# Stefan Phlmann

# List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

27,628 65 165 227 h-index g-index citations papers 259 7.73 35,312 9.7 L-index ext. citations avg, IF ext. papers

#	Paper	IF	Citations
227	The MEK1/2-inhibitor ATR-002 efficiently blocks SARS-CoV-2 propagation and alleviates pro-inflammatory cytokine/chemokine responses <i>Cellular and Molecular Life Sciences</i> , <b>2022</b> , 79, 65	10.3	3
226	No evidence for increased cell entry or antibody evasion by Delta sublineage AY.4.2 <i>Cellular and Molecular Immunology</i> , <b>2022</b> ,	15.4	2
225	Rapid SARS-CoV-2 Adaptation to Available Cellular Proteases <i>Journal of Virology</i> , <b>2022</b> , jvi0218621	6.6	2
224	MCMV-based vaccine vectors expressing full-length viral proteins provide long-term humoral immune protection upon a single-shot vaccination <i>Cellular and Molecular Immunology</i> , <b>2022</b> ,	15.4	1
223	Alternatives to animal models and their application in the discovery of species susceptibility to SARS-CoV-2 and other respiratory infectious pathogens: A review <i>Veterinary Pathology</i> , <b>2022</b> , 300985	8 <del>2</del> 18107	73678
222	Augmented Neutralization of SARS-CoV-2 Omicron Variant by Boost Vaccination and Monoclonal Antibodies <i>European Journal of Immunology</i> , <b>2022</b> ,	6.1	1
221	Functional analysis of polymorphisms at the S1/S2 site of SARS-CoV-2 spike protein <i>PLoS ONE</i> , <b>2022</b> , 17, e0265453	3.7	2
220	Omicron: Master of immune evasion maintains robust ACE2 binding <i>Signal Transduction and Targeted Therapy</i> , <b>2022</b> , 7, 118	21	1
219	SARS-CoV-2 variants C.1.2 and B.1.621 (Mu) partially evade neutralization by antibodies elicited upon infection or vaccination <i>Cell Reports</i> , <b>2022</b> , 110754	10.6	0
218	Comparable neutralisation evasion of SARS-CoV-2 omicron subvariants BA.1, BA.2, and BA.3 <i>Lancet Infectious Diseases, The</i> , <b>2022</b> ,	25.5	7
217	Evidence for an ACE2-Independent Entry Pathway That Can Protect from Neutralization by an Antibody Used for COVID-19 Therapy <i>MBio</i> , <b>2022</b> , e0036422	7.8	O
216	Understanding Omicron: Transmissibility, immune evasion and antiviral intervention <i>Clinical and Translational Medicine</i> , <b>2022</b> , 12, e839	5.7	0
215	Heterologous ChAdOx1 nCoV-19 and BNT162b2 prime-boost vaccination elicits potent neutralizing antibody responses and T cell reactivity against prevalent SARS-CoV-2 variants <i>EBioMedicine</i> , <b>2021</b> , 75, 103761	8.8	24
214	The Omicron variant is highly resistant against antibody-mediated neutralization: Implications for control of the COVID-19 pandemic <i>Cell</i> , <b>2021</b> ,	56.2	156
213	Novel SARS-CoV-2 receptors: ASGR1 and KREMEN1 <i>Cell Research</i> , <b>2021</b> ,	24.7	8
212	Activation of Sphingomyelinase-Ceramide-Pathway in COVID-19 Purposes Its Inhibition for Therapeutic Strategies <i>Frontiers in Immunology</i> , <b>2021</b> , 12, 784989	8.4	2
211	Spike residue 403 affects binding of coronavirus spikes to human ACE2. <i>Nature Communications</i> , <b>2021</b> , 12, 6855	17.4	3

# (2021-2021)

210	Protective mucosal immunity against SARS-CoV-2 after heterologous systemic prime-mucosal boost immunization. <i>Nature Communications</i> , <b>2021</b> , 12, 6871	17.4	22
209	Improved cellular and humoral immunity upon a second BNT162b2 and mRNA-1273 boost in prime-boost vaccination no/low responders with end-stage renal disease. <i>Kidney International</i> , <b>2021</b> , 100, 1335-1337	9.9	4
