Lorenzo A Pinna

List of Publications by Citations

Source: https://exaly.com/author-pdf/6633537/lorenzo-a-pinna-publications-by-citations.pdf

Version: 2024-04-10

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

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

103 12,014 210 57 h-index g-index citations papers 6.32 211 12,959 5.4 avg, IF L-index ext. citations ext. papers

| # | Paper | IF | Citations |
|-------------|--|-----------------|-----------|
| 21 0 | One-thousand-and-one substrates of protein kinase CK2?. FASEB Journal, 2003, 17, 349-68 | 0.9 | 1067 |
| 209 | Casein kinase 2: an 'eminence grise' in cellular regulation?. <i>Biochimica Et Biophysica Acta - Molecular Cell Research</i> , 1990 , 1054, 267-84 | 4.9 | 807 |
| 208 | Protein kinase CK2: a challenge to canons. <i>Journal of Cell Science</i> , 2002 , 115, 3873-8 | 5.3 | 382 |
| 207 | How do protein kinases recognize their substrates?. <i>Biochimica Et Biophysica Acta - Molecular Cell Research</i> , 1996 , 1314, 191-225 | 4.9 | 362 |
| 206 | Selectivity of 4,5,6,7-tetrabromobenzotriazole, an ATP site-directed inhibitor of protein kinase CK2 ('casein kinase-2'). <i>FEBS Letters</i> , 2001 , 496, 44-8 | 3.8 | 290 |
| 205 | The protein kinase CK2 facilitates repair of chromosomal DNA single-strand breaks. <i>Cell</i> , 2004 , 117, 17 | - 2§ 6.2 | 275 |
| 204 | Protein kinase CK2 ("casein kinase-2") and its implication in cell division and proliferation. <i>Progress in Cell Cycle Research</i> , 1997 , 3, 77-97 | | 263 |
| 203 | Addiction to protein kinase CK2: a common denominator of diverse cancer cells?. <i>Biochimica Et Biophysica Acta - Proteins and Proteomics</i> , 2010 , 1804, 499-504 | 4 | 253 |
| 202 | Different susceptibility of protein kinases to staurosporine inhibition. Kinetic studies and molecular bases for the resistance of protein kinase CK2. <i>FEBS Journal</i> , 1995 , 234, 317-22 | | 222 |
| 201 | Protein kinase CK2 inhibitor 4,5,6,7-tetrabromobenzotriazole (TBB) induces apoptosis and caspase-dependent degradation of haematopoietic lineage cell-specific protein 1 (HS1) in Jurkat cells. <i>Biochemical Journal</i> , 2002 , 364, 41-7 | 3.8 | 200 |
| 200 | The selectivity of inhibitors of protein kinase CK2: an update. <i>Biochemical Journal</i> , 2008 , 415, 353-65 | 3.8 | 193 |
| 199 | A Single Kinase Generates the Majority of the Secreted Phosphoproteome. <i>Cell</i> , 2015 , 161, 1619-32 | 56.2 | 187 |
| 198 | Site specificity of casein kinase-2 (TS) from rat liver cytosol. A study with model peptide substrates. <i>FEBS Journal</i> , 1986 , 160, 239-44 | | 175 |
| 197 | 2-Dimethylamino-4,5,6,7-tetrabromo-1H-benzimidazole: a novel powerful and selective inhibitor of protein kinase CK2. <i>Biochemical and Biophysical Research Communications</i> , 2004 , 321, 1040-4 | 3.4 | 160 |
| 196 | Optimization of protein kinase CK2 inhibitors derived from 4,5,6,7-tetrabromobenzimidazole. <i>Journal of Medicinal Chemistry</i> , 2004 , 47, 6239-47 | 8.3 | 151 |
| 195 | Casein kinase 2 down-regulation and activation by polybasic peptides are mediated by acidic residues in the 55-64 region of the beta-subunit. A study with calmodulin as phosphorylatable substrate. <i>Biochemistry</i> , 1994 , 33, 4336-42 | 3.2 | 149 |
| 194 | Unprecedented selectivity and structural determinants of a new class of protein kinase CK2 inhibitors in clinical trials for the treatment of cancer. <i>Biochemistry</i> , 2011 , 50, 8478-88 | 3.2 | 132 |

(2013-2009)

