

# Libby Wood

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/8760168/publications.pdf>

Version: 2024-02-01

9  
papers

238  
citations

1163117

8  
h-index

1372567

10  
g-index

11  
all docs

11  
docs citations

11  
times ranked

694  
citing authors

#	ARTICLE	IF	CITATIONS
1	Reproductive Cancer Risk Factors in Women With Myotonic Dystrophy (DM): Survey Data From the US and UK DM Registries. <i>Frontiers in Neurology</i> , 2019, 10, 1071.	2.4	5
2	RD-Connect, NeurOmics and EUREnOmics: collaborative European initiative for rare diseases. <i>European Journal of Human Genetics</i> , 2018, 26, 778-785.	2.8	55
3	Benign and malignant tumors in the UK myotonic dystrophy patient registry. <i>Muscle and Nerve</i> , 2018, 57, 316-320.	2.2	15
4	Chronic pain has a strong impact on quality of life in facioscapulohumeral muscular dystrophy. <i>Muscle and Nerve</i> , 2018, 57, 380-387.	2.2	33
5	Eight years after an international workshop on myotonic dystrophy patient registries: case study of a global collaboration for a rare disease. <i>Orphanet Journal of Rare Diseases</i> , 2018, 13, 155.	2.7	19
6	The UK Myotonic Dystrophy Patient Registry: facilitating and accelerating clinical research. <i>Journal of Neurology</i> , 2017, 264, 979-988.	3.6	23
7	Respiratory involvement in ambulant and non-ambulant patients with facioscapulohumeral muscular dystrophy. <i>Journal of Neurology</i> , 2017, 264, 1271-1280.	3.6	30
8	Linked Registries: Connecting Rare Diseases Patient Registries through a Semantic Web Layer. <i>BioMed Research International</i> , 2017, 2017, 1-13.	1.9	28
9	Design, set-up and utility of the UK facioscapulohumeral muscular dystrophy patient registry. <i>Journal of Neurology</i> , 2016, 263, 1401-1408.	3.6	25