

MH P 2

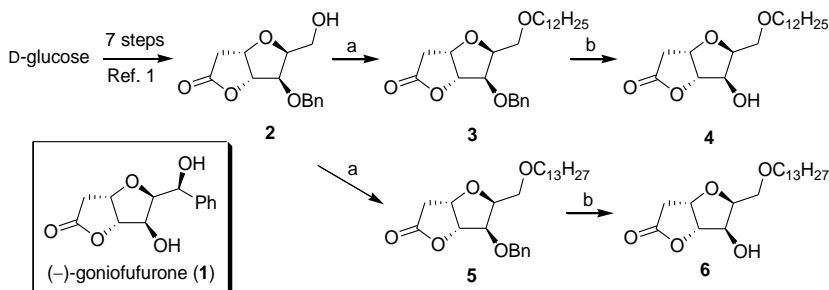
Sinteza i biološka ispitivanja novih analoga (–)-goniofufurona

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Sintetizovani su novi analozi (–)-goniofufurona (**1**), molekuli **4** i **6**, polazeći od komercijalno dostupne D-glukoze (Scheme 1).¹ Furanolaktoni **4** i **6** se mogu smatrati nestirilnim analogizima (–)-goniofufurona (**1**), suprotnog enantiomera prirodnog citotoksičnog laktona (+)-goniofufurona. Sintetizovana jedinjenja su podvrgnuta *in vitro* testovima prema velikom broju tumorskih ćelijskih linija, kao i prema normalnim fetalnim fibroblastima pluća. Osim antiproliferativne aktivnosti, biće predstavljeni i rezultati SAR analize.



Scheme 1. Reagents and conditions: (a) C₁₂H₂₅Br for **3**, C₁₃H₂₇Br for **5**, Ag₂O, AgOTf, Et₂O, reflux; (b) H₂, 10 % Pd/C, EtOH, rt.

Synthesis and biological evaluation of novel (–)-goniofufurone analogues

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New (–)-goniofufurone (**1**) analogues, compounds **4** and **6**, were synthesized starting from commercially available D-glucose (Scheme 1).¹ Furanolactones **4** and **6** might be considered as non-styryl analogues of (–)-goniofufurone (**1**), the opposite enantiomer of naturally occurring cytotoxic lactone (+)-goniofufurone. Synthesized compounds were evaluated by *in vitro* experiments against a number of tumour cell lines, as well as against normal foetal lung fibroblasts. Apart from antiproliferative activity the results of SAR analysis will be presented.

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1. V. Popsavin, B. Srećo, G. Benedeković, M. Popsavin, J. Francuz, V. Kojić, G. Bogdanović, *Bioorg. Med. Chem. Lett.* **2008**, *18*, 5182.