| 193 | Extraordinary pleiotropy of protein kinase CK2 revealed by weblogo phosphoproteome analysis. <i>Biochimica Et Biophysica Acta - Molecular Cell Research</i> , 2009 , 1793, 847-59 | 4.9 | 128 |
|-----|---|-----------------|-----|
| 192 | Quinalizarin as a potent, selective and cell-permeable inhibitor of protein kinase CK2. <i>Biochemical Journal</i> , 2009 , 421, 387-95 | 3.8 | 127 |
| 191 | Biochemical and three-dimensional-structural study of the specific inhibition of protein kinase CK2 by [5-oxo-5,6-dihydroindolo-(1,2-a)quinazolin-7-yl]acetic acid (IQA). <i>Biochemical Journal</i> , 2003 , 374, 639- | 46 ⁸ | 127 |
| 190 | The replacement of ATP by the competitive inhibitor emodin induces conformational modifications in the catalytic site of protein kinase CK2. <i>Journal of Biological Chemistry</i> , 2000 , 275, 29618-22 | 5.4 | 124 |
| 189 | Restoration of CFTR function in patients with cystic fibrosis carrying the F508del-CFTR mutation. <i>Autophagy</i> , 2014 , 10, 2053-74 | 10.2 | 119 |
| 188 | Identification of ellagic acid as potent inhibitor of protein kinase CK2: a successful example of a virtual screening application. <i>Journal of Medicinal Chemistry</i> , 2006 , 49, 2363-6 | 8.3 | 119 |
| 187 | Structural features underlying selective inhibition of protein kinase CK2 by ATP site-directed tetrabromo-2-benzotriazole. <i>Protein Science</i> , 2001 , 10, 2200-6 | 6.3 | 114 |
| 186 | Multiple myeloma cell survival relies on high activity of protein kinase CK2. <i>Blood</i> , 2006 , 108, 1698-707 | 2.2 | 113 |
| 185 | Coumarin as attractive casein kinase 2 (CK2) inhibitor scaffold: an integrate approach to elucidate the putative binding motif and explain structure-activity relationships. <i>Journal of Medicinal Chemistry</i> , 2008 , 51, 752-9 | 8.3 | 109 |
| 184 | Inhibition of protein kinase CK2 by flavonoids and tyrphostins. A structural insight. <i>Biochemistry</i> , 2012 , 51, 6097-107 | 3.2 | 105 |
| 183 | Tetrabromocinnamic acid (TBCA) and related compounds represent a new class of specific protein kinase CK2 inhibitors. <i>ChemBioChem</i> , 2007 , 8, 129-39 | 3.8 | 104 |
| 182 | Inspecting the structure-activity relationship of protein kinase CK2 inhibitors derived from tetrabromo-benzimidazole. <i>Chemistry and Biology</i> , 2005 , 12, 1211-9 | | 104 |
| 181 | Protein kinase CK2alpha' is induced by serum as a delayed early gene and cooperates with Ha-ras in fibroblast transformation. <i>Journal of Biological Chemistry</i> , 1998 , 273, 21291-7 | 5.4 | 103 |
| 180 | A noncanonical sequence phosphorylated by casein kinase 1 in beta-catenin may play a role in casein kinase 1 targeting of important signaling proteins. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2003 , 100, 10193-200 | 11.5 | 100 |
| 179 | Protein kinase CK2 as a druggable target. <i>Molecular BioSystems</i> , 2008 , 4, 889-94 | | 98 |
| 178 | Aurora-A site specificity: a study with synthetic peptide substrates. <i>Biochemical Journal</i> , 2005 , 390, 293- | 302 | 98 |
| 177 | Ribofuranosyl-benzimidazole derivatives as inhibitors of casein kinase-2 and casein kinase-1. <i>FEBS Journal</i> , 1990 , 187, 89-94 | | 95 |
| 176 | Secreted protein kinases. <i>Trends in Biochemical Sciences</i> , 2013 , 38, 121-30 | 10.3 | 94 |

| 175 | The raison d'tre of constitutively active protein kinases: the lesson of CK2. <i>Accounts of Chemical Research</i> , 2003 , 36, 378-84 | 24.3 | 92 |
|-----|--|------|----|
| 174 | The ATP-binding site of protein kinase CK2 holds a positive electrostatic area and conserved water molecules. <i>ChemBioChem</i> , 2007 , 8, 1804-9 | 3.8 | 86 |
| 173 | Subunit structure and autophosphorylation mechanism of casein kinase-TS (type-2) from rat liver cytosol. <i>FEBS Journal</i> , 1984 , 145, 593-9 | | 86 |
| 172 | Development and exploitation of CK2 inhibitors. <i>Molecular and Cellular Biochemistry</i> , 2005 , 274, 69-76 | 4.2 | 83 |
| 171 | Inhibition of protein kinase CK2 by condensed polyphenolic derivatives. An in vitro and in vivo study. <i>Biochemistry</i> , 2004 , 43, 12931-6 | 3.2 | 82 |
| 170 | Phosphorylation by protein kinase CK2 changes the DNA binding properties of the human chromatin protein DEK. <i>Molecular and Cellular Biology</i> , 2004 , 24, 6011-20 | 4.8 | 79 |
| 169 | Multiple phosphorylation of alpha-synuclein by protein tyrosine kinase Syk prevents eosin-induced aggregation. <i>FASEB Journal</i> , 2002 , 16, 210-2 | 0.9 | 79 |
| 168 | Casein kinase: the triple meaning of a misnomer. <i>Biochemical Journal</i> , 2014 , 460, 141-56 | 3.8 | 76 |
| 167 | Golgi apparatus mammary gland casein kinase: monitoring by a specific peptide substrate and definition of specificity determinants. <i>FEBS Letters</i> , 1996 , 382, 149-52 | 3.8 | 72 |
| 166 | Phosphorylated synthetic peptides as tools for studying protein phosphatases. <i>Biochimica Et Biophysica Acta - Molecular Cell Research</i> , 1994 , 1222, 415-31 | 4.9 | 71 |
| 165 | Rat liver Golgi apparatus contains a protein kinase similar to the casein kinase of lactating mammary gland. <i>FEBS Journal</i> , 1997 , 243, 719-25 | | 69 |
| 164 | The consensus sequences for cdc2 kinase and for casein kinase-2 are mutually incompatible. A study with peptides derived from the beta-subunit of casein kinase-2. <i>FEBS Letters</i> , 1992 , 301, 111-4 | 3.8 | 69 |
| 163 | Involvement of protein kinase CK2 in angiogenesis and retinal neovascularization. <i>Investigative Ophthalmology and Visual Science</i> , 2004 , 45, 4583-91 | | 68 |
| 162 | Site specificity of p72syk protein tyrosine kinase: efficient phosphorylation of motifs recognized by Src homology 2 domains of the Src family. <i>FEBS Letters</i> , 1995 , 367, 149-52 | 3.8 | 66 |
| 161 | Phosphorylation of osteopontin by Golgi apparatus casein kinase. <i>Biochemical and Biophysical Research Communications</i> , 1997 , 240, 602-5 | 3.4 | 65 |
| 160 | Inhibition of protein kinase CK2 by anthraquinone-related compounds. A structural insight. <i>Journal of Biological Chemistry</i> , 2003 , 278, 1831-6 | 5.4 | 65 |
| 159 | Autophosphorylation of type 2 casein kinase TS at both its alpha- and beta-subunits. Influence of different effectors. <i>FEBS Letters</i> , 1983 , 160, 203-8 | 3.8 | 62 |
| 158 | Casein kinase-2 structure-function relationship: creation of a set of mutants of the beta subunit that variably surrogate the wildtype beta subunit function. <i>Biochemical and Biophysical Research Communications</i> , 1992 , 188, 228-34 | 3.4 | 61 |

