## James Hardwick

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/11571156/publications.pdf

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933447 1281871 3,262 11 10 11 citations h-index g-index papers 11 11 11 6196 docs citations times ranked citing authors all docs

| #  | Article  | IF   | CITATIONS |
|----|--|------|-----------|
| 1  | Comprehensive Characterization of Oncogenic Drivers in Asian Lung Adenocarcinoma. Journal of Thoracic Oncology, 2016, 11, 2129-2140.   | 1.1  | 70        |
| 2  | Molecular analysis of gastric cancer identifies subtypes associated with distinct clinical outcomes. Nature Medicine, 2015, 21, 449-456.   | 30.7 | 1,592     |
| 3  | Genome-wide identification of RNA editing in hepatocellular carcinoma. Genomics, 2015, 105, 76-82.   | 2.9  | 40        |
| 4  | Genomic landscape and genetic heterogeneity in gastric adenocarcinoma revealed by whole-genome sequencing. Nature Communications, 2014, 5, 5477.                                     | 12.8 | 166       |
| 5  | Whole-genome sequencing identifies recurrent mutations in hepatocellular carcinoma. Genome Research, 2013, 23, 1422-1433.  | 5.5  | 457       |
| 6  | Genome-wide survey of recurrent HBV integration in hepatocellular carcinoma. Nature Genetics, 2012, 44, 765-769.   | 21.4 | 785       |
| 7  | Pyridyl aminothiazoles as potent Chk1 inhibitors: Optimization of cellular activity. Bioorganic and Medicinal Chemistry Letters, 2012, 22, 2613-2619.                                | 2.2  | 8         |
| 8  | Optimization of a pyrazoloquinolinone class of Chk1 kinase inhibitors. Bioorganic and Medicinal Chemistry Letters, 2007, 17, 5989-5994.  | 2.2  | 42        |
| 9  | Synthesis and evaluation of substituted benzoisoquinolinones as potent inhibitors of Chk1 kinase. Bioorganic and Medicinal Chemistry Letters, 2007, 17, 6280-6285.                   | 2.2  | 24        |
| 10 | Development of 6-substituted indolylquinolinones as potent Chek1 kinase inhibitors. Bioorganic and Medicinal Chemistry Letters, 2006, 16, 5907-5912.                                 | 2.2  | 34        |
| 11 | 3-(Indol-2-yl)indazoles as Chek1 kinase inhibitors: Optimization of potency and selectivity via substitution at C6. Bioorganic and Medicinal Chemistry Letters, 2006, 16, 6049-6053. | 2.2  | 44        |