Boe Sandahl Sorensen

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Cell-Free DNA and Clinical Characteristics in Patients with Small Intestinal or Pancreatic Neuroendocrine Tumors. Neuroendocrinology, 2022, 112, 43-50.	1.2	7
2	Alectinib-Induced Pleural and Pericardial Effusions in ALK-Positive NSCLC. Case Reports in Oncology, 2022, 14, 1323-1327.	0.3	4
3	The Diagnostic Value of Circulating Cell-Free HPV DNA in Plasma from Cervical Cancer Patients. Cells, 2022, 11, 2170.	1.8	10
4	Increased Soluble PD-1 Predicts Response to Nivolumab plus Ipilimumab in Melanoma. Cancers, 2022, 14, 3342.	1.7	9
5	Combining tissue and circulating tumor DNA increases the detection rate of a CTNNB1 mutation in hepatocellular carcinoma. BMC Cancer, 2021, 21, 376.	1.1	7
6	Co-occurring MET Amplification Predicts Inferior Clinical Response to First-Line Erlotinib in Advanced Stage EGFR-Mutated NSCLC Patients. Clinical Lung Cancer, 2021, 22, e870-e877.	1.1	6
7	STAT3 is over-activated within CD163pos bone marrow macrophages in both Multiple Myeloma and the benign pre-condition MGUS. Cancer Immunology, Immunotherapy, 2021, , 1.	2.0	7
8	<i>EGFR</i> transcription in nonâ€smallâ€cell lung cancer tumours can be revealed in ctDNA by cellâ€free chromatin immunoprecipitation (cfChIP). Molecular Oncology, 2021, 15, 2868-2876.	2.1	7
9	cGAS-STING pathway expression as a prognostic tool in NSCLC. Translational Lung Cancer Research, 2021, 10, 340-354.	1.3	18
10	Clearing of circulating tumour DNA predicts clinical response to first line tyrosine kinase inhibitors in advanced epidermal growth factor receptor mutated non-small cell lung cancer. Lung Cancer, 2020, 141, 37-43.	0.9	24
11	Neurofilament Light Chain as A Biomarker for Brain Metastases. Cancers, 2020, 12, 2852.	1.7	20
12	Epithelial-to-mesenchymal transition is a resistance mechanism to sequential MET-TKI treatment of MET-amplified EGFR-TKI resistant non-small cell lung cancer cells. Translational Lung Cancer Research, 2020, 9, 1904-1914.	1.3	13
13	Cell-free Chromatin Immunoprecipitation (cfChIP) from blood plasma can determine gene-expression in tumors from non-small-cell lung cancer patients. Lung Cancer, 2020, 147, 244-251.	0.9	12
14	TERT promoter mutated circulating tumor DNA as a biomarker for prognosis in hepatocellular carcinoma. Scandinavian Journal of Gastroenterology, 2020, 55, 1433-1440.	0.6	28
15	Inflammatory Cytokines and ctDNA Are Biomarkers for Progression in Advanced-Stage Melanoma Patients Receiving Checkpoint Inhibitors. Cancers, 2020, 12, 1414.	1.7	15
16	Clearing of circulating tumour DNA predicts clinical response to osimertinib in EGFR mutated lung cancer patients. Lung Cancer, 2020, 143, 67-72.	0.9	17
17	Genomic Profiling of Circulating Tumor DNA Predicts Outcome and Demonstrates Tumor Evolution in ALK-Positive Non-Small Cell Lung Cancer Patients. Cancers, 2020, 12, 947.	1.7	20
18	Correlation between early dynamics in circulating tumour DNA and outcome from FOLFIRI treatment in metastatic colorectal cancer. Scientific Reports, 2019, 9, 11542.	1.6	25

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19	Circulating miR-30b and miR-30c predict erlotinib response in EGFR-mutated non-small cell lung cancer patients. Lung Cancer, 2019, 135, 92-96.	0.9	22
20	Intraâ€individual variation of circulating tumour DNA in lung cancer patients. Molecular Oncology, 2019, 13, 2098-2106.	2.1	14
21	Day-to-day and within-day biological variation of cell-free DNA. EBioMedicine, 2019, 49, 284-290.	2.7	49
22	EGFR Gene Polymorphism Predicts Improved Outcome in Patients With EGFR Mutation-positive Non–small cell Lung Cancer Treated With Erlotinib. Clinical Lung Cancer, 2019, 20, 161-166.e1.	1.1	13
23	Up-Regulated FGFR1 Expression as a Mediator of Intrinsic TKI Resistance in EGFR-Mutated NSCLC. Translational Oncology, 2019, 12, 432-440.	1.7	20
24	The prognostic role of inflammation-scores on overall survival in lung cancer patients. Acta OncolÃ ³ gica, 2019, 58, 371-376.	0.8	15
25	The T790M resistance mutation in EGFR is only found in cfDNA from erlotinib-treated NSCLC patients that harbored an activating EGFR mutation before treatment. BMC Cancer, 2018, 18, 191.	1.1	14