| 157 | Casein kinases as potential therapeutic targets. Expert Opinion on Therapeutic Targets, 2016, 20, 319-40 | 6.4 | 59 |
|-----|---|------|----|
| 156 | Unique activation mechanism of protein kinase CK2. The N-terminal segment is essential for constitutive activity of the catalytic subunit but not of the holoenzyme. <i>Journal of Biological Chemistry</i> , 2002 , 277, 22509-14 | 5.4 | 59 |
| 155 | Chemical derivatization of phosphoserine and phosphothreonine containing peptides to increase sensitivity for MALDI-based analysis and for selectivity of MS/MS analysis. <i>Proteomics</i> , 2006 , 6, 757-66 | 4.8 | 57 |
| 154 | Features and potentials of ATP-site directed CK2 inhibitors. <i>Biochimica Et Biophysica Acta - Proteins and Proteomics</i> , 2005 , 1754, 263-70 | 4 | 57 |
| 153 | Protein kinase CK2 inhibitors: a patent review. Expert Opinion on Therapeutic Patents, 2012, 22, 1081-97 | 6.8 | 56 |
| 152 | Protein kinase CK2 in health and disease: Protein kinase CK2: an ugly duckling in the kinome pond. <i>Cellular and Molecular Life Sciences</i> , 2009 , 66, 1795-9 | 10.3 | 56 |
| 151 | Analysis of the interaction between piD261/Bud32, an evolutionarily conserved protein kinase of Saccharomyces cerevisiae, and the Grx4 glutaredoxin. <i>Biochemical Journal</i> , 2004 , 377, 395-405 | 3.8 | 55 |
| 150 | Dephosphorylation and inactivation of Akt/PKB is counteracted by protein kinase CK2 in HEK 293T cells. <i>Cellular and Molecular Life Sciences</i> , 2009 , 66, 3363-73 | 10.3 | 52 |
| 149 | SH2 domains mediate the sequential phosphorylation of HS1 protein by p72syk and Src-related protein tyrosine kinases. <i>Biochemistry</i> , 1996 , 35, 5327-32 | 3.2 | 51 |
| 148 | Role of phosphorylated aminoacyl residues in generating atypical consensus sequences which are recognized by casein kinase-2 but not by casein kinase-1. <i>Biochemistry</i> , 1992 , 31, 5893-7 | 3.2 | 51 |
| 147 | Tetraiodobenzimidazoles are potent inhibitors of protein kinase CK2. <i>Bioorganic and Medicinal Chemistry</i> , 2009 , 17, 7281-9 | 3.4 | 50 |
| 146 | Structural determinants of protein kinase CK2 regulation by autoinhibitory polymerization. <i>ACS Chemical Biology</i> , 2012 , 7, 1158-63 | 4.9 | 47 |
| 145 | Mutational analysis of residues implicated in the interaction between protein kinase CK2 and peptide substrates. <i>Biochemistry</i> , 1997 , 36, 11717-24 | 3.2 | 47 |
| 144 | CK2-dependent phosphorylation of the E2 ubiquitin conjugating enzyme UBC3B induces its interaction with beta-TrCP and enhances beta-catenin degradation. <i>Oncogene</i> , 2002 , 21, 3978-87 | 9.2 | 46 |
| 143 | Substrate-specificity determinants for a membrane-bound casein kinase of lactating mammary gland. A study with synthetic peptides. <i>FEBS Journal</i> , 1988 , 177, 281-4 | | 46 |
| 142 | Phosphorylation of rat heart ornithine decarboxylase by type-2 casein kinase. <i>Biochemical and Biophysical Research Communications</i> , 1984 , 122, 997-1004 | 3.4 | 44 |
| 141 | Tyrosine phosphorylation of protein kinase CK2 by Src-related tyrosine kinases correlates with increased catalytic activity. <i>Biochemical Journal</i> , 2003 , 372, 841-9 | 3.8 | 42 |
| 140 | Novel consensus sequence for the Golgi apparatus casein kinase, revealed using proline-rich protein-1 (PRP1)-derived peptide substrates. <i>Biochemical Journal</i> , 2000 , 351, 765-768 | 3.8 | 42 |

