ACR Ultrasound Practice
Accreditation and Technical Standard for Ultrasound Performance Monitoring

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Overview of each document
Ongoing evolution of the accreditation programs
Practical aspects of QC testing
Future considerations
Performance evaluations are distinguished from preventive maintenance.

Recommended that performance evaluations be done by a physicist, but flexibility is allowed.

Subjective or objective testing methods may be used.

Phantoms must be used (commercial or custom).

Probe testing systems may be used.

*All* probes must be tested at QC at least semiannually.

Electronic scanner (and primary diagnostic) image displays must be tested.
<table>
<thead>
<tr>
<th>Test</th>
<th>Acceptance Testing** (M)</th>
<th>Quality Control</th>
<th>Annual Testing**</th>
</tr>
</thead>
<tbody>
<tr>
<td>Physical and mechanical inspection</td>
<td>S</td>
<td>M</td>
<td>M</td>
</tr>
<tr>
<td>Image uniformity and artifact survey</td>
<td>S</td>
<td>M</td>
<td>M</td>
</tr>
<tr>
<td>Geometric accuracy</td>
<td>S</td>
<td>M*</td>
<td>M</td>
</tr>
<tr>
<td>System sensitivity</td>
<td>S</td>
<td>S</td>
<td>M</td>
</tr>
<tr>
<td>Spatial resolution</td>
<td>S</td>
<td>S</td>
<td>S</td>
</tr>
<tr>
<td>Contrast resolution</td>
<td>S</td>
<td>S</td>
<td>M</td>
</tr>
<tr>
<td>Fidelity of US scanner electronic image display</td>
<td>S</td>
<td>(S)</td>
<td>M</td>
</tr>
<tr>
<td>Fidelity of display devices used for primary interpretation</td>
<td>S</td>
<td>S</td>
<td>S</td>
</tr>
<tr>
<td>Qualitative evaluations of Doppler functionality</td>
<td>S</td>
<td>S</td>
<td>S</td>
</tr>
</tbody>
</table>

*M* = "must do" / required  
*S* = "should do" / optional

* Only needed for mechanically-scanned probes

** All tests done for QC must be included
General approach: Make the best use of time invested by the practice in routine QC by requiring efficient tests with demonstrated utility.


Four-year experience with a clinical ultrasound quality control program.

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Abstract

Ultrasound (US) quality control (QC) program data over a 4-year period from more than 45 scanners and more than 265 transducers were reviewed to optimize the program in terms of efficiency and effectiveness. Our program included evaluations of mechanical integrity, image uniformity, distance measurement accuracy and maximum depth of penetration (DOP). We computed failure rates and fraction of failures detected by each test. A total of 187 equipment problems were identified. Average annual scanner component and transducer failure rates were 10.5% and 13.9%, respectively. The mechanical integrity and uniformity evaluations detected 25.1% and 66.3% of all failures, respectively. Those evaluations plus defects detected by sonographers accounted for 98.4% of all detected failures. DOP and distance measurement accuracy were not effective at detecting equipment failures. For routine US QC, we recommend quarterly mechanical integrity and uniformity assessments of all transducers. A scanner with five transducers could be tested in an estimated 30 min or less.

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The accreditation program QC requirements are meant to align with those in the technical standard, which should serve as minimum requirements.

It might be expected that QC requirements in accreditation programs involving the same modality would be very similar, if not identical.
Ultrasound Accreditation Program Requirements

- Short list of "must do" tests (sensitivity, uniformity, electrical & mechanical safety, photo and hard-copy)
- Only 2 most commonly used transducers must be tested at QC
- Use of phantom is stated as optional
- Scanner display testing not mentioned (hard copy is included)
- Fairly detailed methods for some tests are embedded in the program document

(Revised 3/23/12)
Continuous Quality Control

Routine quality control testing must occur regularly; *a minimum requirement is semiannually*. The same tests must be performed during each testing period so that changes can be monitored over time and effective corrective action can be taken. Testing results, corrective action, and the effects of corrective action must be documented and the documentation maintained on site. In the event of a site survey, reviewers will expect to see such documentation.

The QC program must evaluate at least the following items in gray-scale imaging mode:

- System sensitivity and/or penetration capability.
- Image uniformity.
- Assurance of electrical and mechanical safety and cleanliness
- Photography and other hard-copy recording.

