



**RETURN BIDS TO:**

**RETOURNER LES SOUMISSIONS À:**

Bid Receiving - PWGSC / Réception des soumissions -  
TPSGC

11 Laurier St. / 11, rue Laurier

Place du Portage, Phase III

Core 0B2 / Noyau 0B2

Gatineau

Quebec

K1A 0S5

Bid Fax: (819) 997-9776

**SOLICITATION AMENDMENT  
MODIFICATION DE L'INVITATION**

The referenced document is hereby revised; unless otherwise indicated, all other terms and conditions of the Solicitation remain the same.

Ce document est par la présente révisé; sauf indication contraire, les modalités de l'invitation demeurent les mêmes.

**Comments - Commentaires**

**Vendor/Firm Name and Address**

Raison sociale et adresse du  
fournisseur/de l'entrepreneur

**Issuing Office - Bureau de distribution**

Information Products/Produits d'information

L'Esplanade Laurier,

East Tower 7th Floor

140 O'Connor, Street

Ottawa

Ontario

K1A 0R5

<b>Title - Sujet</b> Electronic Medical PointofCare Tool	
<b>Solicitation No. - N° de l'invitation</b> W6369-21A003/A	<b>Amendment No. - N° modif.</b> 003
<b>Client Reference No. - N° de référence du client</b> W6369-21A003	<b>Date</b> 2020-07-19
<b>GETS Reference No. - N° de référence de SEAG</b> PW-\$\$PI-035-78842	
<b>File No. - N° de dossier</b> pi035.W6369-21A003	<b>CCC No./N° CCC - FMS No./N° VME</b>
<b>Solicitation Closes - L'invitation prend fin</b> <b>at - à 02:00 PM</b> <b>on - le 2020-08-05</b>	<b>Time Zone</b> <b>Fuseau horaire</b> Eastern Daylight Saving Time EDT
<b>F.O.B. - F.A.B.</b> <b>Plant-Usine:</b> <input type="checkbox"/> <b>Destination:</b> <input type="checkbox"/> <b>Other-Autre:</b> <input type="checkbox"/>	
<b>Address Enquiries to: - Adresser toutes questions à:</b> Henry(pi035), Katelyn	<b>Buyer Id - Id de l'acheteur</b> pi035
<b>Telephone No. - N° de téléphone</b> (343) 550-0484 ( )	<b>FAX No. - N° de FAX</b> ( ) -
<b>Destination - of Goods, Services, and Construction:</b> <b>Destination - des biens, services et construction:</b> See Herein	

Instructions: See Herein

Instructions: Voir aux présentes

<b>Delivery Required - Livraison exigée</b>	<b>Delivery Offered - Livraison proposée</b>
<b>Vendor/Firm Name and Address</b> <b>Raison sociale et adresse du fournisseur/de l'entrepreneur</b>	
<b>Telephone No. - N° de téléphone</b> <b>Facsimile No. - N° de télécopieur</b>	
<b>Name and title of person authorized to sign on behalf of Vendor/Firm</b> <b>(type or print)</b> <b>Nom et titre de la personne autorisée à signer au nom du fournisseur/</b> <b>de l'entrepreneur (taper ou écrire en caractères d'imprimerie)</b>	
<b>Signature</b>	<b>Date</b>

**Amendment 003** is raised to respond to questions received during the solicitation period.

**Questions and Answers**

#	Question	Response
7	Is all information required by the Tool already available as part of any current system at CFHS? If so, are there standard means to read the information from such systems?	In accordance with RFP Section 4.2.2 (M3) "Technical requirements, Mandatory Criteria" and Annex D, Section 1, <b>the tool</b> "must contain the medically relevant, current and thorough information on condition(s) searched. Must provide a summary, general information, diagnosis, treatment, and care." Section 4.2.7 (M8) provides the topics/areas that must be covered by the tool. CFHS is looking to acquire a tool containing the information outlined in the Statement of Work.
8	The mandatory requirements don't refer to the ability to gather newer information - can it be therefore assumed that the Tool will always read from one source for all the information needs?	In accordance with RFP Section 4.2.4 (M5) "Technical requirements, Mandatory Criteria" and Annex D, Section 1, <b>the tool</b> "must provide current information to clinical questions" and <b>the bidder</b> "must provide a document which includes an explanatory paragraph or paragraphs for each of the following aspects of the update methodology of the contents of the Tool: (1) rigour, (2) scope, (3) process, and (4) frequency."
9	Is there a standard to measure the quality of evidence for information in the Tool i.e. since it's an evidence-based Tool for information, whether CFHS has a standard to be followed to measure evidence or rate it as acceptable evidence?	In accordance with RFP Section 4.2.3 (M4) "Technical requirements, Mandatory Criteria" and Annex D, Section 1, the "information published in the tool must be subject to a transparent and comprehensive review", and <b>the bidder</b> "must provide a document which includes an explanatory paragraph or paragraphs for each of the following criteria: (1) rigour, (2) scope, (3) process, and (4) frequency of the peer-review and publication processes. The peer review must be at least a 2-level review process."
10	What are the details of the current network infrastructure available for CFHS clinicians?	In accordance with RFP section 4.1.1 (M1) "Accessibility, Mandatory Criteria" and Annex D, Section 1, CFHS clinicians must be able to access this tool on any web-enabled Defence Wide Area Network (DWAN) computers within designated institutions. (Locations in Canada, the USA, Overseas, and a fluctuating number of deployment locations – current estimate approx. 60 total.)