Example of an Ultrasonic Cleaner (SonoCheck™ only) Policy for Daily Cleaning and Monitoring

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NOTE: This document is an example of a policy that may be instituted in a health-care facility for Daily Monitoring of the cavitation activity of any Ultrasonic Cleaner. The actual policy in a facility must be based on variables, logistics and risk-assessments that are specific to your facility.

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SUBJECT: ULTRASONIC CLEANER MONITORING DAILY (SonoCheck™ only)

DEPARTMENT: CPD, CSSD, SPA, MDRD, dental offices, medical offices and departments using a sonic cleaner.

APPROVED BY:

EFFECTIVE:

REVISED: Jan. 2019

PURPOSE: The purpose of this example policy is to monitor the ultrasonic cavitation process daily to ensure proper cavitation is taking place and reduce risk of infection to personnel or patients. (1,10,13)

POLICY: To inspect and test any ultrasonic cleaner with the SonoCheck™ For cavitation only. The SonoCheck™ is to be used according to the manufacture's guidelines to ensure that cavitation is occurring. (1,6,7,8,9,10,13)

RATIONALE: The SonoCheck™ it to test for the presence of cavitation energy in a sonic bath under normal conditions in an empty tank than has been degassed.
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**Standards and Professional Society Recommendations:**

ST 79;2017 7.6.4.3.1 section states “…. Mechanical cleaning methods minimize personnel risk of cross-contamination, improve cleaning effectiveness, increase productivity, and are more easily monitored for quality performance”. Sonic cleaners are considered mechanical cleaning equipment by AAMI.

“Cleaning, not sterilization (or disinfection) is the first and most important step in any instrument processing protocol. Without first subjecting the instrument to a thorough, validated and standardized (and ideally automated) cleaning process, the likelihood that any disinfection or sterilization process will be effective is significantly reduced”. (11)

The ultrasonic cleaning process cleans dirty surgical instruments so they can be handled safely, repackaged, and sterilized for a future surgery. The danger of handling instruments contaminated with blood is obvious in this age of hepatitis, CJD and HIV. The procedures for sterilizing instruments are based on years of scientific testing of cleaning instruments. If surgical instruments are not clean, the procedures are ineffective. Dried blood on instruments is hazardous to the employees of the hospital and to the next surgical patient upon which the instruments are used. (1,2,3,4,8,)

An ultrasonic cleaner enables thorough cleaning of equipment by producing cavitation and combining with the other factors such as cleaning solution, water quality and time and temperature to produce a clean medical device. Ensuring cavitation is taking place is vital for any ultrasonic cleaner, cavitation is why you purchase an ultrasonic cleaner thus you want to make sure cavitation is being produced.

Ultrasonic cleaners do not disinfect instruments.

Ultrasonic vibration at the frequency used for cleaning does not kill microorganisms and infective aerosols may be produced. It is for this reason that the lid of the tank must be tightly closed during operation.

Ultrasonic cleaners work by subjecting instruments to high frequency, high-energy sound waves. This causes the soil to be dislodged from instruments and drop to the bottom of the tank or be sufficiently loosened that it will be removed during the rinsing process. This is called cavitation.
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The detergent used in the ultrasonic tank must be carefully selected in accordance with advice from the tank's manufacturer. Optimally it will be a neutral, low-foaming product and enzymatic cleaners will have enhanced benefits in this process.

Degassing of cleaning solutions is extremely important in achieving satisfactory cleaning results. Air impedes the cavitation process and thus you need to free as much trapped air in the solution so you can maximize the effect of the cavitation. Degassing is one of the best methods for releasing trapped air in cleaning solutions. Fresh solutions or solutions which have cooled must be degassed before proceeding with cleaning. Degassing is done after the chemical is added and is accomplished by operating the ultrasonic energy and raising the solution temperature. The time required for degassing varies considerably, based on tank capacity and solution temperature, and may range from several minutes for a small tank to an hour or more for a large tank. Degassing is complete when small bubbles of gas cannot be seen rising to the surface of the liquid and a pattern of ripples can be seen.

A Quality Management System (QMS) per ANSI / AAMI ST 90 should be developed for the ultrasonic cleaner. THE QMS should include the IQ, OQ, and PQ of the ultrasonic cleaner.

