Can Changes in Troponin Results Be Useful in Diagnosing Myocardial Infarction?. Clinical Chemistry, 2010, 56, 1047-1049.	3.2	13
158	Are general practitioners characteristics associated with the quality of type 2 diabetes care in general practice? Results from the Norwegian ROSA4 study from 2014. Scandinavian Journal of Primary Health Care, 2018, 36, 170-179.	1.5	13
159	Exploring the relationship between coronary heart disease and type 2 diabetes: a cross-sectional study of secondary prevention among diabetes patients. BJGP Open, 2019, 3, bjgpopen18X101636.	1.8	13
160	Critical review and meta-analysis of biological variation estimates for tumor markers. Clinical Chemistry and Laboratory Medicine, 2022, 60, 494-504.	2.3	13
161	Light-induced redistribution and photobleaching of protoporphyrin in erythrocytes from patients with erythropoietic protoporphyria: An explanation of the rapid fading of fluorocytes. Journal of Photochemistry and Photobiology B: Biology, 1988, 2, 33-41.	3.8	12
162	Diabetes care in Norwegian pharmacies: a descriptive study. International Journal of Clinical Pharmacy, 2008, 30, 191-198.	1.4	12

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163	Do new concepts for deriving permissible limits for analytical imprecision and bias have any advantages over existing consensus?. Clinical Chemistry and Laboratory Medicine, 2011, 49, 637-640.	2.3	12
164	Establishing a network of specialist Porphyria centres - effects on diagnostic activities and services. Orphanet Journal of Rare Diseases, 2012, 7, 93.	2.7	12
165	Performance of 10 Systems for Self-Monitoring of Blood Glucose by Trained Healthcare Professionals and in the Hands of the Users. Clinical Chemistry, 2015, 61, 772-774.	3.2	12
166	Biological variation: Evaluation of methods for constructing confidence intervals for estimates of within-person biological variation for different distributions of the within-person effect. Clinica Chimica Acta, 2017, 468, 166-173.	1.1	12
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