208	The spike protein of SARS-CoV-2 variant A.30 is heavily mutated and evades vaccine-induced antibodies with high efficiency. <i>Cellular and Molecular Immunology</i> , <b>2021</b> , 18, 2673-2675	15.4	12
207	Evidence that two instead of one defective interfering RNA in influenza A virus-derived defective interfering particles (DIPs) does not enhance antiviral activity. <i>Scientific Reports</i> , <b>2021</b> , 11, 20477	4.9	O
206	A novel class of TMPRSS2 inhibitors potently block SARS-CoV-2 and MERS-CoV viral entry and protect human epithelial lung cells. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , <b>2021</b> , 118,	11.5	8
205	Delta variant (B.1.617.2) sublineages do not show increased neutralization resistance. <i>Cellular and Molecular Immunology</i> , <b>2021</b> , 18, 2557-2559	15.4	12
204	Thiol drugs decrease SARS-CoV-2 lung injury and disrupt SARS-CoV-2 spike complex binding to ACE2 <b>2021</b> ,		11
203	assay to evaluate the efficacy of drugs targeting sphingolipids in preventing SARS-CoV-2 infection of nasal epithelial cells. <i>STAR Protocols</i> , <b>2021</b> , 2, 100356	1.4	3
202	Camostat mesylate inhibits SARS-CoV-2 activation by TMPRSS2-related proteases and its metabolite GBPA exerts antiviral activity. <i>EBioMedicine</i> , <b>2021</b> , 65, 103255	8.8	120
201	Mutation D614G increases SARS-CoV-2 transmission. <i>Signal Transduction and Targeted Therapy</i> , <b>2021</b> , 6, 101	21	13
200	Alpha-1 antitrypsin inhibits TMPRSS2 protease activity and SARS-CoV-2 infection. <i>Nature Communications</i> , <b>2021</b> , 12, 1726	17.4	32
199	The SARS-CoV-2 and other human coronavirus spike proteins are fine-tuned towards temperature and proteases of the human airways. <i>PLoS Pathogens</i> , <b>2021</b> , 17, e1009500	7.6	41
198	The sphingosine kinase 1 activator, K6PC-5, attenuates Ebola virus infection. <i>IScience</i> , <b>2021</b> , 24, 102266	6.1	3
197	SARS-CoV-2 variants B.1.351 and P.1 escape from neutralizing antibodies. <i>Cell</i> , <b>2021</b> , 184, 2384-2393.e1	<b>2</b> 56.2	459
196	SARS-CoV-2 mutations acquired in mink reduce antibody-mediated neutralization. <i>Cell Reports</i> , <b>2021</b> , 35, 109017	10.6	42
195	Cell culture-based production and in vivo characterization of purely clonal defective interfering influenza virus particles. <i>BMC Biology</i> , <b>2021</b> , 19, 91	7.3	8
194	Urinary Levels of SARS-CoV-2 Nucleocapsid Protein Associate With Risk of AKI and COVID-19 Severity: A Single-Center Observational Study. <i>Frontiers in Medicine</i> , <b>2021</b> , 8, 644715	4.9	4
193	A novel class of TMPRSS2 inhibitors potently block SARS-CoV-2 and MERS-CoV viral entry and protect human epithelial lung cells <b>2021</b> ,		3

192	Therapeutic Application of Alpha-1 Antitrypsin in COVID-19. <i>American Journal of Respiratory and Critical Care Medicine</i> , <b>2021</b> , 204, 224-227	10.2	15
191	Humoral and Cellular Immune Responses Against Severe Acute Respiratory Syndrome Coronavirus 2 Variants and Human Coronaviruses After Single BNT162b2 Vaccination. <i>Clinical Infectious Diseases</i> , <b>2021</b> , 73, 2000-2008	11.6	17
190	SARS-CoV-2 neutralizing antibodies: Longevity, breadth, and evasion by emerging viral variants. <i>PLoS Medicine</i> , <b>2021</b> , 18, e1003656	11.6	37
189	SARS-CoV-2 variant B.1.617 is resistant to bamlanivimab and evades antibodies induced by infection and vaccination. <i>Cell Reports</i> , <b>2021</b> , 36, 109415	10.6	131
188	Immune responses against SARS-CoV-2 variants after heterologous and homologous ChAdOx1 nCoV-19/BNT162b2 vaccination. <i>Nature Medicine</i> , <b>2021</b> , 27, 1525-1529	50.5	141
