This document provides information on the test method that is used by the U.S. Consumer Product Safety Commission’s (CPSC) testing laboratory (LSC) in the analysis of paint and certain painted products. The method is used to determine the total lead content of paint, or a painted surface, on a dry paint basis. This method supersedes all previously published standard operating procedures for lead in paint testing. Existing accreditations remain valid. The rules for accreditation for lead in paint testing for compliance to the Ban of Lead-Containing Paint and Certain Consumer Products Bearing Lead Containing Paint, Code of Federal Regulations, Title 16, Part 1303 (16 C.F.R §1303) remain unchanged, and do not explicitly require the use of this or any earlier standard operating procedure (http://www.access.gpo.gov/nara/cfr/waisidx_08/16cfr1303_08.html).

This method is provided to inform interested parties of the method used by LSC for assessing the total lead in paint and other surface coatings. Other laboratories making such assessments are not required to follow this method; however, other laboratories should consider using these procedures to ensure they obtain results that are consistent with CPSC staff’s for purposes of compliance to 16 C.F.R. §1303.

CPSC staff has concluded that this test method is sufficient to make appropriate determinations concerning lead in paint, as defined in 16 C.F.R §1303. Screening tests by x-ray fluorescence may sometimes be employed by CPSC staff to determine samples in need of such testing.

Definitions:

1. Sample – an individual consumer product or a group of identical consumer products from a batch to be tested.
2. Component Part – an individual sub-unit within the total sample. Each separate paint color on a sample is a component part.

3. Composite Testing-like parts – combining like paints from several like parts or products to obtain sufficient sample size for analysis when there is an insufficient quantity of paint on one item to perform testing.

4. Composite Testing-different parts – combining different paints (e.g., multiple colors) from one or more samples to reduce the number of digestions and instrumental lead analyses performed.

5. Instrument Detection Limit (IDL) – 3 times the standard deviation of 10 replicate measurements of reagent blank. The IDL for Pb on the Inductively Coupled Plasma Optical Emission Spectrometer (ICP-OES) used by CPSC staff is 0.01 µg/ml.

6. Method Detection Limit (MDL) – reagent blank fortified with 2-3 times the IDL. Seven replicate measurements are made. Calculate the MDL as follows: \[ MDL = t \times S, \quad t = 3.14 \] (99% confidence level for 7 replicates), \( S = \) standard deviation. The MDL determined for Pb is 0.01 µg/ml.

7. Laboratory Reagent Blank (LRB) – an aliquot of the digestion reagents that is treated exactly as a sample including exposure to glassware, digestion media, apparatus, and conditions used for a particular Pb test, but with no added sample. LRB data are used to assess contamination from the laboratory environment.

8. Calibration Blank – deionized water acidified with nitric acid (3 ml concentrated nitric acid diluted to 100 ml with deionized water).

9. Stock Standard Solution – 1000 µg/ml solution of Pb purchased from reputable commercial source, used to prepare calibration standards. Replace before expiration date.

10. Calibration Standards – solutions containing 0 to 25 µg/ml of Pb in 3% nitric acid matrix are used. A minimum of 4 calibration standards are used. Calibration standards should be prepared weekly.

11. Quality Control Sample (QCS) – a solution containing Pb that is used to evaluate the performance of the instrument system. QCS is obtained from a source external to the laboratory and Stock Standard Solution.

12. Certified Reference Material (CRM) – CRMs are materials with similar matrices as test samples with known lead levels. The CRMs are used to verify digestion and analysis methods. For example, standard reference materials (SRMs) are CRMs that are available from the National Institute of Standards and Technology (NIST), such as those listed in the Equipment and Supplies section below.

**Equipment and Supplies:** The materials used for sampling and analysis are as follows:

1. Nitric acid, trace metal grade
2. Disposable plastic digestion vessels, 50 ml, or glass test tubes
3. Hot block digester or hot plate with test tube block
4. Disposable razor blade or scalpel
5. Methylene chloride (optional)
6. Distilled water
7. Microwave digestion apparatus
8. CRMs such as NIST SRMs² 2581 and 2582 lead in powdered paint

² NIST SRMs 2581 and 2582, and other reference materials, are available from the National Institute of Standards and Technology. See [http://ts.nist.gov/measurementservices/referencematerials/index.cfm](http://ts.nist.gov/measurementservices/referencematerials/index.cfm)
9. Internal Standard (such as yttrium, from a stock standard solution of that element appropriate to the instrument parameters of the ICP used for the analysis)

Note on composite testing: Composite testing-like parts (as defined above) is appropriate and may be necessary to obtain valid analytical results. Composite testing-different parts (as defined above) must be done with adequate care, planning, and understanding of the limitations and propagations of error in measurements or the test may fail to detect excessive lead in one individual paint because of dilution. If composite testing-different parts is used, great care must be taken to accurately weigh each component paint and account for the propagation of errors due to weighing and measuring, and the analyst must ensure the compositing will not result in a Pb measurement being below the detection limit for a paint component that would exceed 0.009% Pb.

