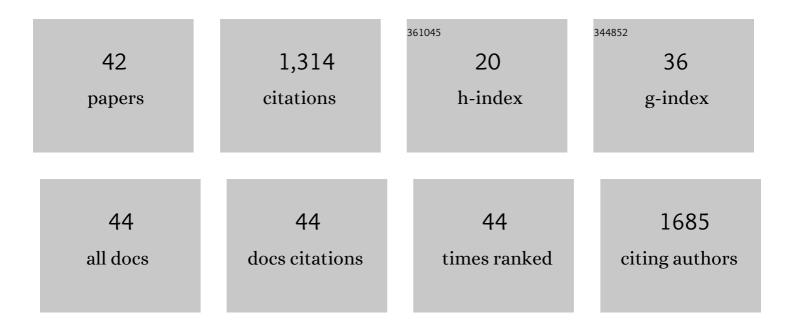
Roberta Noberini

List of Publications by Year in descending order

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#	Article	lF	CITATIONS
1	Mass spectrometryâ€based characterization of histones in clinical samples: applications, progress, and challenges. FEBS Journal, 2022, 289, 1191-1213.	2.2	20
2	Intestinal differentiation involves cleavage of histone H3 N-terminal tails by multiple proteases. Nucleic Acids Research, 2021, 49, 791-804.	6.5	21
3	Anticancer innovative therapy congress: Highlights from the 10th anniversary edition. Cytokine and Growth Factor Reviews, 2021, 59, 1-8.	3.2	4
4	Spatial epi-proteomics enabled by histone post-translational modification analysis from low-abundance clinical samples. Clinical Epigenetics, 2021, 13, 145.	1.8	15
5	LSD1-directed therapy affects glioblastoma tumorigenicity by deregulating the protective ATF4-dependent integrated stress response. Science Translational Medicine, 2021, 13, eabf7036.	5.8	18
6	Enrichment of histones from patient samples for mass spectrometry-based analysis of post-translational modifications. Methods, 2020, 184, 19-28.	1.9	23
7	Label-Free Mass Spectrometry-Based Quantification of Linker Histone H1 Variants in Clinical Samples. International Journal of Molecular Sciences, 2020, 21, 7330.	1.8	8
8	Clinical Application of Mass Spectrometryâ€Based Proteomics in Lung Cancer Early Diagnosis. Proteomics - Clinical Applications, 2020, 14, 1900138.	0.8	14
9	hSWATH: Unlocking SWATH's Full Potential for an Untargeted Histone Perspective. Journal of Proteome Research, 2019, 18, 3840-3849.	1.8	12
10	Epigenetic drug target deconvolution by mass spectrometry–based technologies. Nature Structural and Molecular Biology, 2019, 26, 854-857.	3.6	9
11	Profiling of Epigenetic Features in Clinical Samples Reveals Novel Widespread Changes in Cancer. Cancers, 2019, 11, 723.	1.7	26
12	Mass Spectrometry and Epigenetics. , 2019, , 2251-2268.		0
13	Alternative digestion approaches improve histone modification mapping by mass spectrometry in clinical samples. Proteomics - Clinical Applications, 2019, 13, 1700166.	0.8	11
14	Extensive and systematic rewiring of histone post-translational modifications in cancer model systems. Nucleic Acids Research, 2018, 46, 3817-3832.	6.5	31
15	PAT-H-MS coupled with laser microdissection to study histone post-translational modifications in selected cell populations from pathology samples. Clinical Epigenetics, 2017, 9, 69.	1.8	17
16	Mass Spectrometry and Epigenetics. , 2017, , 1-18.		1
17	Protein kinase A can block EphA2 receptor–mediated cell repulsion by increasing EphA2 S897 phosphorylation. Molecular Biology of the Cell, 2016, 27, 2757-2770.	0.9	59
18	Mass-spectrometry analysis of histone post-translational modifications in pathology tissue using the PAT-H-MS approach. Data in Brief, 2016, 7, 188-194.	0.5	6

