

# Deborah K Hanson

## List of Publications by Year in descending order

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23  
papers

549  
citations

687363

13  
h-index

642732

23  
g-index

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all docs

23  
docs citations

23  
times ranked

362  
citing authors

#	ARTICLE	IF	CITATIONS
1	In Situ, Protein-Mediated Generation of a Photochemically Active Chlorophyll Analogue in a Mutant Bacterial Photosynthetic Reaction Center. <i>Biochemistry</i> , 2021, 60, 1260-1275.	2.5	1
2	Switching sides—Reengineered primary charge separation in the bacterial photosynthetic reaction center. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2020, 117, 865-871.	7.1	11
3	Consequences of saturation mutagenesis of the protein ligand to the B-side monomeric bacteriochlorophyll in reaction centers from <i>Rhodobacter capsulatus</i> . <i>Photosynthesis Research</i> , 2019, 141, 273-290.	2.9	5
4	Manipulating the Energetics and Rates of Electron Transfer in <i>Rhodobacter capsulatus</i> Reaction Centers with Asymmetric Pigment Content. <i>Journal of Physical Chemistry B</i> , 2017, 121, 6989-7004.	2.6	15
5	Species differences in unlocking B-side electron transfer in bacterial reaction centers. <i>FEBS Letters</i> , 2016, 590, 2515-2526.	2.8	8
6	Optimizing multi-step B-side charge separation in photosynthetic reaction centers from <i>Rhodobacter capsulatus</i> . <i>Biochimica Et Biophysica Acta - Bioenergetics</i> , 2016, 1857, 150-159.	1.0	8
7	High yield of secondary B-side electron transfer in mutant <i>Rhodobacter capsulatus</i> reaction centers. <i>Biochimica Et Biophysica Acta - Bioenergetics</i> , 2014, 1837, 1892-1903.	1.0	10
8	High Throughput Engineering to Revitalize a Vestigial Electron Transfer Pathway in Bacterial Photosynthetic Reaction Centers. <i>Journal of Biological Chemistry</i> , 2012, 287, 8507-8514.	3.4	11
9	Determination of the Rate and Yield of B-side Quinone Reduction in <i>Rhodobacter capsulatus</i> Reaction Centers. <i>Biochemistry</i> , 2006, 45, 7314-7322.	2.5	20
10	B-Side Electron Transfer To Form P+HB- in Reaction Centers from the F(L181)Y/Y(M208)F Mutant of <i>Rhodobacter capsulatus</i> . <i>Journal of Physical Chemistry B</i> , 2004, 108, 11827-11832.	2.6	24
11	Lysine substitutions near photoactive cofactors in the bacterial photosynthetic reaction center have opposite effects on the rate of triplet energy transfer. <i>Chemical Physics</i> , 2003, 294, 329-346.	1.9	5
12	Detergent effects on primary charge separation in wild-type and mutant <i>Rhodobacter capsulatus</i> reaction centers. <i>Chemical Physics</i> , 2003, 294, 305-318.	1.9	22
13	Quinone Reduction via Secondary B-Branch Electron Transfer in Mutant Bacterial Reaction Centers. <i>Biochemistry</i> , 2003, 42, 1718-1730.	2.5	71
14	B-Side Charge Separation in Bacterial Photosynthetic Reaction Centers: A Nanosecond Time Scale Electron Transfer from HB-to QB. <i>Biochemistry</i> , 2003, 42, 2016-2024.	2.5	41
15	Comparison of M-Side Electron Transfer in <i>Rb. sphaeroides</i> and <i>Rb. capsulatus</i> Reaction Centers. <i>Journal of Physical Chemistry B</i> , 2002, 106, 1799-1808.	2.6	58
16	Title is missing!. <i>Photosynthesis Research</i> , 1998, 55, 275-280.	2.9	7
17	Title is missing!. <i>Photosynthesis Research</i> , 1998, 55, 267-273.	2.9	21
18	In Bacterial Reaction Centers, a Key Residue Suppresses Mutational Blockage of Two Different Proton Transfer Steps. <i>Biochemistry</i> , 1998, 37, 2077-2083.	2.5	14

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19	Antenna Excited State Decay Kinetics Establish Primary Electron Transfer in Reaction Centers as Heterogeneous. <i>Biochemistry</i> , 1997, 36, 8677-8685.	2.5	25
20	A native electrostatic environment near QB is not sufficient to ensure rapid proton delivery in photosynthetic reaction centers. <i>FEBS Letters</i> , 1997, 407, 159-163.	2.8	9
21	Title is missing!. <i>Photosynthesis Research</i> , 1997, 52, 93-103.	2.9	13
22	Long-range electrostatic interaction in the bacterial photosynthetic reaction centre. <i>Nature Structural and Molecular Biology</i> , 1995, 2, 1057-1059.	8.2	52
23	Recombinant immunoglobulin variable domains generated from synthetic genes provide a system for in vitro characterization of light-chain amyloid proteins. <i>Protein Science</i> , 1995, 4, 421-432.	7.6	98