

# Frauke Christ

## List of Publications by Year in descending order

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49  
papers

2,085  
citations

331259

21  
h-index

233125

45  
g-index

54  
all docs

54  
docs citations

54  
times ranked

2033  
citing authors

#	ARTICLE	IF	CITATIONS
1	Rational design of small-molecule inhibitors of the LEDGF/p75-integrase interaction and HIV replication. <i>Nature Chemical Biology</i> , 2010, 6, 442-448.	3.9	428
2	Transportin-SR2 Imports HIV into the Nucleus. <i>Current Biology</i> , 2008, 18, 1192-1202.	1.8	231
3	The Interaction of LEDGF/p75 with Integrase Is Lentivirus-specific and Promotes DNA Binding. <i>Journal of Biological Chemistry</i> , 2005, 280, 17841-17847.	1.6	182
4	Small-Molecule Inhibitors of the LEDGF/p75 Binding Site of Integrase Block HIV Replication and Modulate Integrase Multimerization. <i>Antimicrobial Agents and Chemotherapy</i> , 2012, 56, 4365-4374.	1.4	158
5	The LEDGF/p75 integrase interaction, a novel target for anti-HIV therapy. <i>Virology</i> , 2013, 435, 102-109.	1.1	96
6	LEDGIN-mediated Inhibition of Integraseâ€“LEDGF/p75 Interaction Reduces Reactivation of Residual Latent HIV. <i>EBioMedicine</i> , 2016, 8, 248-264.	2.7	90
7	Host factors for retroviral integration site selection. <i>Trends in Biochemical Sciences</i> , 2015, 40, 108-116.	3.7	83
8	Multiple cellular proteins interact with LEDGF/p75 through a conserved unstructured consensus motif. <i>Nature Communications</i> , 2015, 6, 7968.	5.8	53
9	Interaction of the HIV-1 Intasome with Transportin 3 Protein (TNPO3 or TRN-SR2). <i>Journal of Biological Chemistry</i> , 2012, 287, 34044-34058.	1.6	52
10	4-Substituted 2-Hydroxyisoquinoline-1,3(2 <i>H</i> )-diones as a Novel Class of HIV-1 Integrase Inhibitors. <i>ACS Medicinal Chemistry Letters</i> , 2013, 4, 606-611.	1.3	52
11	Interplay between HIV Entry and Transportin-SR2 Dependency. <i>Retrovirology</i> , 2011, 8, 7.	0.9	51
12	Lentiviral nuclear import: a complex interplay between virus and host. <i>BioEssays</i> , 2007, 29, 441-451.	1.2	42
13	Kuwanonâ€“ as a New Allosteric HIVâ€“1 Integrase Inhibitor: Molecular Modeling and Biological Evaluation. <i>ChemBioChem</i> , 2015, 16, 2507-2512.	1.3	39
14	Capsid-Labelled HIV To Investigate the Role of Capsid during Nuclear Import and Integration. <i>Journal of Virology</i> , 2020, 94, .	1.5	34
15	LEDGF/p75 is dispensable for hematopoiesis but essential for MLL-rearranged leukemogenesis. <i>Blood</i> , 2018, 131, blood-2017-05-786962.	0.6	32
16	Identification of Novel 3-Hydroxy-pyran-4-One Derivatives as Potent HIV-1 Integrase Inhibitors Using in silico Structure-Based Combinatorial Library Design Approach. <i>Frontiers in Chemistry</i> , 2019, 7, 574.	1.8	32
17	The HIV-1 Integrase Mutant R263A/K264A Is 2-fold Defective for TRN-SR2 Binding and Viral Nuclear Import. <i>Journal of Biological Chemistry</i> , 2014, 289, 25351-25361.	1.6	28
18	Dynamic Oligomerization of Integrase Orchestrates HIV Nuclear Entry. <i>Scientific Reports</i> , 2016, 6, 36485.	1.6	28