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#	Article	IF	CITATIONS
19	The contribution of mass spectrometry-based proteomics to understanding epigenetics. Epigenomics, 2016, 8, 429-445.	1.0	30
20	Recent advances in mass spectrometry analysis of histone post-translational modifications: potential clinical impact of the PAT-H-MS approach. Expert Review of Proteomics, 2016, 13, 245-250.	1.3	3
21	Pathology Tissue-quantitative Mass Spectrometry Analysis to Profile Histone Post-translational Modification Patterns in Patient Samples. Molecular and Cellular Proteomics, 2016, 15, 866-877.	2.5	41
22	Mass Spectrometry for the Identification of Posttranslational Modifications in Histones and Its Application in Clinical Epigenetics. , 2016, , 195-214.		1
23	Design, Synthesis and Bioevaluation of an EphA2 Receptorâ€Based Targeted Delivery System. ChemMedChem, 2014, 9, 1403-1412.	1.6	31
24	Quantitative Chemical Proteomics Identifies Novel Targets of the Anti-cancer Multi-kinase Inhibitor E-3810. Molecular and Cellular Proteomics, 2014, 13, 1495-1509.	2.5	14
25	Design, synthesis and characterization of novel small molecular inhibitors of ephrin-B2 binding to EphB4. Biochemical Pharmacology, 2013, 85, 507-513.	2.0	23
26	Amino Acid Conjugates of Lithocholic Acid As Antagonists of the EphA2 Receptor. Journal of Medicinal Chemistry, 2013, 56, 2936-2947.	2.9	50
27	HTS by NMR of Combinatorial Libraries: A Fragment-Based Approach to Ligand Discovery. Chemistry and Biology, 2013, 20, 19-33.	6.2	72
28	Targeted Delivery of Paclitaxel to EphA2-Expressing Cancer Cells. Clinical Cancer Research, 2013, 19, 128-137.	3.2	53
29	Distinctive binding of three antagonistic peptides to the ephrin-binding pocket of the EphA4 receptor. Biochemical Journal, 2012, 445, 47-56.	1.7	47
30	Profiling Eph receptor expression in cells and tissues. Cell Adhesion and Migration, 2012, 6, 102-156.	1.1	54
31	Inhibition of Eph receptor–ephrin ligand interaction by tea polyphenols. Pharmacological Research, 2012, 66, 363-373.	3.1	18
32	Novel Targeted System To Deliver Chemotherapeutic Drugs to EphA2-Expressing Cancer Cells. Journal of Medicinal Chemistry, 2012, 55, 2427-2436.	2.9	79
33	Targeting Eph receptors with peptides and small molecules: Progress and challenges. Seminars in Cell and Developmental Biology, 2012, 23, 51-57.	2.3	89
34	Structure–Activity Relationships and Mechanism of Action of Eph–ephrin Antagonists: Interaction of Cholanic Acid with the EphA2 Receptor. ChemMedChem, 2012, 7, 1071-1083.	1.6	31
35	A Disalicylic Acidâ€Furanyl Derivative Inhibits Ephrin Binding to a Subset of Eph Receptors. Chemical Biology and Drug Design, 2011, 78, 667-678.	1.5	39
36	Characterization of a novel angiogenic model based on stable, fluorescently labelled endothelial cell lines amenable to scale-up for high content screening. Biology of the Cell, 2011, 103, 467-481.	0.7	15

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37	PECylation Potentiates the Effectiveness of an Antagonistic Peptide That Targets the EphB4 Receptor with Nanomolar Affinity. PLoS ONE, 2011, 6, e28611.	1.1	36
38	Structural Characterization of the EphA4-Ephrin-B2 Complex Reveals New Features Enabling Eph-Ephrin Binding Promiscuity. Journal of Biological Chemistry, 2010, 285, 644-654.	1.6	84
39	Proliferation and Tumor Suppression: Not Mutually Exclusive for Eph Receptors. Cancer Cell, 2009, 16, 452-454.	7.7	19
40	Structural and Binding Study on the Interaction of Small Molecule Antagonists with the EphA4 Receptor. FASEB Journal, 2009, 23, LB297.	0.2	0
41	Small Molecules Can Selectively Inhibit Ephrin Binding to the EphA4 and EphA2 Receptors. Journal of Biological Chemistry, 2008, 283, 29461-29472.	1.6	123
42	Crystal Structure and NMR Binding Reveal That Two Small Molecule Antagonists Target the High Affinity Ephrin-binding Channel of the EphA4 Receptor. Journal of Biological Chemistry, 2008, 283, 29473-29484.	1.6	66