Each individual paint must be weighed individually with sufficient precision and sufficient safety factors to assure that no false negatives are reported.

Sufficient paint must be used for each of the paints giving proper consideration for the weighing capabilities of the balance used and the detection limits and necessary dilution for the subsequent instrumental analysis. The combined paints can then be digested according to the procedure given below for a single paint, before analysis by Inductively Coupled Plasma (ICP) spectroscopy. Appropriate weights for each of the individual paints which are composited depend on final dilution volumes, weighing accuracy, and detection limits. All of the lead found in the digestate for the combined composite can be attributed to the weight of each of the individual paints as though all of the lead came from any one of the individual paints. The measured lead should be treated in calculations as if it originated solely from one part of the composite; the lead concentration calculation should be carried out for each part unless identical weights of each sample are used in the analysis. An example is given at the end of this document.

In considering results from such a composite sample, it is imperative that a sufficient “safety factor” be applied to account for weighing inaccuracy and propagation of errors from each step in the analytical procedure to ensure no violative paints are misrepresented as non-violative. It has been suggested, for example for a composite of 3 samples, that any sample possibly having greater than 80% of the limit for lead in paint in any of the composited paints should be retested. CPSC staff considers this a reasonable practice.

Method: The digestion method is based on the Association of Official Analytical Chemists (AOAC) standard AOAC 974.02 (Lead in Paint). Alternate microwave digestion based on ASTM E1645 may be used as well. Analysis by Inductively Coupled Plasma (ICP) spectroscopy is based on ASTM E1613.

3 Toy Industry Association Inc. Laboratory Testing Technical Work Group, that includes (but is not limited to) representatives of the following laboratories: Specialized Technology Resources (STR), SGS consumer Testing Services; EMC Inc., Bureau Veritas (BV), CMA Testing NSF International, Intertek (ITS) and Toy Safety and Quality, Inc. Compositing Acceptance Method for Lead in Surface Coatings, Comment submitted to CPSC, March 18, 2009.

4 AOAC Official Method 974.02 Lead in Paint
1. For testing of wet paint, apply a thin coating to a glass slide, and dry completely prior to testing by heating in an oven at nominally 105 °C (105 °C ± 2 °C) until the weight is stable for at least two successive readings; the readings should be separated by 30 minutes of heating in the oven.

2. For products coated with paint or a similar surface coating, remove and digest the coating, separately from the substrate material. Care should be taken to remove as little of the substrate as possible. It may be necessary to add a few drops of solvent, such as methylene chloride, to soften the paint and aid in its removal from the substrate. If used, such solvent must be evaporated away prior to analysis. The scraped paint should be finely divided to help in digesting.

3. Scrape approximately 5-100 mg of paint from the product. If it is not possible to collect this much paint, it may be necessary to combine more than one unit of the product to collect sufficient paint.

4. Prepare a LRB and digest and test a standard reference paint material, such as National Institute of Standards and Technology (NIST) SRM 2581 - Powdered Paint Nominal 0.5% Lead or NIST SRM 2582 - Powdered Paint Nominal 0.02% Lead, with each batch of samples tested.

5. Digest appropriately according to AOAC 974.02 or ASTM E1645 in either a disposable glass test tube with a heating block, a disposable plastic digestion vessel in a hot block digester, or in a suitable digestion vessel and digestion microwave oven system.

6. Dilute samples so that Pb results are within the calibration range of the instrument. Be careful not to dilute a sample that could have 0.009% Pb to a level below the method detection limit.

7. Analyze diluted samples for Pb concentration using an ICP spectrometer (or Atomic Absorption spectrometer). Analysis procedures for ICP-OES, flame atomic absorption spectrometry (FLAA), and graphite furnace atomic absorption spectrometry (GFAA) are based on the methodology in ASTM E1613-04. Inductively coupled plasma mass spectrometry (ICP-MS) may also be employed with appropriate procedures, such as EPA 6020A\textsuperscript{7}.

**ICP Analysis Operating Procedures and Quality Control Measures:**

1. Ignite plasma. Perform wavelength calibration or torch alignments per instrument manufacturer recommendations.

2. Allow the instrument to become thermally stable before continuing.

3. Ensure the following element and wavelength are selected in analytical method:
   a. Pb 220.353
   One other Pb line such as Pb 217.00 should be used to ensure spectral interferences are not occurring during analysis.

4. An internal standard such as 2 µg/ml yttrium should be used.

5. Perform calibration using calibration blank and at least 3 standards. Calibration should be performed a minimum of once a day when used for analysis, or each time the

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\textsuperscript{5} ASTM E1645 \textit{Standard Practice for Preparation of Dried Paint Samples by Hotplate or Microwave Digestion for Subsequent Lead Analysis}

\textsuperscript{6} ASTM E1613 \textit{Standard Test Method for Determination of Lead by Inductively Coupled Plasma Atomic Emission Spectrometry (ICP-AES)}

\textsuperscript{7} EPA 6020A \textit{Inductively Coupled Plasma-Mass Spectrometry}
instrument is set up. Results for each standard should be within 5% of the true value. If the values do not fall within this range, recalibration is necessary.

