# Maryam Ehteshami

List of Publications by Year
in descending order



| 14 | 394 | 840776 11 | 1058476 14 |
|---|---|---|---|
| papers | citations | h-index | g-index |

| 14 | 14 | 14 | 750 |
|---|---|---|---|
| all docs | docs citations | times ranked | citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Chutes and ladders in hepatitis C nucleoside drug development. Antiviral Research, 2014, 102, 119-147. | 4.1 | 69 |
| 2 | Characterization of Î²- <scp>d</scp> - N <sup>4</sup> -Hydroxycytidine as a Novel Inhibitor of Chikungunya Virus. Antimicrobial Agents and Chemotherapy, 2017, 61, . | 3.2 | 64 |
| 3 | Towards <scp>HBV</scp> curative therapies. Liver International, 2018, 38, 102-114. | 3.9 | 63 |
| 4 | Approaches to hepatitis C treatment and cure using NS5A inhibitors. Infection and Drug Resistance, 2014, 7, 41. | 2.7 | 51 |
| 5 | Î²-<scp>d</scp>-2â€²-C-Methyl-2,6-diaminopurine Ribonucleoside Phosphoramidates are Potent and Selective Inhibitors of Hepatitis C Virus (HCV) and Are Bioconverted Intracellularly to Bioactive 2,6-Diaminopurine and Guanosine 5â€²-Triphosphate Forms. Journal of Medicinal Chemistry, 2015, 58, 3445-3458. | 6.4 | 30 |
| 6 | Toward Elimination of Hepatitis B Virus Using Novel Drugs, Approaches, and Combined Modalities. Clinics in Liver Disease, 2016, 20, 737-749. | 2.1 | 24 |
| 7 | 2â€²-Chloro,2â€²-fluoro Ribonucleotide Prodrugs with Potent Pan-genotypic Activity against Hepatitis C Virus Replication in Culture. Journal of Medicinal Chemistry, 2017, 60, 5424-5437. | 6.4 | 23 |
| 8 | Synthesis and Evaluation of 2,6-Modified Purine 2â€²-C-Methyl Ribonucleosides as Inhibitors of HCV Replication. ACS Medicinal Chemistry Letters, 2016, 7, 17-22. | 2.8 | 16 |
| 9 | Discovery, characterization, and lead optimization of 7-azaindole non-nucleoside HIV-1 reverse transcriptase inhibitors. Bioorganic and Medicinal Chemistry Letters, 2016, 26, 4101-4105. | 2.2 | 13 |
| 10 | Biochemical Characterization of the Active Anti-Hepatitis C Virus Metabolites of 2,6-Diaminopurine Ribonucleoside Prodrug Compared to Sofosbuvir and BMS-986094. Antimicrobial Agents and Chemotherapy, 2016, 60, 4659-4669. | 3.2 | 11 |
| 11 | Discovery of a Series of 2â€²-Î±-Fluoro,2â€²-Î²-bromo-ribonucleosides and Their Phosphoramidate Prodrugs as Potent Pan-Genotypic Inhibitors of Hepatitis C Virus. Journal of Medicinal Chemistry, 2019, 62, 1859-1874. | 6.4 | 11 |
| 12 | Nucleotide Substrate Specificity of Anti-Hepatitis C Virus Nucleoside Analogs for Human Mitochondrial RNA Polymerase. Antimicrobial Agents and Chemotherapy, 2017, 61, . | 3.2 | 8 |
| 13 | Metabolism of Nucleosides and Nucleotides Prodrugs. Current Pharmaceutical Design, 2018, 23, 6984-7002. | 1.9 | 8 |
| 14 | Intracellular metabolism and potential cardiotoxicity of a Î²-D-2â€™-C-methyl-2,6-diaminopurine ribonucleoside phosphoramidate that inhibits hepatitis C virus replication. Nucleosides, Nucleotides and Nucleic Acids, 2020, 39, 204-224. | 1.1 | 3 |