

Review

Bacterial Protein Toxins as Anticancer Agents: Clinical Potential of *Pseudomonas* and Anthrax Toxins

Richa Misra ¹, Radhika Gupta ², Namita Nayyar ¹, Ritvik Baweja ³, Vishal Sharma ⁴, Yogendra Singh ^{5,*} and Renu Baweja ^{6,*}

¹ Department of Zoology, Sri Venkateswara College, University of Delhi, Delhi 110021, India; richamisra@svc.ac.in (R.M.); namitanayyar@svc.ac.in (N.N.)

² Department of Biochemistry, Daulat Ram College, University of Delhi, Delhi 110007, India; radhikagupta@dr.du.ac.in

³ Maulana Azad Medical College, Delhi 110002, India; ritvikbaweja@gmail.com

⁴ Department of Infectious disease, Albert Einstein College of Medicine, New York, NY 10461, USA; s.vishalzoology@gmail.com

⁵ Delhi School of Public Health, Institution of Eminence, University of Delhi, Delhi 110007, India

⁶ Department of Biochemistry, Shivaji College, University of Delhi, Delhi 110027, India

* Correspondence: ysinghdu@gmail.com (Y.S.); renubaweja@shivaji.du.ac.in (R.B.)

Abstract

Protein toxins are biologically active polypeptides produced by a variety of organisms, including bacteria, plants, fungi, and animals. These molecules exert potent and specific toxic effects on target cells and are primarily associated with pathogenicity and defense mechanisms of the organisms. In the past few decades, significant progress has been made in understanding their structure, mechanisms of action, and regulation. Among these, bacterial protein toxins have emerged as valuable tools particularly in the development of targeted therapies. A notable example is Botulinum toxin, originally known for its neurotoxic effects, which was approved as a therapeutic agent in 1989 for strabismus treatment, paving way for repurposing bacterial toxins for clinical use. This review provides an overview of the different classes of bacterial toxin-based therapeutics, with a particular focus on *Pseudomonas* exotoxin A (PE) from *Pseudomonas aeruginosa* and anthrax toxin from *Bacillus anthracis*. The modular architecture and potent cytotoxicity of these A-B type toxins have enabled their successful adaptation into targeted cancer therapies. The clinical approval of the PE-based immunotoxin, moxetumomab pasudotox, for the treatment of hairy cell leukemia, underscores the potential of this strategy. This review also discusses current challenges and outlines future directions for the advancement of bacterial toxin-based therapeutics.

Keywords: anthrax toxin; PE toxin; protein toxin; AB toxin; cancer therapy; immunotoxin

Key Contribution: This review highlights the therapeutic potential of bacterial protein toxins, focusing on how agents like *Pseudomonas* exotoxin A and anthrax toxin are being engineered to selectively target cancer cells, demonstrating significant clinical promise.



Received: 4 August 2025

Revised: 1 September 2025

Accepted: 9 September 2025

Published: 12 September 2025

Citation: Misra, R.; Gupta, R.; Nayyar, N.; Baweja, R.; Sharma, V.; Singh, Y.; Baweja, R. Bacterial Protein Toxins as Anticancer Agents: Clinical Potential of *Pseudomonas* and Anthrax Toxins. *Toxins* **2025**, *17*, 459. <https://doi.org/10.3390/toxins17090459>

Copyright: © 2025 by the authors.

Licensee MDPI, Basel, Switzerland.

This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (<https://creativecommons.org/licenses/by/4.0/>).

1. Introduction

Bacterial toxins belong to a large, biochemically diverse family of biotoxins produced by numerous living organisms, including plants (phytotoxins), fungi (mycotoxins), and animals (zootoxins). Bacterial toxins, traditionally recognized for their pathogenic roles in several infectious diseases, have increasingly been investigated as potential therapeutic