26	Detection of EGFR Variants in Plasma. Journal of Molecular Diagnostics, 2018, 20, 483-494.	1.2	37
27	A method for treatment monitoring using circulating tumour DNA in cancer patients without targetable mutations. Oncotarget, 2018, 9, 31066-31076.	0.8	18
28	Total cell-free DNA, carcinoembryonic antigen, and C-reactive protein for assessment of prognosis in patients with metastatic colorectal cancer. Tumor Biology, 2018, 40, 101042831881120.	0.8	10
29	Measuring KRAS Mutations in Circulating Tumor DNA by Droplet Digital PCR and Next-Generation Sequencing. Translational Oncology, 2018, 11, 1220-1224.	1.7	63
30	Cell-free DNA levels and correlation to stage and outcome following treatment of locally advanced rectal cancer. Tumor Biology, 2017, 39, 101042831773097.	0.8	18
31	Correlation between circulating mutant DNA and metabolic tumour burden in advanced non-small cell lung cancer patients. British Journal of Cancer, 2017, 117, 704-709.	2.9	45
32	Soluble HER3 predicts survival in bladder cancer patients. Oncology Letters, 2017, 15, 1783-1788.	0.8	5
33	Increased PD-L1 expression in erlotinib-resistant NSCLC cells with <i>MET</i> gene amplification is reversed upon MET-TKI treatment. Oncotarget, 2017, 8, 68221-68229.	0.8	31
34	IGF1R depletion facilitates <i>MET</i> -amplification as mechanism of acquired resistance to erlotinib in HCC827 NSCLC cells. Oncotarget, 2017, 8, 33300-33315.	0.8	23
35	The role of epithelial to mesenchymal transition in resistance to epidermal growth factor receptor tyrosine kinase inhibitors in non-small cell lung cancer. Translational Lung Cancer Research, 2016, 5, 172-182.	1.3	80
36	Dasatinib and Doxorubicin Treatment of Sarcoma Initiating Cells: A Possible New Treatment Strategy. Stem Cells International, 2016, 2016, 1-8.	1.2	12

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37	Early Change in FDG-PET Signal and Plasma Cell-Free DNA Level Predicts Erlotinib Response in EGFR Wild-Type NSCLC Patients. Translational Oncology, 2016, 9, 505-511.	1.7	13
38	Ultra-micro samples can be used for mRNA quantification of lung cancer biomarkers. Scandinavian Journal of Clinical and Laboratory Investigation, 2016, 76, 243-248.	0.6	2
39	Metabolic tumor burden as marker of outcome in advanced EGFR wild-type NSCLC patients treated with erlotinib. Lung Cancer, 2016, 94, 81-87.	0.9	34
40	Increase in soluble PD-1 is associated with prolonged survival in patients with advanced EGFR -mutated non-small cell lung cancer treated with erlotinib. Lung Cancer, 2016, 100, 77-84.	0.9	97
41	Exosomal Proteins as Diagnostic Biomarkers inÂLungÂCancer. Journal of Thoracic Oncology, 2016, 11, 1701-1710.	0.5	213
42	Gene Expression of the EGF System—a Prognostic Model in Non–Small Cell Lung Cancer Patients Without Activating EGFR Mutations. Translational Oncology, 2016, 9, 306-312.	1.7	7
43	Regulatory dissection of the CBX5 and hnRNPA1 bi-directional promoter in human breast cancer cells reveals novel transcript variants differentially associated with HP11± down-regulation in metastatic cells. BMC Cancer, 2016, 16, 32.	1.1	13
44	Exosomal proteins as potential diagnostic markers in advanced nonâ€small cell lung carcinoma. Journal of Extracellular Vesicles, 2015, 4, 26659.	5.5	242
45	Co-expression of HER3 and MUC1 is associated with a favourable prognosis in patients with bladder cancer. BJU International, 2015, 115, 163-165.	1.3	14
46	Genetic polymorphism in the epidermal growth factor receptor gene predicts outcome in advanced non-small cell lung cancer patients treated with erlotinib. Lung Cancer, 2015, 90, 314-320.	0.9	13
47	Abstract 5064: EGFR and HER3 are important in the interaction between lung cancer cells and fibroblasts. , 2015, , .		Ο
48	Expression of the EGF Family in Gastric Cancer: Downregulation of HER4 and Its Activating Ligand NRG4. PLoS ONE, 2014, 9, e94606.	1.1	39
49	Monitoring of epidermal growth factor receptor tyrosine kinase inhibitorâ€sensitizing and resistance mutations in the plasma DNA of patients with advanced non–small cell lung cancer during treatment with erlotinib. Cancer, 2014, 120, 3896-3901.	2.0	180