6. Analyze the QCS immediately after the calibration. The analyzed value of Pb should be within ±10% of the expected value. If the Pb value is outside the ±10% limit, recalibration is required.
   a. At least one LRB must be analyzed with each sample set. If the Pb value exceeds 3 times the MDL, laboratory or reagent contamination should be expected. The source of the contamination should be identified and resolved before continuing analyses. The LRBs should be at the same acid concentration as that of the sample and should be taken through the same digestion procedure as that of a sample.

7. At least one certified reference material (CRM) should be analyzed with each batch of samples. The CRM should have a known amount of Pb and be of a material similar to a test specimen. Analyte recoveries should be within ±15% of expected values. If recoveries are outside this limit, the source of the problem should be identified and resolved before continuing analyses.

8. Dilute any samples that have Pb values exceeding 1.5 times the high calibration standard, and reanalyze.
Calculations and Results Reported: Results for the Pb test methods are calculated and reported as follows:

1. Total Pb concentration: \[ \% \text{ Pb (wt/wt)} = 0.10 \frac{\text{cd}}{\text{w}} \]
   a. \( c \) = concentration of Pb detected (µg/ml)
   b. \( d \) = dilution volume (ml)
   c. \( w \) = weight of aliquot digested (mg)

Example 1: Single Paint Tested

An aliquot of 10 mg of yellow paint scrapings is digested in acid and diluted to a final volume of 50 ml. When analyzed on the ICP-OES, the analytical result is 0.080 µg/ml lead in the acid. The paint therefore contained 0.10 x 0.080 µg/ml x 50 ml / 10 mg = 0.04% (400 ppm).

Table 1: Total Pb Analysis

<table>
<thead>
<tr>
<th>Item</th>
<th>Analytical result Pb (µg/ml)</th>
<th>Dilution volume (ml)</th>
<th>Total Pb (µg)</th>
<th>Sample wt (mg)</th>
<th>Pb (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Yellow Paint</td>
<td>0.080</td>
<td>50</td>
<td>4</td>
<td>10</td>
<td>0.04</td>
</tr>
</tbody>
</table>

Example 2: Composite Testing – Different Parts

One example of composite testing of different paints would be as follows, and considers the case of weighing to the nearest 0.01 mg, digesting in acid, diluting to a final volume of 10 ml, and testing on an ICP-OES with an MDL of 0.01 µg/ml. A sample bears red, green, and orange paint. The paints are removed from the sample and 4.90 mg of red paint is combined with 5.00 mg of green paint, and 5.10 mg of orange paint. The resulting 15 mg of composite paint from this example is digested in acid and diluted to 10 ml, and the diluted digest is found to contain 0.05 µg/ml lead. The combined 3 aliquots of paint contributed to a total of 0.50 µg of lead for the composite sample. Although the average concentration in this case would be 33 ppm, the individual contributions are not known, and one must calculate the lead concentration of each paint as if all of the lead originated from it. Thus, the red paint could contain up to 0.50 µg / 0.0049 g = 102 ppm (µg/g), with similarly calculated results of 100 ppm and 98 ppm for the green and orange paints. See Table 2 below.

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8 The factor 0.10 in this formula is the mathematical conversion of the units used for concentration (µg/ml), and weight of paint (mg) into percent, and represents 100% x 1 mg / 1000 µg.
### Table 2: Total Pb Analysis – Composite Testing

<table>
<thead>
<tr>
<th>Item</th>
<th>Analytical result Pb (µg/ml)</th>
<th>Dilution volume (ml)</th>
<th>Total Pb (µg)</th>
<th>Sample wt (mg)</th>
<th>Potential Pb (%) per component</th>
<th>Pb (%) Composite</th>
</tr>
</thead>
<tbody>
<tr>
<td>Red Paint</td>
<td>*0.05</td>
<td>10</td>
<td>0.5*</td>
<td>4.90</td>
<td>0.0102</td>
<td></td>
</tr>
<tr>
<td>Green Paint</td>
<td>*0.05</td>
<td>10</td>
<td>0.5*</td>
<td>5.00</td>
<td>0.0100</td>
<td></td>
</tr>
<tr>
<td>Orange Paint</td>
<td>*0.05</td>
<td>10</td>
<td>0.5*</td>
<td>5.10</td>
<td>0.0098</td>
<td></td>
</tr>
<tr>
<td>Total Composite</td>
<td>0.05</td>
<td>10</td>
<td>0.5</td>
<td>15.00</td>
<td>0.0033**</td>
<td></td>
</tr>
</tbody>
</table>

*In a composite of different component paints, the analytical result for the total composite would be applied to each component part as if all the Pb was in that component paint.

**In this example, the Pb concentration of the combined paints is <0.009%, but when estimating potential Pb per component, each component could potentially have >0.009% Pb and would require additional testing without compositing to ensure Pb <0.009%.