



Designation: E1093 – 91 (Reapproved 2019)

Standard Specification for Glass Prothrombin Pipet, Disposable¹

This standard is issued under the fixed designation E1093; 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 (ϵ) indicates an editorial change since the last revision or reapproval.

1. Scope

1.1 This specification covers a glass disposable Prothrombin pipet suitable for use in micro techniques for estimation of Prothrombin time.

1.2 The values stated in SI units are to be regarded as standard. No other units of measurement are included in this standard.

1.3 This precautionary statement pertains only to the test method portion, Section 8, of this specification. *This standard does not purport to address all of the safety concerns, if any, associated with its use. It is the responsibility of the user of this standard to establish appropriate safety, health, and environmental practices and determine the applicability of regulatory limitations prior to use.*

1.4 *This international standard was developed in accordance with internationally recognized principles on standardization established in the Decision on Principles for the Development of International Standards, Guides and Recommendations issued by the World Trade Organization Technical Barriers to Trade (TBT) Committee.*

2. Referenced Documents

- 2.1 *ASTM Standards*:²
E438 Specification for Glasses in Laboratory Apparatus
- 2.2 *ISO Standard*:³
1769 Laboratory Glassware—Pipettes—Color Coding

3. Terminology

- 3.1 *Definitions of Terms Specific to This Standard*:
 - 3.1.1 *accuracy*—the expected distribution of mean volumes around the stated volume.

¹ This specification is under the jurisdiction of ASTM Committee E41 on Laboratory Apparatus and is the direct responsibility of Subcommittee E41.01 on Laboratory Ware and Supplies.

Current edition approved Jan. 1, 2019. Published January 2019. Originally approved in 1986. Last previous edition approved in 2012 as E1093 – 91(2012). DOI: 10.1520/E1093-91R19.

² For referenced ASTM standards, visit the ASTM website, www.astm.org, or contact ASTM Customer Service at service@astm.org. For *Annual Book of ASTM Standards* volume information, refer to the standard's Document Summary page on the ASTM website.

³ Available from International Organization for Standardization (ISO), ISO Central Secretariat, BIBC II, Chemin de Blandonnet 8, CP 401, 1214 Vernier, Geneva, Switzerland, http://www.iso.org.

3.1.2 *coefficient of variation*—the expected distribution of individual volumes around the mean volume.

3.1.3 *disposable*—Prothrombin pipets which are intended to be used once only and then discarded.

NOTE 1—Such pipets will only be expected to provide their specified performance during the original operation.

4. Classification

4.1 This specification covers only one glass pipet as illustrated in Fig. X1.1.

5. Materials and Manufacture

5.1 The pipet shall be made of borosilicate glass, Type 1; Class B, or soda lime glass, Type 2, in accordance with Specification E438.

6. Physical Properties

6.1 *Design*—The Prothrombin pipet shall be made of one piece construction glass tubing that is straight and with uniform bore and lightly firepolished on both ends. The pipet shall be made to the dimensions as specified in Fig. X1.1.

6.2 *Dimensions*—The pipet shall be made of tubing with a minimum outside diameter (o.d.) of 2.3 mm with an inside diameter (i.d.) of 1.7 mm. The uniformity of the bore shall be ± 0.05 mm throughout the straight portion of the pipet. The pipet shall be a minimum of 160 mm long.

6.3 *Capacity*—The pipet shall be calibrated “to deliver” (T.D.) 0.1 and 0.2 mL at 20 °C. Marking shall be as specified in 6.5.

6.3.1 *Accuracy* (see 3.1.1)—The accuracy from stated volume shall be ± 2.0 % for the 0.1 and 0.2-mL capacity and shall be determined as specified in 8.1.

6.3.2 *Coefficient of Variation* (see 3.1.2)—The coefficient of variation from stated volume for the 0.1 and 0.2-mL capacity shall not exceed 2.0 % and shall be determined as specified in 8.1.

6.4 *Graduation Lines*—The pipet shall be calibrated and marked with graduation lines at 0.1 and 0.2 mL from the tip of the pipet. The graduation lines shall be 0.3 ± 0.1 mm and shall completely encircle the tube.

6.5 *Pipet Nomenclature*—The pipet shall be marked with 0.1 and 0.2-mL markings slightly above the graduation lines.