187	Molecular mechanism of inhibiting the SARS-CoV-2 cell entry facilitator TMPRSS2 with camostat and nafamostat <i>Chemical Science</i> , <b>2021</b> , 12, 983-992	9.4	27
186	Low serum neutralizing anti-SARS-CoV-2 S antibody levels in mildly affected COVID-19 convalescent patients revealed by two different detection methods. <i>Cellular and Molecular Immunology</i> , <b>2021</b> , 18, 936-944	15.4	62
185	Inhibition of acid sphingomyelinase by ambroxol prevents SARS-CoV-2 entry into epithelial cells. Journal of Biological Chemistry, <b>2021</b> , 296, 100701	5.4	31
184	Natural cystatin C fragments inhibit GPR15-mediated HIV and SIV infection without interfering with GPR15L signaling. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , <b>2021</b> , 118,	11.5	2
183	Synergistic inhibition of SARS-CoV-2 cell entry by otamixaban and covalent protease inhibitors: pre-clinical assessment of pharmacological and molecular properties. <i>Chemical Science</i> , <b>2021</b> , 12, 1260	0-92 <del>1</del> 60:	) <sup>2</sup>
183	Synergistic inhibition of SARS-CoV-2 cell entry by otamixaban and covalent protease inhibitors: pre-clinical assessment of pharmacological and molecular properties. <i>Chemical Science</i> , <b>2021</b> , 12, 1260.  Neutralization of the SARS-CoV-2 Delta variant after heterologous and homologous BNT162b2 or ChAdOx1 nCoV-19 vaccination. <i>Cellular and Molecular Immunology</i> , <b>2021</b> , 18, 2455-2456	0-92 <del>5</del> 09 15.4	9 <sup>2</sup>
	pre-clinical assessment of pharmacological and molecular properties. <i>Chemical Science</i> , <b>2021</b> , 12, 1260  Neutralization of the SARS-CoV-2 Delta variant after heterologous and homologous BNT162b2 or		
182	Pre-clinical assessment of pharmacological and molecular properties. <i>Chemical Science</i> , <b>2021</b> , 12, 1260.  Neutralization of the SARS-CoV-2 Delta variant after heterologous and homologous BNT162b2 or ChAdOx1 nCoV-19 vaccination. <i>Cellular and Molecular Immunology</i> , <b>2021</b> , 18, 2455-2456.  Functional comparison of MERS-coronavirus lineages reveals increased replicative fitness of the	15.4	20
182	Pre-clinical assessment of pharmacological and molecular properties. <i>Chemical Science</i> , <b>2021</b> , 12, 1260.  Neutralization of the SARS-CoV-2 Delta variant after heterologous and homologous BNT162b2 or ChAdOx1 nCoV-19 vaccination. <i>Cellular and Molecular Immunology</i> , <b>2021</b> , 18, 2455-2456.  Functional comparison of MERS-coronavirus lineages reveals increased replicative fitness of the recombinant lineage 5. <i>Nature Communications</i> , <b>2021</b> , 12, 5324.  A pair of noncompeting neutralizing human monoclonal antibodies protecting from disease in a	15.4 17.4	20 0
182 181 180	Pre-clinical assessment of pharmacological and molecular properties. <i>Chemical Science</i> , <b>2021</b> , 12, 1260.  Neutralization of the SARS-CoV-2 Delta variant after heterologous and homologous BNT162b2 or ChAdOx1 nCoV-19 vaccination. <i>Cellular and Molecular Immunology</i> , <b>2021</b> , 18, 2455-2456.  Functional comparison of MERS-coronavirus lineages reveals increased replicative fitness of the recombinant lineage 5. <i>Nature Communications</i> , <b>2021</b> , 12, 5324.  A pair of noncompeting neutralizing human monoclonal antibodies protecting from disease in a SARS-CoV-2 infection model. <i>European Journal of Immunology</i> , <b>2021</b> ,  B.1.617.2 enters and fuses lung cells with increased efficiency and evades antibodies induced by	15.4 17.4 6.1	20 0
