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Designation: D7905/D7905M - 14 D7905/D7905M - 19

Standard Test Method for Determination of the Mode II Interlaminar Fracture Toughness of Unidirectional Fiber-Reinforced Polymer Matrix Composites¹

This standard is issued under the fixed designation D7905/D7905M; 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 test method describes<u>covers</u> the determination of the mode II interlaminar fracture toughness, G_{IIc} , of unidirectional fiber-reinforced polymer matrix composite laminates under mode II shear loading using the end-notched flexure (ENF) test (Fig. 1).

1.2 This method is limited to use with composites consisting of unidirectional carbon-fiber- and glass-fiber-reinforced laminates. This limited scope reflects the experience gained in round robin testing. This test method may prove useful for other types and classes of composite materials; however, certain interferences have been noted (see Section 6).

1.3 The values stated in either SI units or inch-pound units are to be regarded separately as standard. The values stated in each system mayare not benecessarily exact equivalents; therefore, to ensure conformance with the standard, each system shall be used independently of the other. Combiningother, and values from the two systems may result in non-conformance with the standard.shall not be combined.

1.3.1 Within the text the inch-pound units are shown in brackets.

1.4 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 safety, health, and health environmental practices and determine the applicability of regulatory limitations prior to use.

<u>1.5</u> 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

<u>TM D7905/D7905M-19</u>

2.1 ASTM Standards:² catalog/standards/sist/2d7a20c1-bb39-4b6a-8387-db93c48fe4a2/astm-d7905-d7905m-19

D792 Test Methods for Density and Specific Gravity (Relative Density) of Plastics by Displacement

D883 Terminology Relating to Plastics

D2584 Test Method for Ignition Loss of Cured Reinforced Resins

D2734 Test Methods for Void Content of Reinforced Plastics

D3171 Test Methods for Constituent Content of Composite Materials

D3878 Terminology for Composite Materials

D5229/D5229M Test Method for Moisture Absorption Properties and Equilibrium Conditioning of Polymer Matrix Composite Materials

D5687/D5687M Guide for Preparation of Flat Composite Panels with Processing Guidelines for Specimen Preparation D7264/D7264M Test Method for Flexural Properties of Polymer Matrix Composite Materials

E4 Practices for Force Verification of Testing Machines

E6 Terminology Relating to Methods of Mechanical Testing

E18 Test Methods for Rockwell Hardness of Metallic Materials

¹ This test method is under the jurisdiction of ASTM Committee D30 on Composite Materials and is the direct responsibility of Subcommittee D30.06 on Interlaminar Properties.

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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.

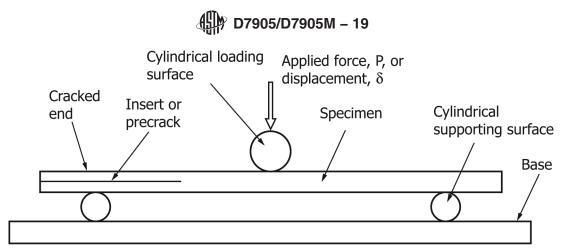


FIG. 1 ENF Test Fixture and Specimen Nomenclature

E122 Practice for Calculating Sample Size to Estimate, With Specified Precision, the Average for a Characteristic of a Lot or Process

E177 Practice for Use of the Terms Precision and Bias in ASTM Test Methods

E456 Terminology Relating to Quality and Statistics

E691 Practice for Conducting an Interlaboratory Study to Determine the Precision of a Test Method

E1309 Guide for Identification of Fiber-Reinforced Polymer-Matrix Composite Materials in Databases (Withdrawn 2015)³

E1434 Guide for Recording Mechanical Test Data of Fiber-Reinforced Composite Materials in Databases (Withdrawn 2015)³

E1471 Guide for Identification of Fibers, Fillers, and Core Materials in Computerized Material Property Databases (Withdrawn 2015)³

3. Terminology

3.1 Terminology D3878 defines terms relating to high-modulous fibers and their composites. Terminology D883 defines terms relating to plastics. Terminology E6 defines terms relating to mechanical testing. Terminology E456 and Practice E177 define terms relating to statistics. In the event of conflict between terms, Terminology D3878 shall have precendence over the other terminology standards.

