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## 白杨素衍生物的合成及其抗肿瘤活性

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**中文摘要:**利用Baker-Venkataraman重排法,以2,4,6-三羟基苯乙酮为原料,经羟基保护、酰化、重排、成环、脱保护5步反应全合成白杨素。以白杨素为骨架,再利用7位羟基的活性,在C-7位引入水溶性基团——烷胺基,设计合成了未见文献报道的21种白杨素系列衍生物**8a-8u**,其结构经<sup>1</sup>H NMR、<sup>13</sup>C NMR、IR和MS确证。采用MTT法评价了目标化合物对HCT-116(人体结肠癌细胞系),HeLa(人体宫颈癌细胞系),DU-145(人体前列腺癌细胞系),SGC-7901(人体胃癌细胞系)和HEK-293(人胚肾细胞系)的抗肿瘤活性。其体外抗肿瘤活性实验表明,**7-(2-哌嗪乙氧基)-5-羟基-2-苯基-4H-1-苯并吡喃-4-酮(8o)**具有较好的抗肿瘤活性。

**中文关键词:**[白杨素](#) [衍生物](#) [合成](#) [抗肿瘤活性](#) [MTT检测](#)

## Synthesis and antitumor activity of novel chrysanthemum derivatives

**Abstract:**Chrysanthemum was synthesized in five steps starting from 2,4,6-trihydroxyacetophenone via methylation, acylation, Baker-Venkataraman rearrangement, cyclization and deprotection. With chrysanthemum as the lead compound, novel chrysanthemum derivatives **8a-8u** were designed and synthesized. The chemical structures of these compounds were confirmed by <sup>1</sup>H NMR, <sup>13</sup>C NMR, IR and MS. Twenty-one new chrysanthemum derivatives were prepared and their antitumor activities were evaluated by MTT method on HCT-116 (human colon cancer cell line), HeLa (human cervical carcinoma cell line), DU-145 (human prostate cell line), SGC-7901 (human gastric cancer cell line), and HEK-293 (human embryonic kidney cell line). Among these derivatives, 5-hydroxy-2-phenyl-7-(2-(piperazin-1-yl) ethoxy)-4H-chromen-4-one (**8o**) had the most potent antitumor activity against HCT-116, HeLa, DU-145, and SGC-7901 cells.

**keywords:**[Chrysanthemum](#) [derivatives](#) [synthesis](#) [antitumor activities](#) [MTT assay](#)

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