1. **Scope**

1.1 This methodology is to be used to perform the VCS STD 5711-102 Salt Spray test standard in an Ascott corrosion chamber.

This should be used in conjunction with the VCS STD 5711-102 standard document. The test standard takes precedence over this method statement and this method may need to be altered to follow/comply with the standard.

1.2 This method is based on VCS STD 5711-102 which consists of:

- Continuous salt spray test exposure at 35°C +/-2°C with a salt concentration of 5% NaCl +/-1%.
- Three variations within the standard give three options:
  - NSS – Neutral Salt Spray
  - AASS – Acetic Acid Salt Spray
  - CASS – Copper-accelerated Acetic acid Salt Spray (50°C +/-2°C).

1.3 The chamber will be loaded with test samples as required by the customer (or in accordance with VCS STD 5711-102).

2. **Instrumentation**

2.1 All measuring equipment must be calibrated. The recalibration renewal date must not fall within the test duration.

2.1.1 The Ascott corrosion chamber should be calibrated for chamber air temperature and relative humidity as a minimum.

If required, the following ‘chamber’ items may also be calibrated:

- Chamber air saturator temperature.
- Chamber air pressure gauge (atomiser pressure).

2.2 Peripheral devices also should be calibrated prior to use and may include the following:

- **Hand Held pH Meter** (Ascott Accessory No: ACC11)
  
  Digital pH meter, for measuring the pH of salt solution fallout over range 0-14 pH with a resolution of 0.01 pH. Supplied complete with buffers for calibration. View all our accessories at www.ascott-analytical.com

- **Salinity Refractometer** (Ascott Accessory No: ACC100)
  
  A salinity refractometer optimized to give a direct reading of percentage sodium chloride in the range 0 to 28%, with automatic temperature compensation. View all our accessories at www.ascott-analytical.com

- **Conductivity meter** is calibrated using standard solution, used for checking the conductivity of the water used for the salt solution.

2.3 The chamber temperature may be continuously monitored if required, using an independently calibrated data logger. For salt spray testing, it may be satisfactory to record the chamber temperature manually using the Ascott chamber display every day.

2.4 **Exposure to Salt Solution**

Collection rates are monitored manually using collection vessels placed at sample height. The collection rates are to be within the range of range of 1-2ml/hr/80cm².
2.5 The salt solution exposure is by means of atomisation using compressed air. The air delivered to the spray nozzle must be ‘heated and moistened’ by passing the air through an air saturator, the temperature of the chamber air saturator is set ‘several degrees higher than the chamber’.

3. Salt Solution Preparation

3.1 Salt solution to be prepared in accordance with VCS STD 5711-102.

- Check that the water conductivity is measured and monitored and is within the requirements of the standard using a conductivity meter. (Less than 20 µS/cm at 25 °C ± 2 °C).
- Salt solution concentration is measured & monitored and is within the requirements of the standard using a calibrated Salinity Refractometer.
- Salt solution pH is measured & monitored and is within the requirements of the standard using a calibrated pH Meter.

3.1.1 NSS - After allowing the solution to stabilise for several hours, the salinity and pH is measured and recorded. Any adjustments to the pH can be made using reagent grade hydrochloric acid (HCL) to increase the acidity or reagent grade Sodium hydroxide (NaOH) to reduce the acidity. Record all results.

3.1.2 AASS – Glacial Acetic acid Salt Spray is added to the salt solution so that the pH meets the required limits stipulated within the standard. Any adjustments to the pH can be made using Glacial Acetic Acid to increase the acidity or reagent grade Sodium hydroxide (NaOH) to reduce the acidity. Record all results.

3.1.3 CASS – Copper (II) Chloride Dihydrate (CuCl2 2H2O) is added to the salt solution to give a concentration of 0.26g/L (+/- 0.01g/L).

Any adjustments to the pH should be made using Glacial Acetic Acid to increase the acidity or reagent grade Sodium hydroxide (NaOH) to reduce the acidity. Record all results.

4. Sample Preparation

4.1 The test samples should be thoroughly cleaned before testing commences. This should not include the use of abrasives or solvents. This process should be agreed with the customer.

Latex gloves must be worn at all times when handling samples.

Photographs should be taken of each sample prior to starting the test.

5. Operation

5.1 Pre-test evaluation.

Refer to the test standard document to create a test profile for the chamber.

- Run a 24 hour cycle of the test with the chamber empty and collection funnels positioned, record the temperature and ensure it remains in tolerance of 35°C/+-2°C (50°C/+-2°C for CASS).
- Ensure the salt fog collection rates are within the expected range of 1.2ml/hr/80cm². Record all results.
- Check that the collected solution pH falls within the requirement of the standard. Record all results.
- If required, adjust the pH of the salt solution within the solution reservoir to offset any change to the pH when collected, so that the collected, the solution is within requirements of the standard.
• This may require additional periods of 24hr testing to prove results before testing with samples commences.
• It may be required to evaluate the chamber corrosivity before commencing a test. Refer to the test standard for full details.

5.2 Starting the test cycle

5.2.1 Test Exposure Conditions

- Position samples within the chamber in accordance of the test standard.
- Set the chamber air saturator temperature to be several degrees higher than the chamber set temperature.
- Ensure that no samples ‘shadow’ other samples and that droplets from one sample cannot fall onto other samples.
- Start the test cycle and record test parameters at start.
- Spray continuously with atomised salt solution at a constant chamber temperature of 35°C +/-2°C (50°C +/-2°C for CASS).
- Exceptions to continuous testing are permitted to evaluate the samples periodically. This should be kept to a minimum.
- Photographs to be taken prior to starting the test and at customer specified times.

5.3 Quality Control

5.3.1 Daily checks to ensure the standard is being followed with variable parameters within limits - Record all parameters.

- Check that all atomiser air pressure is within acceptable limits
- Check that atomiser air pressure is within acceptable limits
- Check that the reservoir salt solution is within 5.0% +/-1.0% NaCl
- Check that salt solution in the reservoir pH is within acceptable limits.
- Record the conductivity of DI water when used.
- Monitor the level of salt solution in the reservoir and ensure that there is enough for the next 24/48 hours. (Allow extra for weekends).

5.4 After Exposure

5.4.1 Remove the samples from the chamber and allow them to dry for 0.5 h to 1 h then rinse or dip the samples gently in clean running water at a temperature not exceeding 40°C and then dry them immediately in a stream of air (at a pressure not exceeding 200 kPa and at approximately 300 mm). Other methods may be agreed with the customer.

5.5 Deviation Handling

5.5.1 General deviations such as downtime, out of tolerance recordings should be noted in the test report, including details of any alterations made.

For further information, please contact us.

Ascott Analytical Equipment Limited
6-8 Gerard, Lichfield Road Industrial Estate, Tamworth, Staffordshire, B79 7UW, Great Britain
T +44 (0)1827 318040
F +44 (0)1827 318049
E sales@ascott-analytical.com
W www.ascott-analytical.com

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## Typical Daily Checks

<table>
<thead>
<tr>
<th>Hours</th>
<th>Chamber Temp °C</th>
<th>Air Sat Temp °C</th>
<th>Collected mi/hr (2 vessels per atomiser)</th>
<th>Collected Solution pH</th>
<th>Reservoir Salinity %</th>
<th>Pump Speed</th>
<th>Atomiser Air Pressure PSI</th>
<th>Initials</th>
<th>Photos Taken</th>
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