NOTE 1—If the term represents a physical quantity, its analytical dimensions are stated immediately following the term (or letter symbol) in fundamental dimension form, using the following ASTM standard symbology for fundamental dimensions, shown within square brackets: [M][M] for mass, [L][L] for length, [T][T] for time, [u][u] for thermodynamic temperature, and [nd][nd] for non-dimensional quantities. Use of these symbols is restricted to analytical dimensions when used with square brackets, as the symbols may have other definitions when used without the brackets.

3.2 Definitions of Terms Specific to This Standard:

3.2.1 <u>Compliance Calibration compliance calibration</u> (CC) <u>Method—method</u> the method of data reduction where the relationship between specimen compliance $[T^2/M]$ and delamination length [L] is determined prior to testing by measuring specimen compliance $[T^2/M]$ at multiple simulated delamination lengths.

3.2.2 <u>Mode II Interlaminar Fracture Toughness, mode II interlaminar fracture toughness</u>, G_{IIc} [M/T^2]—the critical value of strain energy release rate, G, [M/T^2] for delamination growth [L] due to an in-plane shear force [M/T^2] or displacement [L] oriented perpendicular to the delamination front.

3.2.3 <u>Non-precracked non-precracked</u> (NPC) toughness $[M/T^2]$ —an interlaminar fracture toughness value that is determined from the preimplanted insert.

3.2.4 <u>Precracked precracked</u> (PC) <u>Toughness toughness</u> $[M/T^2]$ —an interlaminar fracture toughness value that is determined after the delamination has been advanced from the preimplanted insert.

3.2.5 <u>Strain Energy Release Rate, strain energy release rate</u>, $G [M/T^2]$ —the loss of strain energy, $dU [ML^2/T^2]$, in the test specimen per unit of specimen width [L] for an infinitesimal increase in delamination length, da [L], for a delamination growing self-similarly under constant displacement [L]. In]; in mathematical form,

$$G = -\frac{1}{B}\frac{dU}{da} \tag{1}$$

where:

a = delamination length; and

U = total elastic strain energy in the specimen;

³ The last approved version of this historical standard is referenced on www.astm.org.



B = specimen width.

3.3 Symbols:

3.3.1 A-intercept of the linear fit of compliance versus crack length cubed data

3.3.2 a-delamination length

3.3.3 a_i —insert length in the trimmed specimen

3.3.4 a_i —the jth crack length used during compliance calibration (j = 1,2)

3.3.5 a_o —delamination length used in fracture test

3.3.6 a_{calc}—crack length calculated from an unloading curve after the NPC test

3.3.7 a_{PC} —actual crack length used during the PC test

3.3.8 a_{vis} —visually determined crack length after the NPC test

3.3.9 B-specimen width

3.3.10 C—specimen compliance

3.3.11 C_0 —specimen compliance during load-up of the fracture test (See Figure 6 in 13.1)

3.3.12 C_u —specimen compliance from unloading after the non-precracked test

3.3.13 δ —displacement of loading roller during testing perpendicular to the plane of the specimen (Fig. 1)

3.3.14 E_{If} -flexural modulus of the specimen

3.3.15 G-total strain energy release rate

3.3.16 G_{IIC}-mode II interlaminar fracture toughness

3.3.17 Go-candidate mode II interlaminar fracture toughness

3.3.18 %Go-peak percentage of Go achieved during compliance calibration

3.3.19 h—specimen half-thickness (Fig. 2) Len Standard

3.3.20 L—specimen half-span (Fig. 2)

3.3.21 L_c — distance from the center of the support roller at the cracked end of the specimen to the cracked end of the specimen (Fig. 2)

3.3.22 L_u —distance from the center of the support roller at the uncracked end of the specimen to the uncracked end of the specimen (Fig. 2)

3.3.23 m—slope of the linear fit of compliance versus crack length cubed data

3.3.24 *P*—force applied to center loading roller and perpendicular to the plane of the specimen (Fig. 1)

 $3.3.25 P_c$ —eritical force for mode II fracture

3.3.26 P_i —the compliance calibration force used at crack length a_i

3.3.27 P_{Max}—maximum value of force on the force-displacement curve

3.3.28 r_1 —radius of the loading roller (Fig. 2)