**INSTALLATION QUALIFICATION (IQ) = PROCESS OF OBTAINING AND DOCUMENTING EVIDENCE THAT EQUIPMENT HAS BEEN PROVIDED AND INSTALLED IN ACCORDANCE WITH IT SPECIFICATION.**

**OPERATIONAL QUALIFICATION (OQ) = PROCESS OF OBTAINING AND DOCUMENTING EVIDENCE THAT INSTALLED EQUIPMENT OPERATES WITHIN PREDETERMINED LIMITS WHEN USED IN ACCORDANCE WITH ITS OPERATIONAL PROCEDURES.**

Performance qualification (PQ) = Process of obtaining and documenting evidence that the equipment, as installed and operated in accordance with operational procedures, consistently performs in accordance with predetermined criteria and thereby yields product meeting its specifications

**Routine cleaning:**
Cleaning the ultrasonic cleaner and replacement of the cleaning solution is
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necessary at least daily or more frequently if solution soiled. Follow manufactures guidelines on changing solution.

**Performance testing:**
The efficacy of the ultrasonic cleaner should be tested each day it is used. The results of the testing shall be documented as part of the proof of process or the PQ.

Sonic cleaners fail for many reasons. Tests should provide a means of monitoring the variables that influence the effectiveness of the ultrasonic cleaning process. Some of these variables are water, time, detergent, enzyme, temperature, high pH, agitation, speed, tray selection, initial heat, drying, obstructions, and insufficient amount of chemicals and equipment failure. (7)

Proper cleaning is critical. The SonoCheck™ provides an independent objective test of cavitation and allows the Sterile Processing professional to monitor and ensure proper cavitation in the sonic process. (I)

AAMI does list a sonic cleaner as a piece of medical equipment in section 13.2
ANNEX D states “…users should incorporate test methods that verify the functionality of the mechanical cleaning equipment (if used) and the cleanliness of specific devices after manual or mechanical cleaning is completed. These verification tests are part of continuous quality improvement to demonstrate continued compliance with cleaning benchmarks, once these benchmarks have been defined.”

Section 13.2 states “…Mechanical cleaning equipment should be tested upon installation, each day that it is use, and after major repairs. When evaluating or changing to a new cleaning solution and after all major repairs, all cycles used should be tested to ensure that the cleaning solution and cleaning action are effective. A major repair is a repair that is outside the scope of routine preventive maintenance and that significantly affects the performance of the equipment. Examples include software upgrade or the replacement of the water pump(s), detergent delivery system, heating system, water delivery system, water treatment system, ultrasonic generators, or computer controls”.

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JCAHO in standard E.C.6.20 it states that medical equipment is maintained, tested and inspected

In accordance with the various standards and guidelines, Healthmark recommends daily testing each day the unit is used to monitoring both for cavitation and the cleaning efficacy of any sonic cleaning unit with the following verification tests:

- Verification of sufficient cavitation energy with the SonoCheck™ monitoring vials*. Designed to change color (blue/green to yellow) when the ultrasonic cleaner is supplying sufficient energy and conditions are correct (degassed water, temperature, etc.) the SonoCheck™ is an easy to use and interpret method for monitoring cavitation energy. Failure to change color indicates that either the sonic bath conditions were not correct, or a failure of one or more of the ultrasonic transducers.

**PROCEDURE:**

"An risk assessment should be completed for any issue with any aspect of decontamination that can pose a risk to personnel or patients. The risk assessment should define and resolve the issue, and the system should be monitored to ensure that the issue has been corrected"(21)

*Directions for use:

Daily Inspection & Testing:

- Follow manufacture guidelines concerning the daily inspection of equipment (screens, tank condition, filters…).
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- Remember each Sonic unit is different, and staff should be trained on operation and inspection and cleaning of the equipment at least yearly.
- Inspect the level of the detergent daily (mark the container of the solution daily with the date at the level of the solution in the container) this will allow a visual inspection if the solution is actual being used.
- Log all observations in a report, as daily observations.
- Report any concerns to the proper management staff within the department to address.

**Types of Testing of the Sonic Cleaner with the SonoCheck™**

The functional test (OQ) will check the uniform operation of the empty ultrasonic cleaner’s tank. This testing should be done on installation of the equipment and or after major repairs. This would be considered the OQ in a QMS. The diagram below gives the suggested placement of SonoChecks™ in relation to the sonic tank size.