182 181 180	Pre-clinical assessment of pharmacological and molecular properties. <i>Chemical Science</i> , <b>2021</b> , 12, 1260.  Neutralization of the SARS-CoV-2 Delta variant after heterologous and homologous BNT162b2 or ChAdOx1 nCoV-19 vaccination. <i>Cellular and Molecular Immunology</i> , <b>2021</b> , 18, 2455-2456.  Functional comparison of MERS-coronavirus lineages reveals increased replicative fitness of the recombinant lineage 5. <i>Nature Communications</i> , <b>2021</b> , 12, 5324.  A pair of noncompeting neutralizing human monoclonal antibodies protecting from disease in a SARS-CoV-2 infection model. <i>European Journal of Immunology</i> , <b>2021</b> ,  B.1.617.2 enters and fuses lung cells with increased efficiency and evades antibodies induced by infection and vaccination. <i>Cell Reports</i> , <b>2021</b> , 37, 109825.  The Upper Respiratory Tract of Felids Is Highly Susceptible to SARS-CoV-2 Infection. <i>International</i>	15.4 17.4 6.1	20 0 14 31
182 181 180 179	Pre-clinical assessment of pharmacological and molecular properties. <i>Chemical Science</i> , <b>2021</b> , 12, 1260  Neutralization of the SARS-CoV-2 Delta variant after heterologous and homologous BNT162b2 or ChAdOx1 nCoV-19 vaccination. <i>Cellular and Molecular Immunology</i> , <b>2021</b> , 18, 2455-2456  Functional comparison of MERS-coronavirus lineages reveals increased replicative fitness of the recombinant lineage 5. <i>Nature Communications</i> , <b>2021</b> , 12, 5324  A pair of noncompeting neutralizing human monoclonal antibodies protecting from disease in a SARS-CoV-2 infection model. <i>European Journal of Immunology</i> , <b>2021</b> ,  B.1.617.2 enters and fuses lung cells with increased efficiency and evades antibodies induced by infection and vaccination. <i>Cell Reports</i> , <b>2021</b> , 37, 109825  The Upper Respiratory Tract of Felids Is Highly Susceptible to SARS-CoV-2 Infection. <i>International Journal of Molecular Sciences</i> , <b>2021</b> , 22,	15.4 17.4 6.1 10.6	20 0 14 31 3

# (2019-2020)

174	Pharmacological Inhibition of Acid Sphingomyelinase Prevents Uptake of SARS-CoV-2 by Epithelial Cells. <i>Cell Reports Medicine</i> , <b>2020</b> , 1, 100142	18	76
173	A Multibasic Cleavage Site in the Spike Protein of SARS-CoV-2 Is Essential for Infection of Human Lung Cells. <i>Molecular Cell</i> , <b>2020</b> , 78, 779-784.e5	17.6	965
172	Structural Basis for Potent Neutralization of Betacoronaviruses by Single-Domain Camelid Antibodies. <i>Cell</i> , <b>2020</b> , 181, 1004-1015.e15	56.2	319
171	SARS-CoV-2 Cell Entry Depends on ACE2 and TMPRSS2 and Is Blocked by a Clinically Proven Protease Inhibitor. <i>Cell</i> , <b>2020</b> , 181, 271-280.e8	56.2	10629
170	Polymorphisms in dipeptidyl peptidase 4 reduce host cell entry of Middle East respiratory syndrome coronavirus. <i>Emerging Microbes and Infections</i> , <b>2020</b> , 9, 155-168	18.9	53
169	LY6E impairs coronavirus fusion and confers immune control of viral disease 2020,		12
168	Camostat mesylate inhibits SARS-CoV-2 activation by TMPRSS2-related proteases and its metabolite GBPA exerts antiviral activity <b>2020</b> ,		30
167	Camostat Mesylate May Reduce Severity of Coronavirus Disease 2019 Sepsis: A First Observation <b>2020</b> , 2, e0284		25
166	LY6E impairs coronavirus fusion and confers immune control of viral disease. <i>Nature Microbiology</i> , <b>2020</b> , 5, 1330-1339	26.6	98
165	Sphingosine prevents binding of SARS-CoV-2 spike to its cellular receptor ACE2. <i>Journal of Biological Chemistry</i> , <b>2020</b> , 295, 15174-15182	5.4	19
164	Chloroquine does not inhibit infection of human lung cells with SARS-CoV-2. <i>Nature</i> , <b>2020</b> , 585, 588-590	0 50.4	243
