

# Jacques Teulon

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/9059484/publications.pdf>

Version: 2024-02-01

8  
papers

247  
citations

1478505

6  
h-index

1720034

7  
g-index

8  
all docs

8  
docs citations

8  
times ranked

248  
citing authors

#	ARTICLE	IF	CITATIONS
1	Analysis of <i>CLCNKB</i> mutations at dimer interface, calcium binding site, and pore reveals a variety of functional alterations in ClC-Kb channel leading to Bartter syndrome. <i>Human Mutation</i> , 2020, 41, 774-785.	2.5	6
2	Renal Chloride Channels in Relation to Sodium Chloride Transport. , 2018, 9, 301-342.		12
3	The ClC-K2 Chloride Channel Is Critical for Salt Handling in the Distal Nephron. <i>Journal of the American Society of Nephrology: JASN</i> , 2017, 28, 209-217.	6.1	87
4	In silico model of the human ClC-Kb chloride channel: pore mapping, biostructural pathology and drug screening. <i>Scientific Reports</i> , 2017, 7, 7249.	3.3	15
5	ClC-K chloride channels: emerging pathophysiology of Bartter syndrome type 3. <i>American Journal of Physiology - Renal Physiology</i> , 2015, 308, F1324-F1334.	2.7	52
6	CLCNKB mutations causing mild Bartter syndrome profoundly alter the pH and Ca <sup>2+</sup> dependence of ClC-Kb channels. <i>Pflügers Archiv European Journal of Physiology</i> , 2014, 466, 1713-1723.	2.8	23
7	Characterization of the mouse ClC-K1/Barttin chloride channel. <i>Biochimica Et Biophysica Acta - Biomembranes</i> , 2013, 1828, 2399-2409.	2.6	25
8	Novel <i>CLCNKB</i> Mutations Causing Bartter Syndrome Affect Channel Surface Expression. <i>Human Mutation</i> , 2013, 34, 1269-1278.	2.5	27