| 139 | Urolithin as a converging scaffold linking ellagic acid and coumarin analogues: design of potent protein kinase CK2 inhibitors. <i>ChemMedChem</i> , 2011 , 6, 2273-86 | 3.7 | 41 |
|-----|---|------|----|
| 138 | A structural insight into CK2 inhibition. <i>Molecular and Cellular Biochemistry</i> , 2008 , 316, 57-62 | 4.2 | 40 |
| 137 | Effects of the CK2 inhibitors CX-4945 and CX-5011 on drug-resistant cells. <i>PLoS ONE</i> , 2012 , 7, e49193 | 3.7 | 39 |
| 136 | GRP94 (endoplasmin) co-purifies with and is phosphorylated by Golgi apparatus casein kinase. <i>FEBS Letters</i> , 2000 , 471, 151-5 | 3.8 | 39 |
| 135 | Phosphorylation of HIV-1 Rev protein: implication of protein kinase CK2 and pro-directed kinases. <i>Biochemical and Biophysical Research Communications</i> , 1996 , 226, 547-54 | 3.4 | 39 |
| 134 | Synthetic fragments of beta-casein as model substrates for liver and mammary gland casein kinases. <i>FEBS Journal</i> , 1989 , 186, 459-64 | | 39 |
| 133 | Discrimination between the activity of protein kinase CK2 holoenzyme and its catalytic subunits. <i>FEBS Letters</i> , 2006 , 580, 3948-52 | 3.8 | 38 |
| 132 | Isolation from spleen of a 57-kDa protein substrate of the tyrosine kinase Lyn. Identification as a protein related to protein disulfide-isomerase and localisation of the phosphorylation sites. <i>FEBS Journal</i> , 1996 , 235, 18-25 | | 38 |
| 131 | Modulation of protein kinase CK2 activity by fragments of CFTR encompassing F508 may reflect functional links with cystic fibrosis pathogenesis. <i>Biochemistry</i> , 2008 , 47, 7925-36 | 3.2 | 37 |
| 130 | The regulatory beta subunit of protein kinase CK2 contributes to the recognition of the substrate consensus sequence. A study with an eIF2 beta-derived peptide. <i>Biochemistry</i> , 2008 , 47, 8317-25 | 3.2 | 37 |
| 129 | Unique features of HIV-1 Rev protein phosphorylation by protein kinase CK2 ('casein kinase-2'). <i>FEBS Letters</i> , 2000 , 481, 63-7 | 3.8 | 37 |
| 128 | Structure and properties of casein kinase-2 from Saccharomyces cerevisiae. A comparison with the liver enzyme. <i>FEBS Journal</i> , 1986 , 159, 31-8 | | 37 |
| 127 | Developmental phosphoproteomics identifies the kinase CK2 as a driver of Hedgehog signaling and a therapeutic target in medulloblastoma. <i>Science Signaling</i> , 2018 , 11, | 8.8 | 37 |
| 126 | Casein kinase 2 (CK2) phosphorylates the deubiquitylase OTUB1 at Ser16 to trigger its nuclear localization. <i>Science Signaling</i> , 2015 , 8, ra35 | 8.8 | 36 |
| 125 | Cell-permeable dual inhibitors of protein kinases CK2 and PIM-1: structural features and pharmacological potential. <i>Cellular and Molecular Life Sciences</i> , 2014 , 71, 3173-85 | 10.3 | 36 |
| 124 | Inhibition of protein kinase CK2 with the clinical-grade small ATP-competitive compound CX-4945 or by RNA interference unveils its role in acute myeloid leukemia cell survival, p53-dependent apoptosis and daunorubicin-induced cytotoxicity. <i>Journal of Hematology and Oncology</i> , 2013 , 6, 78 | 22.4 | 35 |
| 123 | Motif analysis of phosphosites discloses a potential prominent role of the Golgi casein kinase (GCK) in the generation of human plasma phospho-proteome. <i>Journal of Proteome Research</i> , 2010 , 9, 3335-8 | 5.6 | 35 |
| 122 | Structure-function analysis of yeast piD261/Bud32, an atypical protein kinase essential for normal cell life. <i>Biochemical Journal.</i> 2002 . 364. 457-63 | 3.8 | 35 |

| 121 | Assessment of CK2 constitutive activity in cancer cells. <i>Methods in Enzymology</i> , 2010 , 484, 495-514 | 1.7 | 34 | |
|-----|---|------|----|--|
| 120 | Identification of novel protein kinase CK1 delta (CK1delta) inhibitors through structure-based virtual screening. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008 , 18, 5672-5 | 2.9 | 34 | |
| 119 | Chemical dissection of the APC Repeat 3 multistep phosphorylation by the concerted action of protein kinases CK1 and GSK3. <i>Biochemistry</i> , 2007 , 46, 11902-10 | 3.2 | 34 | |
| 118 | Biochemical evidence that Saccharomyces cerevisiae YGR262c gene, required for normal growth, encodes a novel Ser/Thr-specific protein kinase. <i>FEBS Letters</i> , 1997 , 414, 171-5 | 3.8 | 33 | |
| 117 | Autophosphorylation at the regulatory beta subunit reflects the supramolecular organization of protein kinase CK2. <i>Molecular and Cellular Biochemistry</i> , 2005 , 274, 23-9 | 4.2 | 33 | |
| 116 | Autocatalytic tyrosine-phosphorylation of protein kinase CK2 and a subunits: implication of Tyr182. <i>Biochemical Journal</i> , 2001 , 357, 563-567 | 3.8 | 33 | |
| 115 | Structural features underlying the unusual mode of calmodulin phosphorylation by protein kinase CK2: A study with synthetic calmodulin fragments. <i>Biochemical and Biophysical Research Communications</i> , 1999 , 256, 442-6 | 3.4 | 33 | |
| 114 | Isolation and identification of two proto-oncogene products related to c-fgr and fyn in a tyrosine-protein-kinase fraction of rat spleen. <i>FEBS Journal</i> , 1993 , 216, 323-7 | | 33 | |
| 113 | Extracellular phosphorylation of C9 by protein kinase CK2 regulates complement-mediated lysis. <i>European Journal of Immunology</i> , 2005 , 35, 1939-48 | 6.1 | 32 | |
| 112 | Spleen protein tyrosine kinases TPK-IIB and CSK display different immunoreactivity and opposite specificities toward c-src-derived peptides. <i>FEBS Letters</i> , 1992 , 313, 291-4 | 3.8 | 32 | |
| 111 | Nanoencapsulated anti-CK2 small molecule drug or siRNA specifically targets malignant cancer but not benign cells. <i>Cancer Letters</i> , 2012 , 315, 48-58 | 9.9 | 31 | |
| 110 | Re-evaluation of protein kinase CK2 pleiotropy: new insights provided by a phosphoproteomics analysis of CK2 knockout cells. <i>Cellular and Molecular Life Sciences</i> , 2018 , 75, 2011-2026 | 10.3 | 31 | |
| 109 | Aberrant signalling by protein kinase CK2 in imatinib-resistant chronic myeloid leukaemia cells: biochemical evidence and therapeutic perspectives. <i>Molecular Oncology</i> , 2013 , 7, 1103-15 | 7.9 | 30 | |
| 108 | Mass spectrometry analysis of a protein kinase CK2beta subunit interactome isolated from mouse brain by affinity chromatography. <i>Journal of Proteome Research</i> , 2008 , 7, 990-1000 | 5.6 | 30 | |
| 107 | Functional homology between yeast piD261/Bud32 and human PRPK: both phosphorylate p53 and PRPK partially complements piD261/Bud32 deficiency. <i>FEBS Letters</i> , 2003 , 549, 63-6 | 3.8 | 30 | |
| 106 | Susceptibility of the prion protein to enzymic phosphorylation. <i>Biochemical and Biophysical Research Communications</i> , 2000 , 271, 337-41 | 3.4 | 30 | |
| 105 | Detection of phospho-sites generated by protein kinase CK2 in CFTR: mechanistic aspects of Thr1471 phosphorylation. <i>PLoS ONE</i> , 2013 , 8, e74232 | 3.7 | 30 | |
| 104 | Generation and quantitative proteomics analysis of CK2/IIIcells. <i>Scientific Reports</i> , 2017 , 7, 42409 | 4.9 | 29 | |

