

## PUBLIC HEALTH UNIT INFECTION PREVENTION AND CONTROL LAPSE REPORT

### Initial Report

Premise/facility under investigation (name and address)	Orangeville Medical Imaging 314 Broadway, Orangeville ON L9W 1L3
Type of premise/facility: (E.g. clinic, personal services setting)	Diagnostic Imaging
Date Board of Health became aware of IPAC lapse	May 20, 2020
Date of Initial Report posting	June 8, 2020
Date of Initial Report update(s) (if applicable)	
How the IPAC lapse was identified	A complaint by a member of the public.
Summary Description of the IPAC Lapse	During the complaint inspection it was documented that the cleaning and disinfection of medical equipment and devices on-site did not follow the Provincial Infectious Diseases Advisory Committee (PIDAC) Best Practice Standards for medical device reprocessing and infection control in the clinical office setting.
<b>IPAC Lapse Investigation</b>	
Did the IPAC lapse involve a member of a regulatory college?	Yes
If yes, was the issue referred to the regulatory college?	Yes
Were any corrective measures recommended and/or implemented?	Yes
Please provide further details/steps	The medical practice is to ensure: <ul style="list-style-type: none"> <li>Invasive vaginal ultrasound/endocavity transducer procedures are halted until Wellington-Dufferin-Guelph Health Unit has given written permission to resume</li> </ul>

- Reusable vaginal ultrasound probes/endocavity transducers are cleaned, disinfected and rinsed as per manufacturer's instructions for use (MIFU), PIDAC Best Practice and CAN/CSA 2314-18 (as current)
- A log to document high level disinfectant solution and vaginal ultrasound probes/endocavity transducers disinfection is provided and completed
- An enzymatic detergent for cleaning the reusable vaginal ultrasound probe/endocavity transducer that is compatible with the reusable vaginal ultrasound probe/endocavity transducer as per MIFU is provided
- A dedicated hand washing sink located in or near the reprocessing area is provided
- All staff members involved in cleaning and disinfecting reusable ultrasound /endocavity probes used for invasive ultrasound procedures must complete the online Public Health Ontario (PHO) Reprocessing Modules on cleaning and high level disinfection (modules 1-4) before cleaning and disinfecting reusable vaginal ultrasound probes/endocavity transducers and resuming invasive vaginal ultrasound/endocavity procedures, and provide the certificates of completion for each staff member to Wellington-Dufferin-Guelph Health Unit
- A plan is provided to Wellington-Dufferin-Guelph Health Unit for the development of site-specific policies and procedures based on current PIDAC Best Practices for clinical office practice and for onsite cleaning and disinfection of reusable vaginal ultrasound probes
- Ventilation requirements for Cidex® OPA used onsite meet Ministry of Labour and manufacturer's instructions requirements.
- An eye washing facility in the vaginal ultrasound probes/endocavity transducer reprocessing area

Date any order(s) or directive(s) were issued to the owners/operators (if applicable)

Written Order to Mr. Kanwal Bhullar: May 20, 2020

**Initial Report Comments and Contact Information**

Any Additional Comments (Do not include any personal information or personal health information)

<b>If you have any further questions, please contact:</b>	
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<b>Final Report</b>	
Date of Final Report posting:	July 20, 2020
Date any order(s) or directive(s) were issued to the owner/operator (if applicable)	
Brief description of corrective measures taken	Staff members at Orangeville Medical Imaging have worked cooperatively with Wellington-Dufferin-Guelph Public Health to make required changes to the onsite infection prevention and control practices. On July 17, 2020 a final re-inspection was conducted. At this time, infection prevention and control (IPAC) practices were in compliance with IPAC Best Practices. Patient care services using invasive vaginal ultrasound probe/endocavity transducer was permitted to resume as all items specified in the section 13 Order under the Health Protection and Promotion Act have been addressed.
Date all corrective measures were confirmed to have been completed	July 17, 2020
<b>Final Report Comments and Contact Information</b>	
Any Additional Comments (Do not include any personal information or personal health information)	
<b>If you have any further questions, please contact:</b>	
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