163	Glycan-Gold Nanoparticles as Multifunctional Probes for Multivalent Lectin-Carbohydrate Binding: Implications for Blocking Virus Infection and Nanoparticle Assembly. <i>Journal of the American Chemical Society</i> , <b>2020</b> , 142, 18022-18034	16.4	20
162	Nafamostat Mesylate Blocks Activation of SARS-CoV-2: New Treatment Option for COVID-19. <i>Antimicrobial Agents and Chemotherapy</i> , <b>2020</b> , 64,	5.9	281
161	H2 influenza A virus is not pathogenic in Tmprss2 knock-out mice. Virology Journal, 2020, 17, 56	6.1	6
160	Interferon-Induced Transmembrane Proteins Mediate Viral Evasion in Acute and Chronic Hepatitis C Virus Infection. <i>Hepatology</i> , <b>2019</b> , 70, 1506-1520	11.2	11
159	Guanylate-Binding Proteins 2 and 5 Exert Broad Antiviral Activity by Inhibiting Furin-Mediated Processing of Viral Envelope Proteins. <i>Cell Reports</i> , <b>2019</b> , 27, 2092-2104.e10	10.6	53
158	Characterization of the Filovirus-Resistant Cell Line SH-SY5Y Reveals Redundant Role of Cell Surface Entry Factors. <i>Viruses</i> , <b>2019</b> , 11,	6.2	6
157	Disease Manifestation and Viral Sequences in a Bonobo More Than 30 Years after Papillomavirus Infection. <i>Pathogens</i> , <b>2019</b> , 8,	4.5	3

156	A system for production of defective interfering particles in the absence of infectious influenza A virus. <i>PLoS ONE</i> , <b>2019</b> , 14, e0212757	3.7	15
155	Calu-3 cells are largely resistant to entry driven by filovirus glycoproteins and the entry defect can be rescued by directed expression of DC-SIGN or cathepsin L. <i>Virology</i> , <b>2019</b> , 532, 22-29	3.6	12
154	Modulation of HIV-1 Gag/Gag-Pol frameshifting by tRNA abundance. <i>Nucleic Acids Research</i> , <b>2019</b> , 47, 5210-5222	20.1	19
153	Inhibitors of signal peptide peptidase and subtilisin/kexin-isozyme 1 inhibit Ebola virus glycoprotein-driven cell entry by interfering with activity and cellular localization of endosomal cathepsins. <i>PLoS ONE</i> , <b>2019</b> , 14, e0214968	3.7	1
152	Analysis of IFITM-IFITM Interactions by a Flow Cytometry-Based FRET Assay. <i>International Journal of Molecular Sciences</i> , <b>2019</b> , 20,	6.3	9
151	Spike proteins of novel MERS-coronavirus isolates from North- and West-African dromedary camels mediate robust viral entry into human target cells. <i>Virology</i> , <b>2019</b> , 535, 261-265	3.6	7
150	Novel Virus Related to Kaposiß Sarcoma-Associated Herpesvirus from Colobus Monkey. <i>Emerging Infectious Diseases</i> , <b>2019</b> , 25, 1548-1551	10.2	2
149	Kaposi Sarcoma in Mantled Guereza. <i>Emerging Infectious Diseases</i> , <b>2019</b> , 25, 1552-1555	10.2	1
148	Analysis of Resistance of Ebola Virus Glycoprotein-Driven Entry Against MDL28170, An Inhibitor of Cysteine Cathepsins. <i>Pathogens</i> , <b>2019</b> , 8,	4.5	3
147	Role of rhesus macaque IFITM3(2) in simian immunodeficiency virus infection of macaques. <i>PLoS ONE</i> , <b>2019</b> , 14, e0224082	3.7	
146	Hemagglutinin Cleavability, Acid Stability, and Temperature Dependence Optimize Influenza B Virus for Replication in Human Airways. <i>Journal of Virology</i> , <b>2019</b> , 94,	6.6	17
146 145	Hemagglutinin Cleavability, Acid Stability, and Temperature Dependence Optimize Influenza B	6.6	17 3
•	Hemagglutinin Cleavability, Acid Stability, and Temperature Dependence Optimize Influenza B Virus for Replication in Human Airways. <i>Journal of Virology</i> , <b>2019</b> , 94,  Seroprevalence of viral infections in captive rhesus and cynomolgus macaques. <i>Primate Biology</i> ,		•