| 103 | A chemogenomic screening identifies CK2 as a target for pro-senescence therapy in PTEN-deficient tumours. <i>Nature Communications</i> , 2015 , 6, 7227 | 17.4 | 29 |
|-----|---|-------------------|----|
| 102 | Eukaryotic translation-initiation factor eIF2beta binds to protein kinase CK2: effects on CK2alpha activity. <i>Biochemical Journal</i> , 2003 , 375, 623-31 | 3.8 | 29 |
| 101 | The crystal structure of the complex of Zea mays alpha subunit with a fragment of human beta subunit provides the clue to the architecture of protein kinase CK2 holoenzyme. <i>FEBS Journal</i> , 2000 , 267, 5184-90 | | 29 |
| 100 | Altered protein kinase activities of lymphoid cells transformed by Abelson and Moloney leukemia viruses. <i>FEBS Letters</i> , 1986 , 206, 59-63 | 3.8 | 29 |
| 99 | Sequence specificity of C-terminal Src kinase (CSK)a comparison with Src-related kinases c-Fgr and Lyn. <i>FEBS Journal</i> , 1997 , 246, 433-9 | | 28 |
| 98 | Phosphorylation of calmodulin fragments by protein kinase CK2. Mechanistic aspects and structural consequences. <i>Biochemistry</i> , 2004 , 43, 12788-98 | 3.2 | 28 |
| 97 | Comparative analysis of CK2 expression and function in tumor cell lines displaying sensitivity vs. resistance to chemical induced apoptosis. <i>Molecular and Cellular Biochemistry</i> , 2008 , 316, 155-61 | 4.2 | 27 |
| 96 | Phosphorylation and activation of protein kinase CK2 by p34cdc2 are independent events. <i>FEBS Journal</i> , 1995 , 230, 1025-31 | | 27 |
| 95 | Isoform specific phosphorylation of p53 by protein kinase CK1. <i>Cellular and Molecular Life Sciences</i> , 2010 , 67, 1105-18 | 10.3 | 26 |
| 94 | Phosphotyrosine as a specificity determinant for casein kinase-2, a growth related Ser/Thr-specific protein kinase. <i>FEBS Letters</i> , 1991 , 279, 307-9 | 3.8 | 26 |
| 93 | Superiority of PLK-2 as Esynuclein phosphorylating agent relies on unique specificity determinants. <i>Biochemical and Biophysical Research Communications</i> , 2012 , 418, 156-60 | 3.4 | 25 |
| 92 | Efficient Fmoc/solid-phase peptide synthesis of O-phosphotyrosyl-containing peptides and their use as phosphatase substrates. <i>International Journal of Peptide and Protein Research</i> , 1994 , 43, 39-46 | | 25 |
| 91 | Programmed cell death protein 5 (PDCD5) is phosphorylated by CK2 in vitro and in 293T cells. <i>Biochemical and Biophysical Research Communications</i> , 2009 , 387, 606-10 | 3.4 | 25 |
| 90 | Hematopoietic lineage cell specific protein 1 associates with and down-regulates protein kinase CK2. <i>FEBS Letters</i> , 1999 , 461, 32-6 | 3.8 | 25 |
| 89 | Structural features underlying the selectivity of the kinase inhibitors NBC and dNBC: role of a nitro group that discriminates between CK2 and DYRK1A. <i>Cellular and Molecular Life Sciences</i> , 2012 , 69, 449- | 60 ^{0.3} | 24 |
| 88 | Exploring the CK2 Paradox: Restless, Dangerous, Dispensable. <i>Pharmaceuticals</i> , 2017 , 10, | 5.2 | 24 |
| 87 | Design, validation and efficacy of bisubstrate inhibitors specifically affecting ecto-CK2 kinase activity. <i>Biochemical Journal</i> , 2015 , 471, 415-30 | 3.8 | 24 |
| 86 | Generation of protein kinase Ck1alpha mutants which discriminate between canonical and non-canonical substrates. <i>Biochemical Journal</i> , 2005 , 391, 417-24 | 3.8 | 24 |

