TITLE: Peripheral Intravenous Infusion of 5-fluorouracil versus Delivery by Central Venous Access Device: Comparative Clinical Effectiveness and Guidelines

DATE: 19 January 2009

RESEARCH QUESTIONS:

1. What is the comparative clinical effectiveness of peripheral intravenous infusion of 5-fluorouracil versus delivery by a central venous access device?

2. What are the guidelines associated with mode of delivery of 5-fluorouracil?

METHODS:

A limited literature search was conducted on key health technology assessment resources, including PubMed, the Cochrane Library (Issue 4, 2008), University of York Centre for Reviews and Dissemination (CRD) databases, ECRI, EuroScan, international HTA agencies, and a focused Internet search. Results include articles published between 2004 and January 2009 and are limited to English language publications only. For question #1, no filters were applied to limit the retrieval by study type. For question #2, a filter was applied to limit the retrieval to clinical guidelines. Internet links are provided, where available.

RESULTS:

HTIS reports are organized so that the higher quality evidence is presented first. Therefore, health technology assessment reports, systematic reviews, and meta-analyses are presented first. These are followed by randomized controlled trials (RCTs), controlled clinical trials, observational studies, and evidence-based guidelines.

One RCT and two observational studies were identified pertaining to the comparative clinical effectiveness of peripheral intravenous infusion of 5-fluorouracil versus delivery by a central venous access device. No HTAs, systematic reviews, clinical controlled studies or evidence based guidelines were identified. Additional information that may be of interest has been included in the appendix.

Disclaimer: The Health Technology Inquiry Service (HTIS) is an information service for those involved in planning and providing health care in Canada. HTIS responses are based on a limited literature search and are not comprehensive, systematic reviews. The intent is to provide a list of sources of the best evidence on the topic that CADTH could identify using all reasonable efforts within the time allowed. HTIS responses should be considered along with other types of information and health care considerations. The information included in this response is not intended to replace professional medical advice, nor should it be construed as a recommendation for or against the use of a particular health technology. Readers are also cautioned that a lack of good quality evidence does not necessarily mean a lack of effectiveness particularly in the case of new and emerging health technologies, for which little information can be found, but which may in future prove to be effective. While CADTH has taken care in the preparation of the report to ensure that its contents are accurate, complete and up to date, CADTH does not make any guarantee to that effect. CADTH is not liable for any loss or damages resulting from use of the information in the report.

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Health technology assessments
No literature identified.

Systematic reviews and meta-analyses
No literature identified.

Randomized controlled trials


Controlled clinical trials
No literature identified.

Observational studies


Guidelines and recommendations
No literature identified.

PREPARED BY:
Kristen Moulton, BA, Research Assistant
Charlene Argáez, MLIS, Information Specialist
Health Technology Inquiry Service
Email: htis@cadth.ca
Tel: 1-866-898-8439
APPENDIX – FURTHER INFORMATION:

Observational studies


Additional references