3.3.29 r_2 —radius of the support rollers (Fig. 2)

3.3.30 r^2 —correlation coefficient of linear fit of compliance versus crack length cubed

3.3.31 As-Maximum measured difference in crack length along the delamination front of the precrack

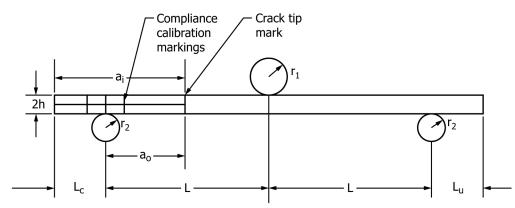


FIG. 2 ENF Specimen, Fixture, and Dimensions

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3.3.32 U-total elastic strain energy in the specimen

3.3 Symbols:

A-intercept of the linear fit of compliance versus crack length cubed data

a-delamination length

acalc-crack length calculated from an unloading curve after the NPC test

a,---insert length in the trimmed specimen

 a_i —the jth crack length used during compliance calibration (j = 1, 2)

a_o-delamination length used in fracture test

 a_{PC} —actual crack length used during the PC test

 a_{vis} —visually determined crack length after the NPC test

B-specimen width

C—specimen compliance

 C_0 —specimen compliance during load-up of the fracture test (see Figure 6 in 13.1)

 \underline{C}_{μ} —specimen compliance from unloading after the non-precracked test

 δ —displacement of loading roller during testing perpendicular to the plane of the specimen (Fig. 1)

 E_{1f} —flexural modulus of the specimen

G-total strain energy release rate

G_{IIC}—mode II interlaminar fracture toughness

Go-candidate mode II interlaminar fracture toughness

 $\frac{\widetilde{G}_Q}{\widetilde{G}_Q}$ peak percentage of G_Q achieved during compliance calibration

h-specimen half-thickness (Fig. 2)

L-specimen half-span (Fig. 2)

L_c-distance from the center of the support roller at the cracked end of the specimen to the cracked end of the specimen (Fig.

<u>2)</u>

 L_{u} —distance from the center of the support roller at the uncracked end of the specimen to the uncracked end of the specimen (Fig. 2)

m-slope of the linear fit of compliance versus crack length cubed data

P—force applied to center loading roller and perpendicular to the plane of the specimen (Fig. 1)

<u>*P_c*</u>—critical force for mode II fracture

 P_i —the compliance calibration force used at crack length a_i

 P_{Max} —maximum value of force on the force-displacement curve

 r_1 —radius of the loading roller (Fig. 2)

 r_2 —radius of the support rollers (Fig. 2)

 r^2 —correlation coefficient of linear fit of compliance versus crack length cubed

 Δs —maximum measured difference in crack length along the delamination front of the precrack /astm-d7905-d7905m-19 U—total elastic strain energy in the specimen

4. Summary of Test Method

4.1 The ENF specimen shown in Fig. 1 consists of a rectangular, uniform thickness, unidirectional laminated composite specimen containing a non-adhesive insert at the midplane that serves as a delamination initiator. Forces are applied to the specimen through an ENF fixture under displacement controlled loading.

4.2 Delamination growth is not stable in the ENF test. A method is presented so that the initiation values of the mode II interlaminar fracture toughness are obtained from the preimplanted insert as well as from a precrack.

4.3 A record of the applied force versus center roller displacement is to be obtained using an *x*-*y* recorder or equivalent real-time plotting device, or else it may be obtained and stored digitally. The mode II interlaminar fracture toughness, G_{Hc} , is obtained using the compliance calibration (CC) method. This is the only acceptable method of data reduction for this test (1).⁴

4.4 This standard recommends that static mode II precracking is performed and a recommended method is described. Other precracking methods may be used provided that a record of the shape of the precracked delamination front is obtained prior to the PC test. Precracking methods that typically leave crack front markings for post-test evaluation of these values include mode I and fatigue mode II.