| 85 | Protein kinase CK2 phosphorylates BAD at threonine-117. Neurochemistry International, 2004, 45, 747-5 | 524.4 | 23 |
|----------------------|---|-------|----------------------|
| 84 | Differential phosphorylation of Akt1 and Akt2 by protein kinase CK2 may account for isoform specific functions. <i>Biochimica Et Biophysica Acta - Molecular Cell Research</i> , 2014 , 1843, 1865-74 | 4.9 | 22 |
| 83 | Cystic fibrosis transmembrane regulator fragments with the Phe508 deletion exert a dual allosteric control over the master kinase CK2. <i>Biochemical Journal</i> , 2010 , 426, 19-29 | 3.8 | 22 |
| 82 | Golgi apparatus casein kinase phosphorylates bioactive Ser-6 of bone morphogenetic protein 15 and growth and differentiation factor 9. <i>FEBS Letters</i> , 2010 , 584, 801-5 | 3.8 | 22 |
| 81 | Sic1 is phosphorylated by CK2 on Ser201 in budding yeast cells. <i>Biochemical and Biophysical Research Communications</i> , 2006 , 346, 786-93 | 3.4 | 22 |
| 80 | Protein kinase CK2 phosphorylates the cell cycle regulatory protein Geminin. <i>Biochemical and Biophysical Research Communications</i> , 2004 , 315, 1011-7 | 3.4 | 22 |
| 79 | Phosphorylation and activation of the atypical kinase p53-related protein kinase (PRPK) by Akt/PKB. <i>Cellular and Molecular Life Sciences</i> , 2007 , 64, 2680-9 | 10.3 | 21 |
| 78 | Targeting Protein Kinase CK2: Evaluating CX-4945 Potential for GL261 Glioblastoma Therapy in Immunocompetent Mice. <i>Pharmaceuticals</i> , 2017 , 10, | 5.2 | 20 |
| 77 | Protein kinase CK2 accumulation in "oncophilic" cells: causes and effects. <i>Molecular and Cellular Biochemistry</i> , 2011 , 356, 5-10 | 4.2 | 19 |
| 76 | From phosphoproteins to phosphoproteomes: a historical account. <i>FEBS Journal</i> , 2017 , 284, 1936-1951 | 5.7 | 18 |
| | | | |
| 75 | The generation of phosphoserine stretches in phosphoproteins: mechanism and significance. <i>Molecular BioSystems</i> , 2015 , 11, 2666-79 | | 18 |
| 75 74 | | 3 | 18 |
| | Molecular BioSystems, 2015 , 11, 2666-79 The Selectivity of CK2 Inhibitor Quinalizarin: A Reevaluation. <i>BioMed Research International</i> , 2015 , | 3 | |
| 74 | Molecular BioSystems, 2015, 11, 2666-79 The Selectivity of CK2 Inhibitor Quinalizarin: A Reevaluation. BioMed Research International, 2015, 2015, 734127 Variable contribution of protein kinases to the generation of the human phosphoproteome: a | | 18 |
| 74 73 | Molecular BioSystems, 2015, 11, 2666-79 The Selectivity of CK2 Inhibitor Quinalizarin: A Reevaluation. BioMed Research International, 2015, 2015, 734127 Variable contribution of protein kinases to the generation of the human phosphoproteome: a global weblogo analysis. Biomolecular Concepts, 2010, 1, 185-95 Ser/Thr phosphorylation of hematopoietic specific protein 1 (HS1): implication of protein kinase | | 18 |
| 74 73 72 | Molecular BioSystems, 2015, 11, 2666-79 The Selectivity of CK2 Inhibitor Quinalizarin: A Reevaluation. BioMed Research International, 2015, 2015, 734127 Variable contribution of protein kinases to the generation of the human phosphoproteome: a global weblogo analysis. Biomolecular Concepts, 2010, 1, 185-95 Ser/Thr phosphorylation of hematopoietic specific protein 1 (HS1): implication of protein kinase CK2. FEBS Journal, 2000, 267, 3065-72 Synthesis and properties of a selective inhibitor of homeodomain-interacting protein kinase 2 | 3.7 | 18 17 17 |
| 74 73 72 71 | Molecular BioSystems, 2015, 11, 2666-79 The Selectivity of CK2 Inhibitor Quinalizarin: A Reevaluation. BioMed Research International, 2015, 2015, 734127 Variable contribution of protein kinases to the generation of the human phosphoproteome: a global weblogo analysis. Biomolecular Concepts, 2010, 1, 185-95 Ser/Thr phosphorylation of hematopoietic specific protein 1 (HS1): implication of protein kinase CK2. FEBS Journal, 2000, 267, 3065-72 Synthesis and properties of a selective inhibitor of homeodomain-interacting protein kinase 2 (HIPK2). PLoS ONE, 2014, 9, e89176 The pleiotropic protein kinase CK2 phosphorylates HTLV-1 Tax protein in vitro, targeting its | 3.7 | 18 17 17 17 |