Remember that degassing should always be done before any testing cycle begins. Record all results for trend analysis and for help in any troubleshooting issues.

- Small up to 5ltrs or (1.5 gals)
- Medium size tank 5 to 20ltrs or (1.6-5 gals)
- Large above 20ltrs or greater than 5 gals.

**Routine Testing of the Sonic Cleaner**
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The routine test (PQ) will also monitor the performance of the sonic cleaner. The routine test is performed under normal conditions in an empty tank that has been degassed. Frequency of testing should be each day it is used. The diagram below gives the suggested placement of SonoCheck™ in relation to the sonic tank size for routine testing. All testing results should be logged and saved for trend analysis and troubleshooting concerns.

- In case of unsatisfactory results of the SONOCHECK™, please refer to the troubleshooting guide
- A more detailed testing may be required based on the results (a more routine testing and/or a functional testing, may be required)
- Record all information in logbook

Because there are many different ultrasonic cleaners on the market for medical facilities to purchase checking for cavitation only according to the standards might not be enough to ensure an ultrasonic cleaner is working properly as described in ANSI / AAMI ST 79 some other test might be needed to be performed daily or when the unit is used such as:

**Test for soil removal (external) in ultrasonic bath:**
**Test for soil removal (internal within lumens) in ultrasonic bath**

**Equipment Daily Inspection:**
- Inspect equipment (screens…) according to the manufacture instructions. Clean as needed.
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Weekly Inspection and Testing of other factors that can impact the Sonic cleaning process

We recommend that these tests are done at least weekly or after any major repair because they are important factors that help in ensuring that the sonic cleaner is working properly.

- these are helpful test of the various inputs of the ultrasonic process (temperature, pH, hardness).

Maintenance on Equipment (6,10):
- After any maintenance on the equipment, perform a test using the SonoCheck™ to ensure that the equipment is functioning properly.
- Follow the OQ test process.
- Have the maintenance person wait until the test results are complete

RESPONSIBILITY:

Central Service personnel are responsible for the proper use, result interpretation, and documentation of the Sonic Test Kit when used on a sonic cleaner. (1,5,16)

In-service and training of the staff should be done at least yearly on the equipment (ultrasonic) and the use of the SonoCheck™.
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**SonoCheck™ trouble-shooting guide**

If the SonoCheck ultrasonic cavitation monitor does not change color or if the time required to generate the color change takes longer than normal, please check the following guide:

<table>
<thead>
<tr>
<th>Problem</th>
<th>Reason</th>
<th>Corrective action</th>
</tr>
</thead>
<tbody>
<tr>
<td>De-gassing</td>
<td>Dissolved gasses will absorb ultrasonic energy</td>
<td>De-gas solution according to equipment manual</td>
</tr>
<tr>
<td>Water level</td>
<td>Ultrasonic energy may reflect off the surface of the solution and change energy distribution</td>
<td>Check equipment manual for correct water level</td>
</tr>
<tr>
<td>Operating cycle time</td>
<td>Time varies with the amount of ultrasonic energy available</td>
<td>Longer operating cycles generally provide better results</td>
</tr>
<tr>
<td>Instrument load</td>
<td>Heavy instrument loading and certain materials can absorb ultrasonic energy</td>
<td>Look for weak points using the periodic functional test and check for ultrasonic absorbent material like silicone or plastics</td>
</tr>
<tr>
<td>Transducer failure</td>
<td>Transducer efficiency may decrease with age. Individual transducers may fail while others in the equipment continue to function</td>
<td>Perform periodic functional test, placing SonoCheck™ monitors in each transducer location (see equipment manual)</td>
</tr>
<tr>
<td>Low energy</td>
<td>Transducer inefficiency or the ultrasonic basket may absorb too much energy</td>
<td>Check performance without basket in place. Compare performance against another ultrasonic cleaner if available. Call for service</td>
</tr>
</tbody>
</table>
REFERENCES:

1. ANSI/AAMI ST79; 2017
2. Blood as a Soil on Surgical Instruments; Cleaning Profile, Cleaning, Detection; M. Pfeifer, Zentr Steril 1998;6 (6):381-385
4. OSAKA REPORT; Importance of the cleaning test; University of Osaka, Department of Medicine, Ryo Fushimi, 2000
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