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Author: Roman S. Esipov Dmitry A. Makarov Vasily N.

Stepanenko Anatoliy I. Miroshnikov

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ACCEPTED MANUSCRIPT

Development of the Intein-mediated method for production of recombinant Thymosin $\beta 4$ from the acetylated *in vivo* fusion protein

Roman S. Esipov, Dmitry A. Makarov, Vasily N. Stepanenko, Anatoliy I. Miroshnikov

M.M. Shemyakin and Yu.A. Ovchinnikov Institute of bioorganic chemistry of the Russian Academy of Sciences, GSP-7, Miklukho-Maklaya str. 16/10, 117997, Moscow, Russian Federation

Corresponding Author: Roman S. Esipov esipov@ibch.ru

Dmitry A. Makarov youngchemist@mail.ru

Vasily N. Stepanenko svn@ibch.ru

Anatoliy I. Miroshnikov aiv@ibch.ru

Highlights

The cleavage reaction took place only in the presence of the thiol reagent

The most efficient acetylation of the peptide was observed in the producer strain constructed on the basis of the *E. Coli* C3030 strain

The efficiency of the protein post-translational acetylation exceeded 90%

The serine acetyltransferase had a higher specificity with respect to thymosin $\beta 4$

The method developed is straightforward and economically viable

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