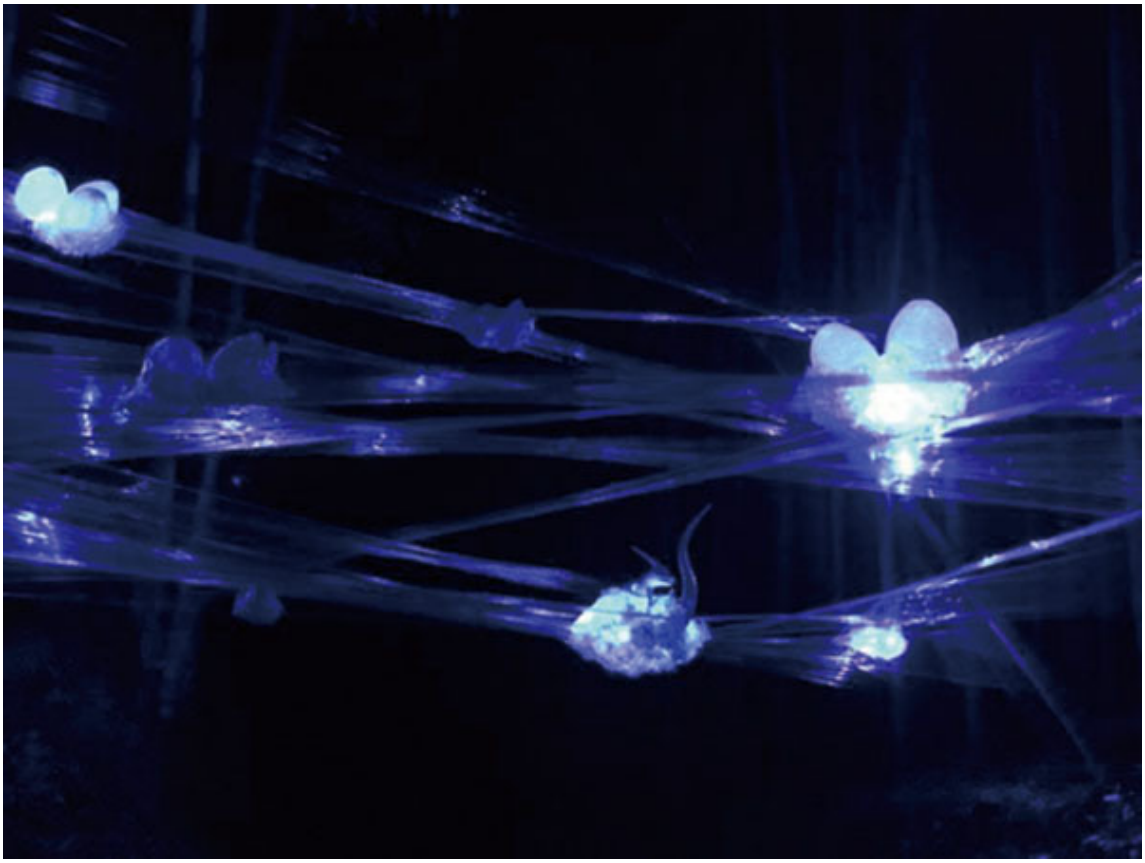
64bit Word Snatcher 1.19 Cracked Full Version Registration



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We provided one example of each compound type. Although few examples are cited in [Figure 2](#fig2){ref-type="fig"}(B), the amount of examples is doubled to provide a comprehensive overview. We chose a small number of compounds for illustration of compound types and of the different code points in a compound in order to keep the text easy to read. The keywords were placed in alphabetical order, and compound types were grouped alphabetically according to the prefix of the compound type. The entry for each compound type is provided in the Supplementary Materials. Biological evaluation of compounds against influenza A virus replication {#sec5}

========== We tested the activity of 21 compounds selected from the screening of [Figure 2](#fig2){ref-type="fig"} against influenza A virus replication in a standard fluorescence-based plaque reduction assay. For each compound we chose concentrations at which compound and virus were present in equal volumes, a concentration that is comparable to the concentration of 4 μM used for the screening. The same assay conditions and assay workflow were used in the screening. We used three batches of A/Vietnam/1203/2004 (H5N1) virus, three of A/duck/Malaysia/5838/2004 (H5N1) virus, and one of A/duck/Hokkaido/172/2008 (H1N1) virus ([@bib7]). Each of the six virus isolates were cloned, plaque-purified, and then expanded in embryonated eggs. Subsequently, the allantoic fluid was harvested and stored in aliquots at --80 °C. The 50% tissue culture infectious dose (TCID~50~) per milliliter of virus in the aliquots was determined by titration of serial dilutions of virus on chicken eggs. The time from clonal isolation until harvesting of aliquots was 7--10 days. Compounds were added to MDCK cells before and after virus inoculation. Before addition of compounds, MDCK cells were washed with PBS and incubated in serum-free Dulbecco's modified Eagle medium (DMEM) for 1 h. After removal of the medium, the cells were incubated for 1 h with or without the compound at the indicated concentrations. The compound was removed, DMEM containing the virus was added, and the cells were incubated for 1 h
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