

### **Insoluble Residue in Carbonate Aggregates**

### 3. Significance and Use

3.1 The amount and size distribution of noncarbonate (insoluble) material in carbonate aggregates is of interest to those conducting research on the polish susceptibility of stone used in pavements or on the friction properties of pavement surfaces. In addition, some agencies utilize the test for acceptance purposes to delineate carbonate aggregates that polish excessively and become slippery and those that will not.

NOTE 1—The quality of the results produced by this standard are dependent upon the competence of the personnel performing the procedure and the capability, calibration, and the maintenance of the equipment used. Agencies that meet the criteria of Specification D3666 are generally considered capable of competent and objective testing/sampling/inspection, etc. Users of this standard are cautioned that compliance with Specification D3666 alone does not completely ensure reliable results. Reliable results depend on many factors: following the suggestions of Specification D3666 or similar acceptable guideline provides a means of evaluating and controlling some of those factors.

## 4. Apparatus

9.5-mm (3⁄8-in.)
4.75-mm (No. 4)
2.36-mm (No. 8)
1.18-mm (No. 16)
600-µm (No. 30)
300-µm (No. 50)
150-µm (No. 100)
75-μm (No. 200)

4.2 <i>Balance</i> —A balance having a minimum capacity of 15 kg and meeting the requirements of Guide
D4753, Class GP5
4.3 <i>Oven,</i> of appropriate size capable of being maintained at 110 <u>+</u> 5 °C
4.4 Agitation Equipment, such as a jar mill or a magnetic stirrer
4.5 Bunsen Burner or Hot Plate
4.6 <i>Receiving Tank,</i> of nonreactive material to receive wash water and residue washed through the 8-in.
diameter sieves
4.7 Rapid Filter Paper, acid-resistant
4.8 Safety Eyeglasses, Rubber Gloves, Aprons, and so forth
4.9 Borosilicate Glass Containers, for performing acid leaching test
4.10 pH Paper or pH Meter

### 5. Reagent

5.1 Purity of Reagents—Reagent grade chemicals shall be used in all tests
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#### 6. Samples

6.1 Samples shall be representative of the sources from which they are obtained and shall be reduced to
appropriate size by use of sample splitters or by quartering in accordance with the Sampling section of
Test Method C136
6.1.1 Normally, the aggregate used in the test sample shall be retained after washing over the 4.75-mm
(No. 4) sieve, and passing the 9.5-mm (3/8-in.) sieve
6.1.2 Test separate fractions or combined aggregate gradings as appropriate
6.2 Dry sample aggregates to constant mass at 110 <u>+</u> 5 °C
6.3 Use an oven-dry sample having a minimum mass of 500 g

### 7. Procedure for Determining Only the Plus 75-µm (No.200) Size Fraction of Insoluble Residue

7.1 For many applications, only the plus 75-μm (No. 200) size insoluble residue fraction is of interest.
This section provides a procedure for determination of the amount and size distribution of only the plus
75-μm (No. 200) size insoluble residue
If the total amount of insoluble residue is desired, follow the procedure in Section 8 and only such
portions of Section 7 as are indicated in Section 8
7.2 Prepare and test samples in triplicate for each aggregate sampled
7.3 Warning—Hydrogen chloride gas is released during this procedure. Exercise extreme care to prevent
corrosive gas from entering the laboratory atmosphere
The hydrogen chloride gas that may be produced in the chemical reaction can be destructive to
laboratory equipment and can also be a definite safety hazard to laboratory personnel. Under no
circumstances shall the test be performed without proper and adequate ventilation
Operators shall be required to wear appropriate personal protective equipment (PPE) when performing
this test
7.4 Place an aggregate sample weighing a minimum of 500 g in a suitable borosilicate glass container
The container shall be vented to allow gas from the reaction to escape and to be drawn away
Slowly add 1000 mL of 6 N hydrochloric acid (HCl) allowing for excessive effervescence or foaming to
subside before further addition of solution
Agitate the container and contents gently by hand until excessive effervescence has subsided
7.5 Place the container and contents on an agitating device
7.6 Check the container periodically, and when the reaction appears to have subsided (no obvious
bubbling), decant the solution and add approximately 300 mL of the 6 N HCl and check for a reaction
Collect all decant solution in a receiving tank for future use only if determination of the total residue
content is desired as in Section 8
7.6.1 If a reaction (bubbling) is observed, continue agitation until the reaction subsides and repeat the
procedure beginning at 7.6.