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19	Affinity switching of the LEDGF/p75 IBD interactome is governed by kinase-dependent phosphorylation. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2018, 115, E7053-E7062.	3.3	27
20	Insight in HIV Integration Site Selection Provides a Block-and-Lock Strategy for a Functional Cure of HIV Infection. <i>Viruses</i> , 2019, 11, 12.	1.5	26
21	Phenotyping of Rare CFTR Mutations Reveals Distinct Trafficking and Functional Defects. <i>Cells</i> , 2020, 9, 754.	1.8	23
22	The mutation of Transportin 3 gene that causes limb girdle muscular dystrophy 1F induces protection against HIV-1 infection. <i>PLoS Pathogens</i> , 2019, 15, e1007958.	2.1	22
23	Impact of LEDGIN treatment during virus production on residual HIV-1 transcription. <i>Retrovirology</i> , 2019, 16, 8.	0.9	22
24	Synthesis, Molecular Modelling and Biological Studies of 3-hydroxypyran- 4-one and 3-hydroxy-pyridine-4-one Derivatives as HIV-1 Integrase Inhibitors. <i>Medicinal Chemistry</i> , 2019, 15, 755-770.	0.7	22
25	Protein-protein and protein-chromatin interactions of LEDGF/p75 as novel drug targets. <i>Drug Discovery Today: Technologies</i> , 2017, 24, 25-31.	4.0	21
26	Diketoacid chelating ligands as dual inhibitors of HIV-1 integration process. <i>European Journal of Medicinal Chemistry</i> , 2014, 78, 425-430.	2.6	17
27	A new potential approach to block HIV-1 replication via protein-protein interaction and strand-transfer inhibition. <i>Bioorganic and Medicinal Chemistry</i> , 2014, 22, 2269-2279.	1.4	17
28	GS-9822, a Preclinical LEDGIN Candidate, Displays a Block-and-Lock Phenotype in Cell Culture. <i>Antimicrobial Agents and Chemotherapy</i> , 2021, 65, .	1.4	17
29	Design and discovery of 5-hydroxy-6-oxo-1,6-dihydropyrimidine-4-carboxamide inhibitors of HIV-1 integrase. <i>Bioorganic and Medicinal Chemistry</i> , 2014, 22, 5446-5453.	1.4	15
30	Inhibitors of the integrase-transportin-SR2 interaction block HIV nuclear import. <i>Retrovirology</i> , 2018, 15, 5.	0.9	14
31	De novo design of small molecule inhibitors targeting the LEDGF/p75-HIVintegrase interaction. <i>RSC Advances</i> , 2012, 2, 974-984.	1.7	13
32	Structure of transportin SR2, a karyopherin involved in human disease, in complex with Ran. <i>Acta Crystallographica Section F, Structural Biology Communications</i> , 2014, 70, 723-729.	0.4	11
33	2-hydroxyisoquinoline-1,3(2 H ,4 H )-diones (HIDs) as human immunodeficiency virus type 1 integrase inhibitors: Influence of the alkylcarboxamide substitution of position 4. <i>European Journal of Medicinal Chemistry</i> , 2016, 117, 256-268.	2.6	11
34	N-terminal half of transportin SR2 interacts with HIV integrase. <i>Journal of Biological Chemistry</i> , 2017, 292, 9699-9710.	1.6	11
35	Synthesis of new pyridazine derivatives as potential anti-HIV agents. <i>Journal of Heterocyclic Chemistry</i> , 2009, 46, 1420-1424.	1.4	9
36	HIV-1 integrase inhibition: looking at cofactor interactions. <i>Future Medicinal Chemistry</i> , 2015, 7, 2407-2410.	1.1	9

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37	Towards a Functional Cure of HIV-1: Insight Into the Chromatin Landscape of the Provirus. <i>Frontiers in Microbiology</i> , 2021, 12, 636642.	1.5	9
38	Lessons Learned: HIV Points the Way Towards Precision Treatment of Mixed-Lineage Leukemia. <i>Trends in Pharmacological Sciences</i> , 2016, 37, 660-671.	4.0	8
39	LEDGF/p75-mediated chemoresistance of mixed-lineage leukemia involves cell survival pathways and super enhancer activators. <i>Cancer Gene Therapy</i> , 2022, 29, 133-140.	2.2	7
40	LEDGINs, Inhibitors of the Interaction Between HIV-1 Integrase and LEDGF/p75, Are Potent Antivirals with a Potential to Cure HIV Infection. <i>Advances in Experimental Medicine and Biology</i> , 2021, 1322, 97-114.	0.8	6
41	Role of Transportin-SR2 in HIV-1 Nuclear Import. <i>Viruses</i> , 2021, 13, 829.	1.5	6
42	Unlike its Paralog LEDGF/p75, HRP-2 Is Dispensable for MLL-R Leukemogenesis but Important for Leukemic Cell Survival. <i>Cells</i> , 2021, 10, 192.	1.8	5
43	Validation of host factors of HIV integration as novel drug targets for anti-HIV therapy. <i>MedChemComm</i> , 2014, 5, 314-320.	3.5	4
44	Optimization of rhodanine scaffold for the development of protein-protein interaction inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2015, 23, 3208-3214.	1.4	4
45	Molecular Mechanism of LEDGF/p75 Dimerization. <i>Structure</i> , 2020, 28, 1288-1299.e7.	1.6	4
46	LEDGF/p75 and transportin-SR2 are cellular cofactors of HIV integrase and novel targets for antiviral therapy. <i>HIV Therapy</i> , 2009, 3, 171-188.	0.6	4
47	CRISPR/Cas9-Induced Mutagenesis Corroborates the Role of Transportin-SR2 in HIV-1 Nuclear Import. <i>Microbiology Spectrum</i> , 2021, 9, e0133621.	1.2	3
48	Insight into HIV-2 latency may disclose strategies for a cure for HIV-1 infection. <i>Journal of Virus Eradication</i> , 2017, 3, 7-14.	0.3	3
49	Design, synthesis, in silico studies, and antiproliferative evaluations of novel indolin-2-one derivatives containing 3-hydroxy-4-pyridinone fragment. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2022, 70, 128784.	1.0	3