50	Expression of the epidermal growth factor system in human middle ear cholesteatoma. Acta Oto-Laryngologica, 2014, 134, 124-134.	0.3	6
51	HER4 and its cytoplasmic isoforms are associated with progression-free survival of malignant melanoma. Melanoma Research, 2014, 24, 88-91.	0.6	15
52	EGFR mutation frequency and effectiveness of erlotinib: A prospective observational study in Danish patients with non-small cell lung cancer. Lung Cancer, 2014, 83, 224-230.	0.9	41
53	EGFR CA repeat polymorphism predict clinical outcome in EGFR mutation positive NSCLC patients treated with erlotinib. Lung Cancer, 2014, 85, 435-441.	0.9	11
54	Detection of EGFR mutations in plasma and biopsies from non-small cell lung cancer patients by allele-specific PCR assays. BMC Cancer, 2014, 14, 294.	1.1	135

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55	The HER4 isoform JM-a/CYT2 relates to improved survival in bladder cancer patients but only if the estrogen receptor \hat{I}_{\pm} is not expressed. Scandinavian Journal of Clinical and Laboratory Investigation, 2013, 73, 503-513.	0.6	5
56	Expression of PIK3CA, PTEN mRNA and PIK3CA mutations in primary breast cancer: association with lymph node metastases. SpringerPlus, 2013, 2, 464.	1.2	12
57	Estrogen receptor \hat{I}_{\pm} is the major driving factor for growth in tamoxifen-resistant breast cancer and supported by HER/ERK signaling. Breast Cancer Research and Treatment, 2013, 139, 71-80.	1.1	59
58	A Single Rainbow Trout Cobalamin-binding Protein Stands in for Three Human Binders. Journal of Biological Chemistry, 2012, 287, 33917-33925.	1.6	12
59	Hypoxia Changes the Expression of the Epidermal Growth Factor (EGF) System in Human Hearts and Cultured Cardiomyocytes. PLoS ONE, 2012, 7, e40243.	1.1	28
60	Erlotinib Accumulation in Brain Metastases from Non-small Cell Lung Cancer: Visualization by Positron Emission Tomography in a Patient Harboring a Mutation in the Epidermal Growth Factor Receptor. Journal of Thoracic Oncology, 2011, 6, 1287-1289.	0.5	124
61	Complete Pathologic Response in Lung Tumors in Two Patients with Metastatic Non-small Cell Lung Cancer Treated with Erlotinib. Journal of Thoracic Oncology, 2011, 6, 1946-1949.	0.5	12
62	Calcium-induced apoptosis is delayed by HER1 receptor signalling through the Akt and PLCÎ ³ pathways in bladder cancer cells. Scandinavian Journal of Clinical and Laboratory Investigation, 2011, 71, 45-51.	0.6	2
63	Identifying responders to trastuzumab therapy in breast cancer. Future Oncology, 2011, 7, 767-773.	1.1	5
64	Mouse Transcobalamin Has Features Resembling both Human Transcobalamin and Haptocorrin. PLoS ONE, 2011, 6, e20638.	1.1	34
65	Transcobalamin deficiency caused by compound heterozygosity for two novel mutations in the <i>TCN2</i> gene: a study of two affected siblings, their brother, and their parents. Journal of Inherited Metabolic Disease, 2010, 33, 269-274.	1.7	11
66	Circulating HER2 DNA after trastuzumab treatment predicts survival and response in breast cancer. Anticancer Research, 2010, 30, 2463-8.	0.5	16
67	Positron Emission Tomography (PET) Imaging with [11C]-Labeled Erlotinib: A Micro-PET Study on Mice with Lung Tumor Xenografts. Cancer Research, 2009, 69, 873-878.	0.4	164
68	Expression of the Epidermal Growth Factor System in Eutopic Endometrium from Women with Endometriosis Differs from That in Endometrium from Healthy Women. Gynecologic and Obstetric Investigation, 2009, 67, 118-126.	0.7	16
69	Activation of ErbB3, EGFR and Erk is essential for growth of human breast cancer cell lines with acquired resistance to fulvestrant. Breast Cancer Research and Treatment, 2009, 114, 263-75.	1.1	129
70	Quantitative real-time RT-PCR in sentinel lymph nodes from melanoma patients Apmis, 2008, 116, 199-205.	0.9	7
71	A comparison among HER2, <i>TP53</i> , PAI-1, angiogenesis, and proliferation activity as prognostic variables in tumours from 408 patients diagnosed with early breast cancer. Acta Oncológica, 2008, 47, 618-632.	0.8	24