| 67 | Exploiting the repertoire of CK2 inhibitors to target DYRK and PIM kinases. <i>Biochimica Et Biophysica Acta - Proteins and Proteomics</i> , 2013 , 1834, 1402-9 | 4 | 15 |
|----|--|------|----|
| 66 | Phosphorylation of the Saccharomyces cerevisiae Grx4p glutaredoxin by the Bud32p kinase unveils a novel signaling pathway involving Sch9p, a yeast member of the Akt / PKB subfamily. <i>FEBS Journal</i> , 2008 , 275, 5919-33 | 5.7 | 14 |
| 65 | Heterogeneity of CK2 phosphorylation sites in the NS5A protein of different hepatitis C virus genotypes. <i>Journal of Hepatology</i> , 2007 , 47, 768-76 | 13.4 | 14 |
| 64 | CK2 regulates in vitro the activity of the yeast cyclin-dependent kinase inhibitor Sic1. <i>Biochemical and Biophysical Research Communications</i> , 2005 , 336, 1040-8 | 3.4 | 14 |
| 63 | Inhibition of protein kinase CK2 by CX-5011 counteracts imatinib-resistance preventing rpS6 phosphorylation in chronic myeloid leukaemia cells: new combined therapeutic strategies. <i>Oncotarget</i> , 2016 , 7, 18204-18 | 3.3 | 14 |
| 62 | CFTR mutations altering CFTR fragmentation. <i>Biochemical Journal</i> , 2013 , 449, 295-305 | 3.8 | 13 |
| 61 | Linear and cyclic synthetic peptides related to the main autophosphorylation site of the Src tyrosine kinases as substrates and inhibitors of Lyn. <i>International Journal of Peptide and Protein Research</i> , 1995 , 45, 529-39 | | 13 |
| 60 | Protein Kinase CK2 Subunits Differentially Perturb the Adhesion and Migration of GN11 Cells: A Model of Immature Migrating Neurons. <i>International Journal of Molecular Sciences</i> , 2019 , 20, | 6.3 | 13 |
| 59 | Proteomics perturbations promoted by the protein kinase CK2 inhibitor quinalizarin. <i>Biochimica Et Biophysica Acta - Proteins and Proteomics</i> , 2015 , 1854, 1676-86 | 4 | 12 |
| 58 | Chimeric peptides as modulators of CK2-dependent signaling: Mechanism of action and off-target effects. <i>Biochimica Et Biophysica Acta - Proteins and Proteomics</i> , 2015 , 1854, 1694-707 | 4 | 12 |
| 57 | A new role for sphingosine: Up-regulation of Fam20C, the genuine casein kinase that phosphorylates secreted proteins. <i>Biochimica Et Biophysica Acta - Proteins and Proteomics</i> , 2015 , 1854, 1718-26 | 4 | 12 |
| 56 | Acidophilic character of yeast PID261/BUD32, a putative ancestor of eukaryotic protein kinases. <i>Biochemical and Biophysical Research Communications</i> , 2002 , 296, 1366-71 | 3.4 | 12 |
| 55 | Pharmacophore-guided discovery of CDC25 inhibitors causing cell cycle arrest and tumor regression. <i>Scientific Reports</i> , 2019 , 9, 1335 | 4.9 | 11 |
| 54 | CK2 involvement in ESCRT-III complex phosphorylation. <i>Archives of Biochemistry and Biophysics</i> , 2014 , 545, 83-91 | 4.1 | 11 |
| 53 | An exploration of the effects of constraints on the phosphorylation of synthetic protein tyrosine kinase peptide substrates. <i>Journal of Peptide Science</i> , 1996 , 2, 325-38 | 2.1 | 11 |
| 52 | Phosphorylation of ornithine decarboxylase in intact erythroleukemia cells. <i>Biochimica Et Biophysica Acta - Molecular Cell Research</i> , 1990 , 1052, 345-7 | 4.9 | 11 |
| 51 | The p23 co-chaperone protein is a novel substrate of CK2 in Arabidopsis. <i>Molecular and Cellular Biochemistry</i> , 2011 , 356, 245-54 | 4.2 | 10 |
| 50 | Understanding protein kinase CK2 mis-regulation upon F508del CFTR expression. Naunyn-Schmiedebergrs Archives of Pharmacology, 2011 , 384, 473-88 | 3.4 | 10 |

(2017-2003)

| 49 | Conformational constraints of tyrosine in protein tyrosine kinase substrates: Information about preferred bioactive side-chain orientation. <i>Biopolymers</i> , 2003 , 71, 478-88 | 2.2 | 10 |
|----|---|---------|----|
| 48 | Specific monitoring of Syk protein kinase activity by peptide substrates including constrained analogs of tyrosine. <i>FEBS Letters</i> , 2002 , 523, 48-52 | 3.8 | 10 |
| 47 | A Comparative Analysis and Review of lysyl Residues Affected by Posttranslational Modifications. <i>Current Genomics</i> , 2015 , 16, 128-38 | 2.6 | 10 |
| 46 | The Golgi 'casein kinase' Fam20C is a genuine 'phosvitin kinase' and phosphorylates polyserine stretches devoid of the canonical consensus. <i>FEBS Journal</i> , 2018 , 285, 4674-4683 | 5.7 | 10 |
| 45 | Protein kinase CK2 potentiates translation efficiency by phosphorylating eIF3j at Ser127. <i>Biochimica Et Biophysica Acta - Molecular Cell Research</i> , 2015 , 1853, 1693-701 | 4.9 | 9 |
| 44 | Prevalence and significance of the commonest phosphorylated motifs in the human proteome: a global analysis. <i>Cellular and Molecular Life Sciences</i> , 2020 , 77, 5281-5298 | 10.3 | 9 |
| 43 | "Janus" efficacy of CX-5011: CK2 inhibition and methuosis induction by independent mechanisms. <i>Biochimica Et Biophysica Acta - Molecular Cell Research</i> , 2020 , 1867, 118807 | 4.9 | 9 |
| 42 | Dependence of HSP27 cellular level on protein kinase CK2 discloses novel therapeutic strategies. <i>Biochimica Et Biophysica Acta - General Subjects</i> , 2018 , 1862, 2902-2910 | 4 | 9 |
| 41 | A proteomics analysis of CK2IC2C12 cells provides novel insights into the biological functions of the non-catalytic is ubunit. <i>FEBS Journal</i> , 2019 , 286, 1561-1575 | 5.7 | 8 |
| 40 | Different Persistence of the Cellular Effects Promoted by Protein Kinase CK2 Inhibitors CX-4945 and TDB. <i>BioMed Research International</i> , 2015 , 2015, 185736 | 3 | 8 |
| 39 | Spatial conformation and topography of the tyrosine aromatic ring in substrate recognition by protein tyrosine kinases. <i>Journal of Medicinal Chemistry</i> , 2006 , 49, 1916-24 | 8.3 | 8 |
| 38 | Fam20C is under the control of sphingolipid signaling in human cell lines. FEBS Journal, 2017, 284, 1246 | -3;2,57 | 7 |
| 37 | A "SYDE" effect of hierarchical phosphorylation: possible relevance to the cystic fibrosis basic defect. <i>Cellular and Molecular Life Sciences</i> , 2014 , 71, 2193-6 | 10.3 | 7 |
| 36 | Phosphorylation of phosvitin by casein kinase-2 provides the evidence that phosphoserines can replace carboxylic amino acids as specificity determinants. <i>Biochimica Et Biophysica Acta - Bioenergetics</i> , 1988 , 971, 227-231 | 4.6 | 7 |
| 35 | Under-expression of CK2Isubunit in ccRCC represents a complementary biomarker of p-STAT3 Ser727 that correlates with patient survival. <i>Oncotarget</i> , 2018 , 9, 5736-5751 | 3.3 | 7 |
| 34 | The Interactome of Protein Kinase CK276-116 | | 7 |
| 33 | Identification of the PLK2-dependent phosphopeptidome by quantitative proteomics [corrected]. <i>PLoS ONE</i> , 2014 , 9, e111018 | 3.7 | 6 |
| 32 | Protein kinase CK2 modulates HSJ1 function through phosphorylation of the UIM2 domain. <i>Human Molecular Genetics</i> , 2017 , 26, 611-623 | 5.6 | 6 |