5. Significance and Use

5.1 Susceptibility to delamination is one of the major design concerns for many advanced laminated composite structures. Knowledge of a laminated composite material's resistance to interlaminar fracture is useful for product development and material selection. Furthermore, a measurement of the mode II interlaminar fracture toughness that is independent of specimen geometry

⁴ The boldface numbers in parentheses refer to <u>athe</u> list of references at the end of this standard.

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or method of force introduction is useful for establishing design allowables used in damage tolerance analyses of composite structures. Knowledge of both the non-precracked and precracked toughnesses allows the appropriate value to be used for the application of interest.

5.2 This test method can serve the following purposes:

5.2.1 To establish quantitatively the effect of fiber surface treatment, local variations in fiber volume fraction, and processing and environmental variables on G_{IIc} of a particular composite material;

5.2.2 To compare quantitatively the relative values of G_{IIc} for composite materials with different constituents;

5.2.3 To compare quantitatively the values of G_{IIc} obtained from different batches of a specific composite material, for example, to use as a material screening criterion or to develop a design allowable; and

5.2.4 To develop delamination failure criteria for composite damage tolerance and durability analyses.

6. Interferences

6.1 Linear elastic behavior is assumed in the calculation of G used in this method. This assumption is valid when the zone of damage or nonlinear deformation at the delamination front, or both, is small relative to the smallest specimen dimension, which is typically the specimen's thickness for the ENF test.

6.2 G_{IIc} is obtained for both non-precracked and precracked specimens based on the maximum load point. G_{IIc} based on the nonlinear load point or other measures, such as a compliance offset, may also be obtained if desired. However, definitions of this type have not been related to any specific physical occurrences in the ENF test.

6.3 The three loading noses in the ENF test fixture may be fixed, rotatable, or rolling. Fixed loading noses or pins supported in a v-groove are recommended, and loading noses of this type were used in the interlaboratory test program that was conducted in support of this standard. The type of supports that are used is to be reported as described in Section 14. The loading noses should uniformly contact the specimen across its width. Lack of uniform contact can affect results, most commonly due to non-uniform loading across the width of the specimen. Formulas used in this standard assume a uniform line loading across the entire specimen width at the loading nose and at the specimen supports; deviations from this type of loading are beyond the scope of this standard.

6.4 There is an inherent error associated with the use of Eq 7 to obtain the calculated crack length, and it is not expected that the calculated crack length will exactly correspond to the true length of the precrack. However, since toughness is computed by CC, it has been shown (2) that this error in crack length will not affect the accuracy of the computed toughness provided that the recommended approach is followed.

6.6 A precracking method that only produces a short crack "jump," e.g., for example, by positioning a specimen with a crack tip close to the center loading roller, may produce precracked toughness values that are significantly higher than those that will be produced for a long crack jump following the recommended procedure (2, 3).

6.7 The toughness measured using this method is sensitive to reinforcement volume and void content. Consequently, the test results may reflect manufacturing quality as much as material properties.

6.8 Number of Points for CC—The use of a three-point CC was studied extensively in References (2, 4) and resulted in the recommended approach (subsection (11.9)). However, equivalent results will be obtained with a five-point CC, and one may use this approach following Note 4 (11.9).

6.9 The toughness values obtained by this test method for delamination growth at $0^{\circ}/0^{\circ}$ interfaces may not be representative of the toughness required for delamination growth at interfaces with different relative ply orientations.

7. Apparatus

7.1 *Testing Machine*—A properly calibrated test machine shall be used which can be operated in a displacement control mode with a constant displacement rate in the range from 0.025 to 1.6 mm/min [0.001 to 0.063 in./min]. The testing machine will conform to the requirements of Practices E4.

7.2 The testing machine shall be equipped with a loading fixture as shown in Fig. 1 and Fig. 2.



7.2.1 A fixture geometry with a nominal specimen span length (2L) of 100 mm [4 in.] and a nominal half-span length (L) of 50 mm [2 in.] is required.

