

CADTH Reference List

Monitoring for Drug-Induced Anaphylaxis With Injectable Antibiotics

August 2023

Summary of Abstracts



Key Messages

- We did not find any studies describing the risk of drug-induced anaphylaxis with a first dose of injectable antibiotics administered intravenously versus intramuscularly.
- We did not find any studies describing monitoring for drug-induced anaphylaxis with a first dose of antibiotics in a nonacute health care setting.
- We did not find any evidence-based recommendations concerning monitoring for drug-induced anaphylaxis for a first dose of antibiotics.
- We identified other references on this topic that may be of interest, which are listed in the report.

Research Questions

- 1. What is the clinical evidence describing risk of drug-induced anaphylaxis with a first dose of injectable antibiotics administered intravenously versus intramuscularly?
- 2. What is the clinical evidence describing monitoring for drug-induced anaphylaxis with a first dose of antibiotics in a nonacute health care setting?
- 3. What are the evidence-based recommendations concerning monitoring for drug-induced anaphylaxis for a first dose of antibiotics?

Methods

Literature Search Methods

An information specialist conducted a literature search on key resources including MEDLINE, Embase, the Cochrane Database of Systematic Reviews, the International HTA Database, the websites of Canadian and major international health technology agencies, as well as a focused internet search. The search approach was customized to retrieve a limited set of results, balancing comprehensiveness with relevancy. The search strategy comprised both controlled vocabulary, such as the National Library of Medicine's MeSH (Medical Subject Headings), and keywords. Search concepts were developed based on the elements of the research questions and selection criteria. The main search concepts were antibiotics, anaphylaxis, and injection. The search was completed on July 20, 2023, and limited to English-language documents published since January 1, 2018. Internet links were provided, where available.

Selection Criteria and Summary Methods

One reviewer screened literature search results (titles and abstracts) and selected publications according to the inclusion criteria presented in <u>Table 1</u>. Full texts of study publications were not reviewed. The Overall Summary of Findings was based on information available in the abstracts of selected publications. Open access full-text versions of evidence-based guidelines were reviewed when available, and relevant recommendations were summarized.



Criteria	Description
Population	Q1, Q2, and Q3: Individuals receiving a first dose of injectable antibiotics (i.e., IV, IM)
Intervention	Q1: Injectable antibiotics administered IV Q2 and Q3: Monitoring for drug-induced anaphylaxis in a nonacute care setting (e.g., home care, community- based care, long-term care)
Comparator	Q1: Injectable antibiotics administered IM Q2: Monitoring for drug-induced anaphylaxis in an acute care setting (e.g., hospital, tertiary care facility) Q1 and Q3: No comparator
Outcomes	Q1: Drug-induced anaphylaxis Q2: Clinical evidence of benefit (e.g., timely receipt of emergency medical services); or harm (e.g., morbidity, urticaria, shock); mortality Q3: Evidence-based recommendations
Study designs	Health technology assessments, systematic reviews, randomized controlled trials, nonrandomized studies, evidence-based guidelines

Table 1: Selection Criteria

IM = intramuscular.

Results

No relevant literature was identified regarding the risk of drug-induced anaphylaxis (DIA) with a first dose of injectable antibiotics administered intravenously versus intramuscularly. No relevant literature was identified regarding monitoring for DIA with a first dose of antibiotics in a nonacute health care setting. No evidence-based guidelines were identified regarding monitoring for DIA for a first dose of antibiotics. No health technology assessments, systematic reviews, randomized controlled trials, or nonrandomized studies were identified.

References of potential interest that did not meet the inclusion criteria are provided in Appendix 1.

Overall Summary of Findings

No relevant literature was found regarding risk of DIA with a first dose of injectable antibiotics administered intravenously versus intramuscularly. No relevant literature was found regarding monitoring for DIA with a first dose of antibiotics in a nonacute health care setting. Additionally, no evidence-based guidelines were found regarding monitoring for DIA for first dose of antibiotics; therefore, no summary can be provided.



References

Health Technology Assessments No literature identified.

Systematic Reviews No literature identified.

Randomized Controlled Trials No literature identified. Non-Randomized Studies No literature identified.

Guidelines and Recommendations No literature identified.



Appendix 1: References of Potential Interest

Previous CADTH Reports

Vu T, Horton J. First dose of antibiotics administered using intravenous push versus intravenous mini bag systems. Ottawa (ON): CADTH; 2021: <u>https://www.cadth.ca/sites/default/files/pdf/htis/2021/RB1576%20IV%20antibiotics%20Final.pdf</u>. Accessed 2023 Jul 24.