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7.7 When no reaction is observed, heat the container gently over a Bunsen burner or hot plate
Some carbonate materials will require heating to temperatures of up to 110 °C and maintaining this
temperature level for approximately 1 h in order to completely react the carbonate fraction
7.7.1 If no reaction is observed upon heating, proceed with the remainder of the test procedure
7.7.2 If a reaction is observed upon heating, continue the agitation and repeat the procedure beginning at 7.7.
7.8 When the reaction has stopped, decant the acid solution and add water to the aggregate in the glass container to thoroughly dilute the acid
The residue in the container shall have a pH as indicated by pH paper or pH meter of more than 5.5
7.9 Wash the diluted solution and residue from the aggregate sample over a preweighed 75-µm (No. 200) sieve placed over a receiving tank
200) sieve placed over a receiving tank
7.10 Thoroughly rinse the material on the 75- $\mu$ m (No. 200) sieve and into the receiving tank. Dry the
sieve and the residue in a 110 <u>+</u> 5 °C oven
7.11 When the sieve and residue have cooled to room temperature, weigh the sieve plus the residue
and subtract the original mass of the dry clean sieve to obtain the mass of the plus 75-μm (No. 200) sieve size residue
7.12 Visually inspect the residue to determine if there is particle agglomeration because of the presence
of plastic clay fractions
7.12.1 If agglomeration has occurred, physical reduction of the agglomerated particles shall be
performed by use of a mortar and rubber-tipped pestle. Exercise care to prevent crushing of individual
particle grains during this process
7.12.2 If no agglomeration has occurred, proceed with the remainder of the test procedure
7.13 Dry sieve the residue from the aggregate sample over preweighed nested sieves
The nested sieves shall be of the following series:



# 8. Procedure for Determining the Total Acid Insoluble Residue Content

8.1 Perform the steps in 7.1 – 7.8
8.2 Wash the diluted solution and residue from the aggregate sample over pre-weighed nested sieves
placed over a receiving tank
The nested sieves shall be of the following series:



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4.75-mm (No. 4)
2.36-mm (No. 8)
1.18-mm (No. 16)
600-µm (No. 30)
300-µm (No. 50)
150-µm (No. 100)
75-μm (No. 200)

8.3 Rinse each sieve with water individually and thoroughly over the lower sieves and into the receiving tank..... Dry the sieves and the residue in a 110 + 5 °C oven..... 8.4 When the sieves have cooled to room temperature, weigh each sieve plus the residue and subtract the original mass of the dry, clean sieve to obtain the mass of the residue..... 8.5 Dilute the decant acid solution in the receiving tank with water. The solution in the receiving tank shall have a pH as indicated by pH paper (or a pH meter) of more than 5.5..... 8.6 Pour the solution contained in the receiving tank through a 75- $\mu$ m (No. 200) sieve and then through rapid filtering paper placed in glass funnels..... Thoroughly rinse with additional water..... Pre-weigh the filter papers and the evaporating dish..... 8.7 When filtration is complete, rinse by filtering clean water, then transfer the paper and the residue into an evaporating dish and place in the oven at 110 ± 5 °C for 24 h..... 8.8 Weigh the dish, dried filters, and residue..... Subtract the mass of the clean filters and dish from the mass of the dish, filters, and residue to obtain the mass of the residue passing the 75-μm (No. 200) sieve..... 8.9 Dry the 75- $\mu$ m (No. 200) sieve and residue obtained during filtering process (see 8.6) in a 110  $\pm$  5 °C oven..... 8.10 When the 75-μm (No. 200) sieve and residue have cooled to room temperature, weigh the sieve plus the residue and subtract the original mass of the dry, clean sieve to obtain the mass of the residue (if any) retained on the 75-μm (No. 200) sieve from the decant solution..... Add this quantity to the quantity established in 8.2. 8.11 Examine the residue and describe it with the aid of such mineral and rock identification tests as are necessary and appropriate. Refer to Descriptive Nomenclature C294.....

### 9. Calculation



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## 10. Report

- 10.1 Report the following information:
- 10.1.1 Identification and description of the aggregate samples......

   10.1.2 Calculated results for each sample and average results for triplicate tests.....

#### Comments