GENERAL ULTRASOUND

DIAGNOSTIC MEDICAL SONOGRAPHER OBSERVATION FORM

Please complete one form for each examination observed

DMS OBSERVED:		
CMRTO #:		
PATIENT IDENTIFIER:		
PATIENT WRITTEN CONSENT OBTAINED:		
TYPE OF EXAMINATION OBS	ERVED?	

	С	NC	NA
1.8.1 DUTIES AND RESPONSIBILITIES OF DMSs			
Follow facility policy regarding situations where the use of chaperones may be			
appropriate.			
Post appropriate signage to restrict access to the patient exam room. (If not, is there a policy to restrict access?)			
Ensure the room is prepared for the procedure specified in the order.			
Select and set up the equipment and materials needed for the procedure			
specified in the order.			
Ensure correct patient identification (e.g. confirmation of patient name, date			
of birth, examination to be performed, and physician/authorized health			
professional authorization is present).			
Confirm that the order is appropriate based on the patient history.			
Inquire about and record any contraindications (e.g. latex allergy) before			
starting the exam.			
Ensure that the worklist contains the correct patient information (if			
applicable).			

	С	NC	NA
Obtain informed consent (oral or written as per facility policy) before each			
examination (after explaining the procedure and answering any questions).			
Ensure pertinent clinical history is available, supplement as necessary and			
record on the technical impression worksheet.			
Instruct the patient to remove only the clothing and items that will interfere			
with the procedure, providing the patient with a gown or sheet to cover areas			
where clothing was removed and explaining to the patient when and where			
the DMS may touch them and why.			
Follow the facility examination protocols.			
Write a technical impression as per site protocol.			
Follow facility protocols when unexpected findings are found that would			
require immediate attention (e.g. appendicitis, ectopic pregnancy).			
Allergies to latex must be identified and non-latex transducer covers must be			
utilized. This information must be recorded on the sonographer's technical			
impression worksheet.			
THROUGHOUT THE EXAMINATION:			
Assess the patient's condition before, during and after the procedure or			
course of treatment and make modifications to procedures based on the			
patient's physical, medical and/or emotional status and needs.			
Maintain patient comfort, privacy and dignity at all times.			
Stop procedure if at any time the patient withdraws consent and record			
withdrawal of consent and reason as per site protocol.			
Use PPE (personal protective equipment masks/gloves/gown etc.) and devices			
as required for the procedure and as indicated by personal risk assessment.			
Ensure that patient examination images and data contains patient name, ID			
number, date of examination and type of examination and number of images.			
Ensure images were scanned as per site protocol and include:			
o correct annotation			
o fine & total gain controls set correctly			
o appropriate magnification			
o focal zone set correctly			
o proper use of calipers			
o measurements documented			
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o scan correctly annotated			
o scan through the entire organ appropriately			
o the technical worksheet is suitable for regions examined			
Ensure the processed image provides diagnostic image quality while			
minimizing patient exposure to soundwaves (ALARA – As Low As Reasonably			
Achievable). Take corrective action if necessary and record explanation of sub-optimal imaging.			
Ensure that each patient record (including the technical impression			
worksheet) has the DMS identifier to verify who performed the examination.			
Comply with privacy and confidentiality legislation such as the <i>Personal Health</i>			
Information Protection Act (Ontario). Was patient privacy maintained at all			
times?			
TRANSVAGINAL/ENDOCAVITY ULTRASOUNDS: include the criteria above plus:			
Transvaginal/endocavity transducer ID number (individual to each transducer)			
must be identified on the reprocessing sheet.			
Upon exam completion follow Provincial Infectious Diseases Advisory			
Committee (PIDAC) or manufacturers guidelines for transducer cleaning.			
Ensure Internal & External Gel use meets PIDAC guidelines.			
IMAGE REVIEW:			
Are there enough images to allow 3 rd party interpretation?			
Ensure the examination includes interrogation of all relevant anatomy using			
appropriate transducers and gain settings.			
General Comments: (Please use this section to provide overall comments regards	ing the	e	
technologist's performance, attitude, competency infection control procedures in	ncludir	ng gel	and
probe cleaning. etc.). Document products used.			

NC

NA

Recommendations: These recommendations must be documented in the Final Assessment
Report