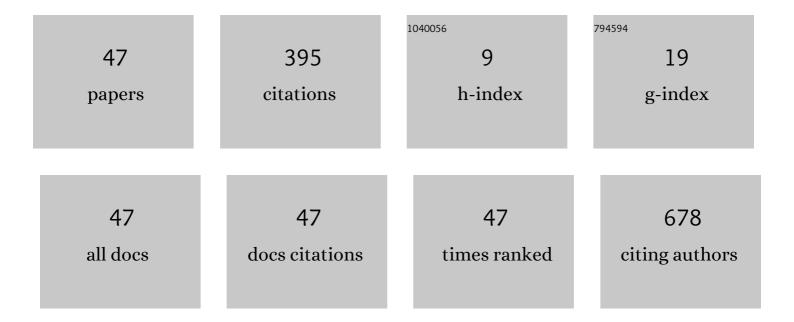
Sriram Krishnaswamy

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Generation of an anticoagulant aptamer that targets factor V/Va and disrupts the FVa-membrane interaction in normal and COVID-19 patient samples. Cell Chemical Biology, 2022, 29, 215-225.e5.	5.2	5
2	ISTH congress 2021. Journal of Thrombosis and Haemostasis, 2021, 19, 1385-1385.	3.8	0
3	Exosite binding drives substrate affinity for the activation of coagulation factor X by the intrinsic Xase complex. Journal of Biological Chemistry, 2020, 295, 15198-15207.	3.4	6
4	FcRn augments induction of tissue factor activity by IgG-containing immune complexes. Blood, 2020, 135, 2085-2093.	1.4	19
5	Occlusion of anion-binding exosite 2 in meizothrombin explains its impaired ability to activate factor V. Journal of Biological Chemistry, 2019, 294, 2422-2435.	3.4	6
6	Advances in Clinical and Basic Science of Coagulation: Illustrated abstracts of the 9th Chapel Hill Symposium on Hemostasis. Research and Practice in Thrombosis and Haemostasis, 2018, 2, 407-428.	2.3	5
7	Combination of aptamer and drug for reversible anticoagulation in cardiopulmonary bypass. Nature Biotechnology, 2018, 36, 606-613.	17.5	52
8	JNJ-64179375 Inhibits Exosite I-Mediated Thrombin Activity While Preserving Exosite II and Active Site Function in Vitro. Blood, 2018, 132, 24-24.	1.4	2
9	A Novel Variant of Factor VIII Yields Cofactor Activity without Proteolysis between the A1 and A2 Domains. Blood, 2018, 132, 3771-3771.	1.4	0
10	Phosphatidylinositol transfer protein-α in platelets is inconsequential for thrombosis yet is utilized for tumor metastasis. Nature Communications, 2017, 8, 1216.	12.8	22
11	Selective factor VIII activation by the tissue factor–factor VIIa–factor Xa complex. Blood, 2017, 130, 1661-1670.	1.4	58
12	The Fragment 1 Region of Prothrombin Facilitates the Favored Binding of Fragment 12 to Zymogen and Enforces Zymogen-like Character in the Proteinase. Journal of Biological Chemistry, 2016, 291, 11114-11123.	3.4	3
13	A rapid pro-hemostatic approach to overcome direct oral anticoagulants. Nature Medicine, 2016, 22, 924-932.	30.7	39
14	FVIII-VWF dos-Ã-dos. Blood, 2015, 126, 923-924.	1.4	6
15	The NET Effect: Platelet Factor 4 and DNA-Histone Interactions in Sepsis. Blood, 2015, 126, 2197-2197.	1.4	1
16	The Affinity of Factor X for the Intrinsic Xase Is Determined By Exosite Interactions Between the Substrate and the Enzyme Complex. Blood, 2015, 126, 1065-1065.	1.4	0
17	Long-Range Allosteric Linkage Between Exosites Reciprocally Regulates the Zymogenicity of Prothrombin Derivatives. Blood, 2015, 126, 122-122.	1.4	1
18	Prothrombin Membrane Binding and Gla-Dependent Function Are Not Required for Effective Hemostasis In Vivo. Blood, 2015, 126, 124-124.	1.4	1