145	Hemagglutinin Cleavability, Acid Stability, and Temperature Dependence Optimize Influenza B Virus for Replication in Human Airways. <i>Journal of Virology</i> , <b>2019</b> , 94,  Seroprevalence of viral infections in captive rhesus and cynomolgus macaques. <i>Primate Biology</i> , <b>2019</b> , 6, 1-6  Tmprss2 knock-out mice are resistant to H10 influenza A virus pathogenesis. <i>Journal of General</i>	0.9	3
145	Hemagglutinin Cleavability, Acid Stability, and Temperature Dependence Optimize Influenza B Virus for Replication in Human Airways. <i>Journal of Virology</i> , <b>2019</b> , 94,  Seroprevalence of viral infections in captive rhesus and cynomolgus macaques. <i>Primate Biology</i> , <b>2019</b> , 6, 1-6  Tmprss2 knock-out mice are resistant to H10 influenza A virus pathogenesis. <i>Journal of General Virology</i> , <b>2019</b> , 100, 1073-1078  Release of Immunomodulatory Ebola Virus Glycoprotein-Containing Microvesicles Is Suppressed by	0.9	3
145 144 143	Hemagglutinin Cleavability, Acid Stability, and Temperature Dependence Optimize Influenza B Virus for Replication in Human Airways. <i>Journal of Virology</i> , <b>2019</b> , 94,  Seroprevalence of viral infections in captive rhesus and cynomolgus macaques. <i>Primate Biology</i> , <b>2019</b> , 6, 1-6  Tmprss2 knock-out mice are resistant to H10 influenza A virus pathogenesis. <i>Journal of General Virology</i> , <b>2019</b> , 100, 1073-1078  Release of Immunomodulatory Ebola Virus Glycoprotein-Containing Microvesicles Is Suppressed by Tetherin in a Species-Specific Manner. <i>Cell Reports</i> , <b>2019</b> , 26, 1841-1853.e6  Mutations in the Spike Protein of Middle East Respiratory Syndrome Coronavirus Transmitted in	0.9 4.9 10.6	3 19 7
145 144 143	Hemagglutinin Cleavability, Acid Stability, and Temperature Dependence Optimize Influenza B Virus for Replication in Human Airways. <i>Journal of Virology</i> , <b>2019</b> , 94,  Seroprevalence of viral infections in captive rhesus and cynomolgus macaques. <i>Primate Biology</i> , <b>2019</b> , 6, 1-6  Tmprss2 knock-out mice are resistant to H10 influenza A virus pathogenesis. <i>Journal of General Virology</i> , <b>2019</b> , 100, 1073-1078  Release of Immunomodulatory Ebola Virus Glycoprotein-Containing Microvesicles Is Suppressed by Tetherin in a Species-Specific Manner. <i>Cell Reports</i> , <b>2019</b> , 26, 1841-1853.e6  Mutations in the Spike Protein of Middle East Respiratory Syndrome Coronavirus Transmitted in Korea Increase Resistance to Antibody-Mediated Neutralization. <i>Journal of Virology</i> , <b>2019</b> , 93,  Tetherin Inhibits Nipah Virus but Not Ebola Virus Replication in Fruit Bat Cells. <i>Journal of Virology</i> ,	0.9 4.9 10.6 6.6	3 19 7 84

# (2016-2018)

138	Cell Entry of Influenza A Viruses: Sweet Talk between HA and Ca1.2. <i>Cell Host and Microbe</i> , <b>2018</b> , 23, 697-699	23.4	4
137	Functional analysis of potential cleavage sites in the MERS-coronavirus spike protein. <i>Scientific Reports</i> , <b>2018</b> , 8, 16597	4.9	94
136	Priming Time: How Cellular Proteases Arm Coronavirus Spike Proteins <b>2018</b> , 71-98		48
135	A Polymorphism within the Internal Fusion Loop of the Ebola Virus Glycoprotein Modulates Host Cell Entry. <i>Journal of Virology</i> , <b>2017</b> , 91,	6.6	28
134	Herpes B virus replication and viral lesions in the liver of a cynomolgus macaque which died from severe disease with rapid onset. <i>Journal of Medical Primatology</i> , <b>2017</b> , 46, 256-259	0.7	1
133	pH Optimum of Hemagglutinin-Mediated Membrane Fusion Determines Sensitivity of Influenza A Viruses to the Interferon-Induced Antiviral State and IFITMs. <i>Journal of Virology</i> , <b>2017</b> , 91,	6.6	46
