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(54) **Title:** METHODS AND COMPOSITIONS FOR GENETICALLY ENGINEERING CLOSTRIDIA SPECIES

(57) **Abstract:** The present invention relates to methods and compositions for engineering Clostridia species. In particular, embodiments of the present invention relate to the expression of recombinant resolvase proteins in Clostridia species.

## INTERNATIONAL SEARCH REPORT

International application No.  
**PCT/US2009/043314****A. CLASSIFICATION OF SUBJECT MATTER***C12N 15/31(2006.01)i, C12N 15/09(2006.01)i*

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**Minimum documentation searched (classification system followed by classification symbols)  
IPC as aboveDocumentation searched other than minimum documentation to the extent that such documents are included in the fields searched  
Korean Utility models and applications for Utility models  
Japanese Utility models and application for Utility modelsElectronic data base consulted during the international search (name of data base and, where practicable, search terms used)  
eKOMPASS (KIPO internal), Google, NCBI PubMed (clostridi\*, recU, resolvase, and similar terms.)**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X ----- A	E. P. C. ROCHA, et al. "Comparative and evolutionary analysis of the bacterial homologous recombination systems." In PLoS Genetics Vol.1(2):e15 (August 2005). See figure 2. Table 1.	1-17 ----- 18-22
A ----- X	D. Lyras, et al. "The large resolvase TnpX is the only transposon-encoded protein required for transposition of the Tn4451/3 family of integrative mobilizable elements." In Molecular Microbiology. Vol.51(6):1787-1800 (2004). See the abstract, figure 1, table 2, page 1791, Experimental procedure.	1-17 ----- 18-22
A ----- X	P. K. CRELLIN AND J. I. ROOD. "The resolvase/invertase domain of the site-specific recombinase TnpX is functional and recognizes a target sequence that resembles the junction of the circular form of the Clostridium perfringens transposon Tn4451." In Journal of Bacteriology. Vol.179(16):5148-5156 (August 1997). See the abstract, tables 1-2, figure 1, Materials and Methods.	1-17 ----- 18-22

 Further documents are listed in the continuation of Box C. See patent family annex.

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C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A ----- X	N. MCGREGOR, et al. "The structure of Bacillus subtilis RecU Holliday junction resolvase and its role in substrate selection and sequence-specific cleavage." In Structure. Vol.13:1341-1351 (September 2005). See the abstract, page 1349, left column, first paragraph.	1-17 ----- 18-22
A	T. Garnier, et al. "Molecular characterization of the resolvase gene, res, carried by a multicopy plasmid from Clostridium perfringens: common evolutionary origin for prokaryotic site-specific recombinases." In Molecular Microbiology. Vol.1(3):371-376 (1987). See the abstract, page 371.	1-22