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## Development of new peptide antiagregatsionnyh heteromeric with imidazo[4,5-e]benzo[1,2-c;3,4-c']difuroxan moiety

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### Abstract

With the application of program Algotomb, mathematical simulation of heteromeric peptides comprising imidazo[4,5-e]benzo[1,2-c;3,4-c']difuroxan moiety. Effectiveness of their binding with GP IIb/IIIa-receptors of platelets is confirmed. The generated compounds were synthesized in the conditions of automatic peptide synthesizer Applied Biosystems 433A with the use of Fmoc-strategy. Evaluation of antiplatelet activity modeled heteromeric peptides showed the presence of dose-dependent inhibition of ADF-induced platelet aggregations.