132	The glycoprotein of vesicular stomatitis virus promotes release of virus-like particles from tetherin-positive cells. <i>PLoS ONE</i> , <b>2017</b> , 12, e0189073	3.7	26
131	Dissecting Multivalent Lectin-Carbohydrate Recognition Using Polyvalent Multifunctional Glycan-Quantum Dots. <i>Journal of the American Chemical Society</i> , <b>2017</b> , 139, 11833-11844	16.4	41
130	Virion Background and Efficiency of Virion Incorporation Determine Susceptibility of Simian Immunodeficiency Virus Env-Driven Viral Entry to Inhibition by IFITM Proteins. <i>Journal of Virology</i> , <b>2017</b> , 91,	6.6	7
129	Rhesus macaque IFITM3 gene polymorphisms and SIV infection. <i>PLoS ONE</i> , <b>2017</b> , 12, e0172847	3.7	5
128	Non-human primate orthologues of TMPRSS2 cleave and activate the influenza virus hemagglutinin. <i>PLoS ONE</i> , <b>2017</b> , 12, e0176597	3.7	9
127	Different residues in the SARS-CoV spike protein determine cleavage and activation by the host cell protease TMPRSS2. <i>PLoS ONE</i> , <b>2017</b> , 12, e0179177	3.7	57
126	Detection systems for antibody responses against herpes B virus. Primate Biology, 2017, 4, 9-16	0.9	2
125	The Tetherin Antagonism of the Ebola Virus Glycoprotein Requires an Intact Receptor-Binding Domain and Can Be Blocked by GP1-Specific Antibodies. <i>Journal of Virology</i> , <b>2016</b> , 90, 11075-11086	6.6	17
124	Compact, Polyvalent Mannose Quantum Dots as Sensitive, Ratiometric FRET Probes for Multivalent Protein-Ligand Interactions. <i>Angewandte Chemie</i> , <b>2016</b> , 128, 4816-4820	3.6	5
123	Compact, Polyvalent Mannose Quantum Dots as Sensitive, Ratiometric FRET Probes for Multivalent Protein-Ligand Interactions. <i>Angewandte Chemie - International Edition</i> , <b>2016</b> , 55, 4738-42	16.4	45
122	The Proteolytic Activation of (H3N2) Influenza A Virus Hemagglutinin Is Facilitated by Different Type II Transmembrane Serine Proteases. <i>Journal of Virology</i> , <b>2016</b> , 90, 4298-4307	6.6	34
121	Evidence that Processing of the Severe Fever with Thrombocytopenia Syndrome Virus Gn/Gc Polyprotein Is Critical for Viral Infectivity and Requires an Internal Gc Signal Peptide. <i>PLoS ONE</i> , <b>2016</b> , 11, e0166013	3.7	17

120	The Role of Phlebovirus Glycoproteins in Viral Entry, Assembly and Release. Viruses, 2016, 8,	6.2	33
119	The Glycoproteins of All Filovirus Species Use the Same Host Factors for Entry into Bat and Human Cells but Entry Efficiency Is Species Dependent. <i>PLoS ONE</i> , <b>2016</b> , 11, e0149651	3.7	27
118	The Hemagglutinin of Bat-Associated Influenza Viruses Is Activated by TMPRSS2 for pH-Dependent Entry into Bat but Not Human Cells. <i>PLoS ONE</i> , <b>2016</b> , 11, e0152134	3.7	19
117	Tetherin Sensitivity of Influenza A Viruses Is Strain Specific: Role of Hemagglutinin and Neuraminidase. <i>Journal of Virology</i> , <b>2015</b> , 89, 9178-88	6.6	24
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109	The clinically approved drugs amiodarone, dronedarone and verapamil inhibit filovirus cell entry.		140 62
	The clinically approved drugs amiodarone, dronedarone and verapamil inhibit filovirus cell entry. Journal of Antimicrobial Chemotherapy, 2014, 69, 2123-31  DESC1 and MSPL activate influenza A viruses and emerging coronaviruses for host cell entry.	5.1	
108	The clinically approved drugs amiodarone, dronedarone and verapamil inhibit filovirus cell entry.  Journal of Antimicrobial Chemotherapy, 2014, 69, 2123-31  DESC1 and MSPL activate influenza A viruses and emerging coronaviruses for host cell entry.  Journal of Virology, 2014, 88, 12087-97  Toll-like receptor 3 signalling up-regulates expression of the HIV co-receptor G-protein coupled	5.1 6.6	62