Nonrandomized Studies

No Comparator

Kovacik CN, Shah MD, Thomas TA, Eby JC. First-dose antimicrobial infusion reactions in patients enrolled in outpatient parenteral antimicrobial therapy services. *Open Forum Infect Dis*. 2023 Jun;10(6):ofad239. <u>PubMed</u>

Guidelines and Recommendations

Drug Challenges Not Specific to Monitoring in Non-Acute Care Settings

Kahn DA, Banerji A, Blumenthal KG, et al. Drug allergy: a 2022 practice parameter update. J Allergy Immunol. 2022;150(6):1333-1393. Available from: <u>https://www.jacionline.org/article/S0091-6749(22)01186-1/pdf</u>. Accessed 2023 Jul 24. <u>PubMed</u> Refer to: Consensus-Based Statement 12 (p. 1357), Statement 13 to 14 (p. 1359), Statement 17 (p. 1360)

Review Articles

Macy E, Adkinson NF Jr. The evolution of our understanding of penicillin allergy: 1942-2022. J Allergy Clin Immunol Pract. 2023 Feb;11(2):405-413. PubMed



Authors: Weiyi Xie, Carolyn Spry

Contributor: Camille Santos

Cite As: Monitoring for Drug-Induced Anaphylaxis with Injectable Antibiotics. (CADTH reference list: summary of abstracts). Ottawa: CADTH; 2023 Aug.

Disclaimer: The information in this document is intended to help Canadian health care decision-makers, health care professionals, health systems leaders, and policymakers make well-informed decisions and thereby improve the quality of health care services. While patients and others may access this document, the document is made available for informational purposes only and no representations or warranties are made with respect to its fitness for any particular purpose. The information in this document should not be used as a substitute for professional medical advice or as a substitute for the application of clinical judgment in respect of the care of a particular patient or other professional judgment in any decision-making process. The Canadian Agency for Drugs and Technologies in Health (CADTH) does not endorse any information, drugs, therapies, treatments, products, processes, or services.

While care has been taken to ensure that the information prepared by CADTH in this document is accurate, complete, and up to date as at the applicable date the material was first published by CADTH, CADTH does not make any guarantees to that effect. CADTH does not guarantee and is not responsible for the quality, currency, propriety, accuracy, or reasonableness of any statements, information, or conclusions contained in any third-party materials used in preparing this document. The views and opinions of third parties published in this document do not necessarily state or reflect those of CADTH.

CADTH is not responsible for any errors, omissions, injury, loss, or damage arising from or relating to the use (or misuse) of any information, statements, or conclusions contained in or implied by the contents of this document or any of the source materials.

This document may contain links to third-party websites. CADTH does not have control over the content of such sites. Use of third-party sites is governed by the third-party website owners' own terms and conditions set out for such sites. CADTH does not make any guarantee with respect to any information contained on such third-party sites and CADTH is not responsible for any injury, loss, or damage suffered as a result of using such third-party sites. CADTH has no responsibility for the collection, use, and disclosure of personal information by third-party sites.

Subject to the aforementioned limitations, the views expressed herein do not necessarily reflect the views of Health Canada, Canada's provincial or territorial governments, other CADTH funders, or any third-party supplier of information.

This document is prepared and intended for use in the context of the Canadian health care system. The use of this document outside of Canada is done so at the user's own risk.

This disclaimer and any questions or matters of any nature arising from or relating to the content or use (or misuse) of this document will be governed by and interpreted in accordance with the laws of the Province of Ontario and the laws of Canada applicable therein, and all proceedings shall be subject to the exclusive jurisdiction of the courts of the Province of Ontario, Canada.

The copyright and other intellectual property rights in this document are owned by CADTH and its licensors. These rights are protected by the Canadian *Copyright Act* and other national and international laws and agreements. Users are permitted to make copies of this document for non-commercial purposes only, provided it is not modified when reproduced and appropriate credit is given to CADTH and its licensors.

About CADTH: CADTH is an independent, not-for-profit organization responsible for providing Canada's health care decision-makers with objective evidence to help make informed decisions about the optimal use of drugs, medical devices, diagnostics, and procedures in our health care system.

Funding: CADTH receives funding from Canada's federal, provincial, and territorial governments, with the exception of Quebec.

Questions or requests for information about this report can be directed to requests@cadth.ca