#	Article	IF	CITATIONS
19	Platelet Pitp-Alpha Promotes Thrombin Generation and the Dissemination of Tumor Metastasis, but Has Minimal Effect on Vascular Plug Formation. Blood, 2015, 126, 418-418.	1.4	1
20	The X-Ray Structure of a Variant of Human Factor V Provides Structural Insights into the Procofactor Activation Paradox. Blood, 2015, 126, 121-121.	1.4	1
21	Assembly of Prothrombinase on Endothelial Cells: Receptor-Mediated or Phospholipid-Driven?. Blood, 2015, 126, 2267-2267.	1.4	Ο
22	New insights into the spatiotemporal localization of prothrombinase in vivo. Blood, 2014, 124, 1705-1714.	1.4	85
23	Distinct Mechanism of Anti-Hemophilic FVIII Activation By Nascent FXa in the Tissue Factor-FVIIa Coagulation Initiation Complex Resistant to FXa-Directed Inhibition. Blood, 2014, 124, 2803-2803.	1.4	1
24	X-Ray Structure of an Anticoagulant RNA Aptamer Bound to Factor Xa. Structural Basis for Its Ability to Disrupt Interactions Between Xa and Va within Prothrombinase. Blood, 2014, 124, 4232-4232.	1.4	0
25	A Proposed Role for Platelet Factor 4 in Histone Pathobiology in Sepsis. Blood, 2014, 124, 98-98.	1.4	Ο
26	New Structural Insights into High Affinity Membrane Binding By Coagulation Factor V/Va. Blood, 2014, 124, 4216-4216.	1.4	0
27	Structural and Functional Studies of Î ³ -Carboxyglutamic Acid Domains of Factor VIIa and Activated Protein C: Role of Magnesium at Physiological Calcium. Journal of Molecular Biology, 2013, 425, 1961-1981.	4.2	26
28	Restoring the Procofactor State of Factor Va-like Variants by Complementation with B-domain Peptides. Journal of Biological Chemistry, 2013, 288, 30151-30160.	3.4	28
29	Killing 2 proteinases with 1 (dual-acting) stone. Blood, 2012, 119, 2182-2183.	1.4	0
30	Conformational Dynamics of Prothrombin Dictate Its Ordered Cleavage by Prothrombinase. Blood, 2012, 120, 3359-3359.	1.4	0
31	High Resolution X-Ray Structure of Snake Venom Factor V: Evolution of a Hemostatic Cofactor to a Toxin Poised to Inflict Maximal Damage to Mammalian Blood Coagulation. Blood, 2011, 118, 375-375.	1.4	1
32	Mg2+ Is Required for Optimal Folding of the γ-Carboxyglutamic Acid (Gla) Domains of Vitamin K-Dependent Clotting Factors At Physiological Ca2+. Blood, 2011, 118, 1172-1172.	1.4	1
33	Meizothrombin: Zymogen or Proteinase? Slow, Ligand-Dependent Equilibration Between Equally Populated Zymogen-Like and Proteinase-Like Forms Explains Its Selectively Anticoagulant Function. Blood, 2011, 118, 533-533.	1.4	Ο
34	Imaging of Coagulation Reactions During Thrombus Formation In Vivo with Novel Fluorescent Protein Derivatives. Blood, 2010, 116, 817-817.	1.4	1
35	Heparin-Induced Thrombocytopenia Antibodies Inhibit PF4-Dependent Enhancement of Activated Protein C Formation by Binding to Antigenic Complexes Formed with the Chondroitin Sulfate Side-Chain of Thrombomodulin. Blood, 2010, 116, 721-721.	1.4	0
36	Meizothrombin Is Unexpectedly Zymogen-Like: Its Slow Conversion to Proteinase Dominates Thrombin Production by Prothrombinase. Blood, 2010, 116, 2215-2215.	1.4	1

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#	Article	IF	CITATIONS
37	Driving Thrombin From a Protease-Like State to a Zymogen-Like State and Back Blood, 2009, 114, 852-852.	1.4	0
38	Structural and Functional Studies to Define the Molecular Basis by Which Platelet Factor 4 (PF4) Increases Survival of Mice in Lipopolysaccharide (LPS)-Induced Endotoxicity. Blood, 2008, 112, 19-19.	1.4	21
39	Effects of Substrate Geometry on Active Site Engagement Govern the Preferential Action of Prothrombinase on One of the Two Cleavage Sites in Prothrombin Blood, 2008, 112, 2022-2022.	1.4	Ο
40	Fluorescence Resonance Energy Transfer Studies of Prothrombin Recognition by Prothrombinase Blood, 2008, 112, 2014-2014.	1.4	0
41	Conformational Activation of Zymogen-Like Thrombin Variants by Tight Binding Ligands. Blood, 2008, 112, 3070-3070.	1.4	0
42	Human Platelets and Endothelial Cells Differentially Regulate the Pathway for Prothrombin Cleavage by Prothrombinase Blood, 2007, 110, 269-269.	1.4	0
43	Anticoagulation by a thrombin precursor. Blood, 2004, 104, 301-301.	1.4	Ο
44	Active Site-Dependent Substrate Recognition Plays a Primary Role in Determining the Affinity of the Thrombin-Thrombomodulin Complex for Protein C Blood, 2004, 104, 125-125.	1.4	1
45	The Substrate -Membrane Interaction Constrains Bond Presentation and Cleavage in the Action of Prothrombinase on Prothrombin Blood, 2004, 104, 1712-1712.	1.4	Ο
46	Ratcheting between Two Distinct Conformations of Substrate Drives the Sequential Cleavage of Prothrombin by Prothrombinase Blood, 2004, 104, 1717-1717.	1.4	0
47	Clotting in whole blood: analysis of a biochemical reaction network. Blood, 2002, 100, 1-1.	1.4	1