72	Insulin induces a transcriptional activation of epiregulin, HB-EGF and amphiregulin, by a PI3K-dependent mechanism: Identification of a specific insulin-responsive promoter element. Biochemical and Biophysical Research Communications, 2007, 354, 885-891.	1.0	17

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73	Inhibition of the epidermal growth factor receptor in bladder cancer cells treated with the DNA-damaging drug etoposide markedly increases apoptosis. BJU International, 2007, 99, 196-201.	1.3	5
74	Expression of the epidermal growth factor system in endometrioid endometrial cancer. Gynecologic Oncology, 2007, 104, 158-167.	0.6	49
75	Serum YKL-40 Predicts Relapse-Free and Overall Survival in Patients With American Joint Committee on Cancer Stage I and II Melanoma. Journal of Clinical Oncology, 2006, 24, 798-804.	0.8	71
76	Insulin-induced proliferation of bladder cancer cells is mediated through activation of the epidermal growth factor system. FEBS Journal, 2006, 273, 5479-5489.	2.2	16
77	Circulating Tyrosinase and MART-1 mRNA does not Independently Predict Relapse or Survival in Patients with AJCC Stage l–II Melanoma. Journal of Investigative Dermatology, 2006, 126, 849-854.	0.3	8
78	The chemotherapeutic agent VP16 increases the stability of HB-EGF mRNA by a mechanism involving the 3′-UTR. Experimental Cell Research, 2006, 312, 3651-3658.	1.2	8
79	Tyrosinase messenger RNA in peripheral blood is related to poor survival in patients with metastatic melanoma following interleukin-2-based immunotherapy. Melanoma Research, 2005, 15, 409-416.	0.6	30
80	Increase in amphiregulin and epiregulin in prostate cancer xenograft after androgen deprivation—impact of specific HER1 inhibition. Prostate, 2005, 64, 1-8.	1.2	20
81	Pathologic Assessment of Melanoma Sentinel Nodes: A Role for Molecular Analysis Using Quantitative Real-Time Reverse Transcription-PCR for MART-1 and Tyrosinase Messenger RNA. Clinical Cancer Research, 2005, 11, 1425-1433.	3.2	23
82	The DNA damaging agent VP16 induces the expression of a subset of ligands from the EGF system in bladder cancer cells, whereas none of the four EGF receptors are induced. Molecular and Cellular Biochemistry, 2004, 260, 129-135.	1.4	11
83	S100β protein in peripheral blood may predict progressive disease during interleukin-2 based immunotherapy in patients with metastatic melanoma. Melanoma Research, 2004, 14, 211-215.	0.6	8
84	The Influence of Immunohistochemistry on mRNA Recovery from Microdissected Frozen and Formalin-Fixed, Paraffin-Embedded Sections. Diagnostic Molecular Pathology, 2004, 13, 224-233.	2.1	27
85	ErbB1 and prostate cancer: ErbB1 activity is essential for androgen-induced proliferation and protection from the apoptotic effects of LY294002. Prostate, 2003, 56, 142-149.	1.2	39
86	Transcellular Transport of Vitamin B12in LLC-PK1 Renal Proximal Tubule Cells. Journal of the American Society of Nephrology: JASN, 2001, 12, 1099-1106.	3.0	23
87	Simultaneous Quantitation of Several mRNA Species by Calibrated Reverse Transcription Polymerase Chain Reaction and Capillary Electrophoresis: Analysis of the Epidermal Growth Factor Receptor and its Activating Ligands EGF, TGF-1±, and HB-EGF in Rat Liver. Laboratory Investigation, 2000, 80, 983-986.	1.7	6
88	Quantitation of the mRNA expression of the epidermal growth factor system: Selective induction of heparin-binding epidermal growth factor–like growth factor and amphiregulin expression by growth factor stimulation of prostate stromal cells. Translational Research, 2000, 136, 209-217.	2.4	13
89	Epidermal growth factor and insulin-like growth factor I upregulate the expression of the epidermal growth factor system in rat liver. Journal of Hepatology, 2000, 32, 645-654.	1.8	8
90	Different modes of anthracycline interaction with topoisomerase II. Biochemical Pharmacology, 1993, 45, 2025-2035.	2.0	60

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91	Antagonistic effect of the cardioprotector (+)-1,2-BIS(3,5-dioxopiperazinyl-1-YL)propane(ICRF-187) on dna breaks and cytotoxicity induced by the topoisomerase ii directed drugs daunorubicin and etoposide (VP-16). Biochemical Pharmacology, 1993, 46, 389-393.	2.0	80