

# Holly A Black

## List of Publications by Year in Descending Order

**Source:** <https://exaly.com/author-pdf/8736692/holly-a-black-publications-by-year.pdf>

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

11  
papers

84  
citations

5  
h-index

9  
g-index

11  
ext. papers

107  
ext. citations

6.7  
avg, IF

1.13  
L-index

| #  | Paper  | IF  | Citations |
|----|--|-----|-----------|
| 11 | Absolute measurement of the tissue origins of cell-free DNA in the healthy state and following paracetamol overdose. <i>BMC Medical Genomics</i> , <b>2020</b> , 13, 60                            | 3.7 | 6         |
| 10 | A sensitive and affordable multiplex RT-qPCR assay for SARS-CoV-2 detection. <i>PLoS Biology</i> , <b>2020</b> , 18, e3001030  | 9.7 | 17        |
| 9  | A sensitive and affordable multiplex RT-qPCR assay for SARS-CoV-2 detection <b>2020</b> , 18, e3001030   |     |           |
| 8  | A sensitive and affordable multiplex RT-qPCR assay for SARS-CoV-2 detection <b>2020</b> , 18, e3001030   |     |           |
| 7  | A sensitive and affordable multiplex RT-qPCR assay for SARS-CoV-2 detection <b>2020</b> , 18, e3001030   |     |           |
| 6  | A sensitive and affordable multiplex RT-qPCR assay for SARS-CoV-2 detection <b>2020</b> , 18, e3001030   |     |           |
| 5  | A sensitive and affordable multiplex RT-qPCR assay for SARS-CoV-2 detection <b>2020</b> , 18, e3001030   |     |           |
| 4  | A sensitive and affordable multiplex RT-qPCR assay for SARS-CoV-2 detection <b>2020</b> , 18, e3001030   |     |           |
| 3  | Genetic epidemiology of motor neuron disease-associated variants in the Scottish population. <i>Neurobiology of Aging</i> , <b>2017</b> , 51, 178.e11-178.e20                                      | 5.6 | 21        |
| 2  | De novo mutations in autosomal recessive congenital malformations. <i>Genetics in Medicine</i> , <b>2016</b> , 18, 1325-1326   | 8.1 | 5         |
| 1  | Targeted next-generation sequencing makes new molecular diagnoses and expands genotype-phenotype relationship in Ehlers-Danlos syndrome. <i>Genetics in Medicine</i> , <b>2016</b> , 18, 1119-1127 | 8.1 | 35        |