

Hong-Yu Hu

List of Publications by Year in descending order

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Version: 2024-02-01

31
papers

1,051
citations

430874

18
h-index

454955

30
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34
all docs

34
docs citations

34
times ranked

1674
citing authors

#	ARTICLE	IF	CITATIONS
1	RNA-assisted sequestration of RNA-binding proteins by cytoplasmic inclusions of the C-terminal 35-kDa fragment of TDP-43. <i>Journal of Cell Science</i> , 2022, 135, .	2.0	8
2	PolyQ-expanded proteins impair cellular proteostasis of ataxin-3 through sequestering the co-chaperone HSP90 into aggregates. <i>Scientific Reports</i> , 2021, 11, 7815.	3.3	11
3	O-GlcNAcylation of TDP-43 suppresses proteinopathies and promotes TDP-43's mRNA splicing activity. <i>EMBO Reports</i> , 2021, 22, e51649.	4.5	15
4	Solid-State NMR Reveals the Structural Transformation of the TDP-43 Amyloidogenic Region upon Fibrillation. <i>Journal of the American Chemical Society</i> , 2020, 142, 3412-3421.	13.7	51
5	Domain interactions reveal auto-inhibition of the deubiquitinating enzyme USP19 and its activation by HSP90 in the modulation of huntingtin aggregation. <i>Biochemical Journal</i> , 2020, 477, 4295-4312.	3.7	5
6	Structural and dynamic studies reveal that the Ala-rich region of ataxin-7 initiates α -helix formation of the polyQ tract but suppresses its aggregation. <i>Scientific Reports</i> , 2019, 9, 7481.	3.3	13
7	PolyQ-expanded huntingtin and ataxin-3 sequester ubiquitin adaptors hHR23B and UBQLN2 into aggregates via conjugated ubiquitin. <i>FASEB Journal</i> , 2018, 32, 2923-2933.	0.5	24
8	Structural and Functional Investigations of the N-Terminal Ubiquitin Binding Region of Usp25. <i>Biophysical Journal</i> , 2017, 112, 2099-2108.	0.5	6
9	The N-terminal dimerization is required for TDP-43 splicing activity. <i>Scientific Reports</i> , 2017, 7, 6196.	3.3	78
10	HSP90 recognizes the N-terminus of huntingtin involved in regulation of huntingtin aggregation by USP19. <i>Scientific Reports</i> , 2017, 7, 14797.	3.3	35
11	Editorial: Structural Aspects of Protein Aggregation. <i>Protein and Peptide Letters</i> , 2017, 24, 280-280.	0.9	2
12	Sequestration of cellular interacting partners by protein aggregates: implication in a loss-of-function pathology. <i>FEBS Journal</i> , 2016, 283, 3705-3717.	4.7	70
13	Two mutations G335D and Q343R within the amyloidogenic core region of TDP-43 influence its aggregation and inclusion formation. <i>Scientific Reports</i> , 2016, 6, 23928.	3.3	64
14	Cytoplasmic Ubiquitin-Specific Protease 19 (USP19) Modulates Aggregation of Polyglutamine-Expanded Ataxin-3 and Huntingtin through the HSP90 Chaperone. <i>PLoS ONE</i> , 2016, 11, e0147515.	2.5	34
15	Study of Protein Amyloid-Like Aggregates by Solid-State Circular Dichroism Spectroscopy. <i>Current Protein and Peptide Science</i> , 2016, 18, 100-103.	1.4	5
16	The N-terminal ubiquitin-binding region of ubiquitin-specific protease 28 modulates its deubiquitination function: NMR structural and mechanistic insights. <i>Biochemical Journal</i> , 2015, 471, 155-165.	3.7	8
17	TDP-35 sequesters TDP-43 into cytoplasmic inclusions through binding with RNA. <i>FEBS Letters</i> , 2015, 589, 1920-1928.	2.8	40
18	Aggregation of Polyglutamine-expanded Ataxin 7 Protein Specifically Sequesters Ubiquitin-specific Protease 22 and Deteriorates Its Deubiquitinating Function in the Spt-Ada-Gcn5-Acetyltransferase (SAGA) Complex. <i>Journal of Biological Chemistry</i> , 2015, 290, 21996-22004.	3.4	30

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19	A Ubiquitin Shuttle DC-Ubp/UBTD2 Reconciles Protein Ubiquitination and Deubiquitination via Linking UbE1 and USP5 Enzymes. PLoS ONE, 2014, 9, e107509.	2.5	4
20	Autoinhibitory Structure of the WW Domain of HYPB/SETD2 Regulates Its Interaction with the Proline-Rich Region of Huntingtin. Structure, 2014, 22, 378-386.	3.3	39
21	Aggregation of polyglutamine-expanded ataxin-3 sequesters its specific interacting partners into inclusions: Implication in a loss-of-function pathology. Scientific Reports, 2014, 4, 6410.	3.3	110
22	Structural Transformation of the Amyloidogenic Core Region of TDP-43 Protein Initiates Its Aggregation and Cytoplasmic Inclusion. Journal of Biological Chemistry, 2013, 288, 19614-19624.	3.4	124
23	Length of the active-site crossover loop defines the substrate specificity of ubiquitin C-terminal hydrolases for ubiquitin chains. Biochemical Journal, 2012, 441, 143-149.	3.7	48
24	The C-terminal Helices of Heat Shock Protein 70 Are Essential for J-domain Binding and ATPase Activation. Journal of Biological Chemistry, 2012, 287, 6044-6052.	3.4	26
25	Domain Analysis Reveals That a Deubiquitinating Enzyme USP13 Performs Non-Activating Catalysis for Lys63-Linked Polyubiquitin. PLoS ONE, 2011, 6, e29362.	2.5	41
26	Aggregation of the 35 kDa fragment of TDP-43 causes formation of cytoplasmic inclusions and alteration of RNA processing. FASEB Journal, 2011, 25, 2344-2353.	0.5	62
27	Differential ubiquitin binding of the UBA domains from human Cbl and Cblb: NMR structural and biochemical insights. Protein Science, 2008, 17, 1805-1814.	7.6	23
28	Highly efficient expression and purification system of small-size protein domains in Escherichia coli for biochemical characterization. Protein Expression and Purification, 2006, 47, 599-606.	1.3	63
29	The concentration of hydrogen peroxide generated during aggregation of α -synuclein <i>in vitro</i> is lower than 5 nmol/L. Chinese Journal of Chemistry, 2004, 22, 1440-1443.	4.9	0
30	Novel Secondary Structure of Calcitonin in Solid State as Revealed by Circular Dichroism Spectroscopy. Chinese Journal of Chemistry, 2002, 20, 697-698.	4.9	0
31	Structural Transformation of the Peptide Fragments from the Reactive Center Loops of Serpins: Circular Dichroic Studies. Chinese Journal of Chemistry, 2001, 19, 954-959.	4.9	2