| 31 | CK2 Suppression of Apoptosis and Its Implication in Cancer Biology and Therapy319-343 | | 6 |
|----|--|---------------|---|
| 30 | CK2 Contribution to the Generation of the Human Phosphoproteome117-128 | | 6 |
| 29 | Cross talk between protein kinase CK2 and eukaryotic translation initiation factor eIF2beta subunit. <i>Molecular and Cellular Biochemistry</i> , 2005 , 274, 53-61 | 4.2 | 5 |
| 28 | Targeting CK2 in cancer: a valuable strategy or a waste of time?. <i>Cell Death Discovery</i> , 2021 , 7, 325 | 6.9 | 5 |
| 27 | A N-terminally deleted form of the CK2[catalytic subunit is sufficient to support cell viability. <i>Biochemical and Biophysical Research Communications</i> , 2020 , 531, 409-415 | 3.4 | 5 |
| 26 | How can a traffic light properly work if it is always green? The paradox of CK2 signaling. <i>Critical Reviews in Biochemistry and Molecular Biology</i> , 2021 , 56, 321-359 | 8.7 | 5 |
| 25 | Comparing the efficacy and selectivity of Ck2 inhibitors. A phosphoproteomics approach. <i>European Journal of Medicinal Chemistry</i> , 2021 , 214, 113217 | 6.8 | 5 |
| 24 | CK2 in Embryonic Development129-168 | | 5 |
| 23 | Synthetic Tyr-phospho and non-hydrolyzable phosphonopeptides as PTKs and TC-PTP inhibitors. <i>International Journal of Peptide and Protein Research</i> , 1995 , 46, 535-46 | | 4 |
| 22 | Deciphering the role of protein kinase CK2 in the maturation/stability of F508del-CFTR. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2020 , 1866, 165611 | 6.9 | 4 |
| 21 | Up-Regulation of the Alpha Prime Subunit of Protein Kinase CK2 as a Marker of Fast Proliferation in GL261 Cultured Cells. <i>Pathology and Oncology Research</i> , 2019 , 25, 1659-1663 | 2.6 | 4 |
| 20 | Contribution of the CK2 Catalytic Isoforms and ato the Glycolytic Phenotype of Tumor Cells. <i>Cells</i> , 2021 , 10, | 7.9 | 4 |
| 19 | Specific Features of Plant CK2 2013 , 267-289 | | 3 |
| 18 | Effects of CK2lbubunit down-regulation on Akt signalling in HK-2 renal cells. PLoS ONE, 2020 , 15, e0227 | 3 <u>4</u> ,θ | 3 |
| 17 | Structural Bases of Protein Kinase CK2 Function and Inhibition1-75 | | 3 |
| 16 | Linear and cyclic peptides as substrates for Lyn tyrosine kinase. <i>Journal of Peptide Science</i> , 1998 , 4, 33-4 | 52.1 | 2 |
| 15 | ©enuine Casein Kinase: The False Sister of CK2 That Phosphorylates Secreted Proteins at S-x-E/pS Motifs 2015 , 227-237 | | 2 |
| 14 | Genuine Casein Kinase (Fam 20C): The Mother of the Phosphosecretome 47-62 | | 2 |

LIST OF PUBLICATIONS

| 13 | Synthesis and biological activities of cyclic lactam peptides as substrates for non-receptors PTKs. <i>International Journal of Peptide Research and Therapeutics</i> , 1999 , 6, 117-121 | | 1 |
|----|--|-----|---|
| 12 | Direct detection of protein kinases on electropherograms. FEBS Letters, 1975, 51, 130-2 | 3.8 | 1 |
| 11 | Multiple Myeloma Cells Survival and Proliferation Rely on High Levels and Activity of the Serine-Threonine Kinase CK2 <i>Blood</i> , 2004 , 104, 643-643 | 2.2 | 1 |
| 10 | The importance of negative determinants as modulators of CK2 targeting. The lesson of Akt2 S131. <i>PLoS ONE</i> , 2018 , 13, e0193479 | 3.7 | 1 |
| 9 | Addiction of Cancer Cells to CK2: Survival at All Costs or Achilles' Heel?305-318 | | 1 |
| 8 | Protein Kinase CK2 in Normal and Malignant Hematopoiesis344-362 | | 1 |
| 7 | CK2: A Global Regulator of Cell Survival239-266 | | 1 |
| 6 | The Pivotal Role of CK2 in the Kinome-Targeting Hsp90 Chaperone Machinery205-238 | | 1 |
| 5 | CK2 as a Logical Target in Cancer Therapy: Potential for Combining CK2 Inhibitors with Various Classes of Cancer Therapeutic Agents383-439 | | 1 |
| 4 | IPK2019: David Shugar and the genesis of the IPK conferences. IUBMB Life, 2020, 72, 1097-1102 | 4.7 | |
| 3 | Synthesis and biological activities of cyclic lactam peptides as substrates for non-receptors PTKs. <i>International Journal of Peptide Research and Therapeutics</i> , 1999 , 6, 117-121 | | |
| 2 | 1954-2006: the long march of protein kinase CK2. <i>FASEB Journal</i> , 2006 , 20, A499 | 0.9 | |

The Role of Protein Kinase CK2 in the p53 Response190-204