



# Standard Practice for Preparing Test Specimens from Basic Refractory Castable Products by Casting<sup>1</sup>

This standard is issued under the fixed designation C 974; the number immediately following the designation indicates the year of original adoption or, in the case of revision, the year of last revision. A number in parentheses indicates the year of last reapproval. A superscript epsilon ( $\epsilon$ ) indicates an editorial change since the last revision or reapproval.

## 1. Scope

1.1 This practice covers a procedure for preparing test specimens from a basic refractory castable by casting prepared material in a mold. Specimens prepared in accordance with this procedure are intended for use in ASTM test methods.

1.2 *This standard does not purport to address all of the safety problems, if any, associated with its use. It is the responsibility of the user of this standard to establish appropriate safety and health practices and determine the applicability of regulatory limitations prior to use.*

## 2. Significance and Use

2.1 This practice defines a procedure that ensures consistent preparation of specimens for product testing and evaluation.

2.2 This practice can be used in the laboratories of producers, users, and general-interest parties for research and development or quality-control work. It is particularly useful for interlaboratory comparisons or products, for repetitive evaluations or comparisons of products or product quality, and in specifying a uniform preparation practice for specimens for acceptance testing.

2.3 In using this practice it must be recognized that the structure of laboratory-cast specimens may differ significantly from the structure of material cast in field applications.

## 3. Apparatus

3.1 *Mixer*, bench-type, Hobart or equivalent.

3.2 *Steel Molds*.

3.3 *Vibration Table*, spring or air-cushioned, with a pneumatic or electric driver.<sup>2</sup>

3.4 *Oven*, air circulating.

<sup>1</sup> This practice is under the jurisdiction of ASTM Committee C08 on Refractories and is the direct responsibility of Subcommittee C08.09 on Monolithics.

Current edition approved March 1, 2004. Published March 2006. Originally approved in 1982. Last previous edition approved in 1998 as C 974 – 87 (1998).

<sup>2</sup> Martin Engineering Model CV35 or Syntron Model VP-80 has been found suitable.

## 4. Procedure

4.1 Store the material to be tested at  $85 \pm 2^\circ\text{F}$  ( $29.5 \pm 1^\circ\text{C}$ ) for 24 h prior to testing.

4.2 Reduce the sample to the desired batch size with a sample splitter or by quartering, taking precautions to prevent segregation. Sufficient material should be batched to provide at least a 10 % excess over test specimen requirements.

4.3 Add the weighed, dry batch to the mixer and dry-mix at low speed for 30 s. Continue to mix at low speed while adding the required amount of water (see 4.3.1) within 30 s. Wet-mix at low speed for 4 min.

4.3.1 Use the amount of water and mixing time (if different from above) recommended by the manufacturer. The water must be potable and at a temperature of  $85 \pm 2^\circ\text{F}$  ( $29.5 \pm 1^\circ\text{C}$ ). Measure the water addition to the nearest 0.1 % by weight.


4.4 Specimens should be cast in a mold immediately after the batch is mixed so that all specimens made from a batch are cast at the same time. The mold should be securely clamped to the vibration table top. Fill the mold to overflowing distributing the material uniformly between each cavity. Vibrate in accordance with the manufacturer's instructions. Approximately 15 s before the end of the vibration period, remove excess material (approximately flush with top of mold).

4.4.1 If the batch is too wet, water will run off the specimens while if the batch is too dry, the material will not fill the mold. In either case the batch must be discarded and another run made at a suitable water content.

4.5 Test specimens should be air-dried in the molds a minimum of 16 h and a maximum of 24 h, and then removed from the molds and dried for a minimum of 8 h at  $225 \pm 5^\circ\text{F}$  ( $107 \pm 2.5^\circ\text{C}$ ) in a circulating air dryer prior to testing.

## 5. Calculation and Report

5.1 In the report on specimen preparation, include the percent water addition, drying time, and test specimen size.

 **C 974 – 87 (2004)**

*ASTM International takes no position respecting the validity of any patent rights asserted in connection with any item mentioned in this standard. Users of this standard are expressly advised that determination of the validity of any such patent rights, and the risk of infringement of such rights, are entirely their own responsibility.*

*This standard is subject to revision at any time by the responsible technical committee and must be reviewed every five years and if not revised, either reapproved or withdrawn. Your comments are invited either for revision of this standard or for additional standards and should be addressed to ASTM International Headquarters. Your comments will receive careful consideration at a meeting of the responsible technical committee, which you may attend. If you feel that your comments have not received a fair hearing you should make your views known to the ASTM Committee on Standards, at the address shown below.*

*This standard is copyrighted by ASTM International, 100 Barr Harbor Drive, PO Box C700, West Conshohocken, PA 19428-2959, United States. Individual reprints (single or multiple copies) of this standard may be obtained by contacting ASTM at the above address or at 610-832-9585 (phone), 610-832-9555 (fax), or [service@astm.org](mailto:service@astm.org) (e-mail); or through the ASTM website ([www.astm.org](http://www.astm.org)).*