108	The clinically approved drugs amiodarone, dronedarone and verapamil inhibit filovirus cell entry. <i>Journal of Antimicrobial Chemotherapy</i> , <b>2014</b> , 69, 2123-31  DESC1 and MSPL activate influenza A viruses and emerging coronaviruses for host cell entry. <i>Journal of Virology</i> , <b>2014</b> , 88, 12087-97  Toll-like receptor 3 signalling up-regulates expression of the HIV co-receptor G-protein coupled receptor 15 on human CD4+ T cells. <i>PLoS ONE</i> , <b>2014</b> , 9, e88195  Influenza A virus encoding secreted Gaussia luciferase as useful tool to analyze viral replication and	5.1 6.6 3.7	62
108	The clinically approved drugs amiodarone, dronedarone and verapamil inhibit filovirus cell entry. <i>Journal of Antimicrobial Chemotherapy</i> , <b>2014</b> , 69, 2123-31  DESC1 and MSPL activate influenza A viruses and emerging coronaviruses for host cell entry. <i>Journal of Virology</i> , <b>2014</b> , 88, 12087-97  Toll-like receptor 3 signalling up-regulates expression of the HIV co-receptor G-protein coupled receptor 15 on human CD4+ T cells. <i>PLoS ONE</i> , <b>2014</b> , 9, e88195  Influenza A virus encoding secreted Gaussia luciferase as useful tool to analyze viral replication and its inhibition by antiviral compounds and cellular proteins. <i>PLoS ONE</i> , <b>2014</b> , 9, e97695  IFITM proteins inhibit entry driven by the MERS-coronavirus spike protein: evidence for	5.1 6.6 3.7 3.7	62 10 39

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15	The Omicron variant is highly resistant against antibody-mediated neutralization Implications for control of the COVID-19 pandemic		13
14	The novel coronavirus 2019 (2019-nCoV) uses the SARS-coronavirus receptor ACE2 and the cellular protease TMPRSS2 for entry into target cells		284
13	Novel surrogate virus neutralization test reveals low serum neutralizing anti-SARS-CoV-2-S antibodies levels in mildly affected COVID-19 convalescents		5

12	Rapid SARS-CoV-2 Adaptation to Available Cellular Proteases	7
11	The SARS-CoV-2 and other human coronavirus spike proteins are fine-tuned towards temperature and proteases of the human airways	4
10	Evidence for influenza B virus hemagglutinin adaptation to the human host: high cleavability, acid-stability and preference for cool temperature	1
9	Humoral and cellular immune responses against SARS-CoV-2 variants and human coronaviruses after single BNT162b2 vaccination	4
8	A pair of non-competing neutralizing human monoclonal antibodies protecting from disease in a SARS-CoV-2 infection model	3
7	SARS-CoV-2 variant B.1.617 is resistant to Bamlanivimab and evades antibodies induced by infection and vaccination	48
6	Humoral and cellular immune response against SARS-CoV-2 variants following heterologous and homologous ChAdOx1 nCoV-19/BNT162b2 vaccination	6
5	Increased lung cell entry of B.1.617.2 and evasion of antibodies induced by infection and BNT162b2 vaccination	
4	Heterologous ChAdOx1 nCoV-19 and BNT162b2 prime-boost vaccination elicits potent neutralizing antibody responses and T cell reactivity	20
3	SARS-CoV-2 variants B.1.351 and B.1.1.248: Escape from therapeutic antibodies and antibodies induced by infection and vaccination	39
2	SARS-CoV-2 mutations acquired in mink reduce antibody-mediated neutralization	3
1	BNT162b2 boosted immune responses six months after heterologous or homologous ChAdOx1nCoV-19/BNT162b2 vaccination against COVID-19	1