In addition, it is recommended that users verify the accuracy of vertical and horizontal distance measurement when a QC program is initiated for an ultrasound unit.

These items may be assessed using a commercially available phantom test object. At the present time, no one type of phantom is preferred; users should select one that is commercially available. Using a phantom will be helpful in responding to questions about low-contrast detectability in the quality control part of the testing material. However, the use of a phantom is *optional* at this time. Questions relating to characteristics associated with system sensitivity and image uniformity may be answered without the use of a phantom as a test object.
Breast Ultrasound Accreditation Program Requirements

- Long list of “recommended” tests, but no “must do” tests
- Scanner display testing not mentioned (but hard copy testing is included)
- Large loophole is offered…

Revised 6/21/12
Quality Control

The following routine QC should be performed on all ultrasound units used for breast imaging as recommended in the ACR Technical Standard for Diagnostic Medical Physics Performance Monitoring of Real Time Ultrasound Equipment:

<table>
<thead>
<tr>
<th>Recommended Quality Control for Breast Ultrasound</th>
<th>Frequency</th>
<th>Performed By</th>
</tr>
</thead>
<tbody>
<tr>
<td>Maximum depth of visualization and hardcopy recording with a tissue-mimicking phantom</td>
<td>Semiannually</td>
<td>Service engineer/medical physicist</td>
</tr>
<tr>
<td>Vertical and horizontal distance accuracy</td>
<td>Semiannually</td>
<td>Service engineer/medical physicist</td>
</tr>
<tr>
<td>Uniformity</td>
<td>Semiannually</td>
<td>Service engineer/medical physicist</td>
</tr>
<tr>
<td>Electrical-mechanical cleanliness condition</td>
<td>Semiannually</td>
<td>Service engineer/medical physicist</td>
</tr>
<tr>
<td>Anechoic void perception</td>
<td>Semiannually</td>
<td>Service engineer/medical physicist</td>
</tr>
<tr>
<td>Ring down</td>
<td>Semiannually</td>
<td>Service engineer/medical physicist</td>
</tr>
<tr>
<td>Lateral resolution</td>
<td>Semiannually</td>
<td>Service engineer/medical physicist</td>
</tr>
<tr>
<td>Quality control checklist</td>
<td>Semiannually</td>
<td>Service engineer/medical physicist</td>
</tr>
<tr>
<td>Adherence to universal infection control procedures</td>
<td>After each biopsy</td>
<td>Technologist</td>
</tr>
<tr>
<td>Clean transducers</td>
<td>After each patient</td>
<td>Technologist</td>
</tr>
<tr>
<td>Vertical and horizontal distance accuracy</td>
<td>Quarterly</td>
<td>Technologist</td>
</tr>
<tr>
<td>Grey-scale photography</td>
<td>Quarterly</td>
<td>Technologist</td>
</tr>
</tbody>
</table>

As part of accreditation, facilities must submit a copy of the service engineer’s most recent preventive maintenance report or the medical physicist’s most recent equipment survey. Although the ACR will not initially use this information to determine whether a facility passes or fails accreditation, it may be used in the future to set criteria.
Ongoing evolution of the accreditation programs

- Currently developing a new ultrasound performance testing section
  - Used in both US-related Accreditation Programs
  - Correlate closely with the Technical Standard
  - Consider acceptance testing, quality control, and an annual survey
  - No additional specific testing for re/application
  - Include an appendix describing sample methods for performing QC tests
Ultrasound Quality Control Manual
- Standard phantom?
- Standard testing methodology?
- Specific performance targets?
Practical aspects of QC testing

- Physical and mechanical inspection
- Image uniformity and artifact survey
  
  sonog reported problems > 98% of failures

- Ultrasound scanner electronic image display
Physical and mechanical inspection
Image uniformity and artifact survey
BK FlexFocus bi-plane prostate probe

Sagittal Array

Transverse Array
Ultrasound scanner electronic image display

- Ultrasound scanner monitor is a primary diagnostic display device
  - Overall display quality
  - Luminance calibration
Future considerations

- Continued assessment of the utility of existing QC tests and tests proposed in the future
  - Spectral and color Doppler?
- Increased availability of software tools: migration from subjective to objective methods
  - E.g. median or mean processing of US clips (AAPM)
- Improved correlation of equipment flaws and impacts on clinical utility
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