7.2.2 The <u>cylindrical</u> loading rollersurface shall have a radius, r_1 , in the range of 4.7 to 9.6 mm [0.185 to 0.378 in.]. The support rollers 0.378 in.]. The cylindrical supporting surfaces shall have the same radius, r_2 , which shall be in the range of 3.0 to 6.4 mm [0.118 to 0.250 in.]. 0.250 in.]. The loading rollersurface shall be centered between the two support rollerssupporting surfaces (Fig. 2). All rollers shall haveload and support surfaces shall be finely ground surfacesand free of indentation and burrs with all sharp edges relieved, with a hardness of 60 to 62 HRC, as specified in Test Methods 55 HRC or greater. E18. Loading and support rollerssurfaces may be arranged in a fixed, rotatable, or rolling arrangement, where rotation may occur only about the roller cylindrical surfaces shall be restrained, and loading rollerssurface shall only be free to move vertically when viewed in the orientation of Fig. 1 and Fig. 2 (i.e., (that is, perpendicular to the plane of the specimen).

7.2.3 The system compliance, defined as the compliance of the load frame with the test fixture installed, shall be less than $\frac{3\%_3 \%}{9}$ of the measured compliance of the specimens that are tested. The system compliance shall be determined by using an essentially rigid calibration bar with the ENF test fixture and a span length (2L) of 100 mm [4.0 in.]. It is recommended that the calibration bar is at least as stiff as a steel bar with a moment of inertia, *I*, equal to 6 cm⁴ [0.144 in.⁴]. When this is the case, the system compliance can be determined as the slope of the deflection versus force data from the test of the calibration bar in the ENF fixture. For calibration bars with a lower moment of inertia, the bar's compliance should be accounted for. Here, the system compliance of the calibration bar, defined as $L^3/(6EI)$, where *L* is the half-span length and *E* and *I* are the Young's modulus and moment of inertia, respectively, of the calibration bar. The system compliance shall then be compared to the minimum compliance from all specimens tested to ensure that the $\frac{3\%_3 \%}{9}$ requirement is met. It is recommended that the system compliance tests be performed with a nominal loading rate of $\frac{0.050.05 \text{ mm}}{0.02030.003 \text{ in}}$. $\frac{-[0.002 \text{ in}/\text{min}]}{-(\text{min}]}$, but rates in the range of 0.02 to $\frac{0.080.08 \text{ mm}}{-\text{mm}/\text{min}/\text{min}}$ [0.0008 to $\frac{0.0030.003 \text{ in}}{-\text{mm}/\text{min}}/\text{min}]$ are acceptable.

7.2.4 The fixture cannot have rotational bearings that allow rotation about an axis parallel to the length direction of the specimen.

7.2.5 It is recommended that the test fixture be equipped with alignment features to ensure that (1) the loading and support rollers are parallel, and (2) the longitudinal direction of the specimen is perpendicular to the roller direction (3).

7.3 Force Indicator—The testing machine's force-sensing device shall be capable of indicating the total force carried by the test specimen. This device shall be essentially free from inertia-lag at the specified rate of testing and shall indicate the force with an accuracy over the force range(s) of interest of within $\pm 1\% \pm 1\%$ of the indicated value. Forces are dependent on the specimen geometry and toughness. A method to calculate the expected forces can be found in Annex A1.

7.4 Load Point Displacement Indicator—The load point displacement may be obtained from the crosshead separation of the load frame provided that the compliance requirement of subsection 7.2.3 is satisfied. Otherwise, the load point displacement shall be taken from a properly calibrated external gagegauge or transducer and/or a stiffer test fixture and/or fixture, or both, or load frame should be used. used, or both. The load point displacement indicator shall indicate the load point displacement with an accuracy of $\pm 1\% \pm 1\%$ at the displacement at which delamination growth occurs.

7.5 Force versus Load Point Displacement Record—A digital record of force versus load point displacement shall be stored for subsequent post-processing.

7.6 <u>Micrometers and Calipers</u>—The micrometer(s) or caliper(s) used to measure specimens prior to testing shall have A micrometer with a 4 to 7 mm [0.16 to 0.28 in.] nominal diameter ball interface or a flat anvil interface for the measurement of smooth surfaces or a suitably sized ball interface for the measurement of rough surfaces, such as the bag side of a laminate. shall be used to measure the specimen thickness. A ball interface is recommended for thickness measurements when at least one surface is irregular (for example, a course peel ply surface, which is neither smooth nor flat). A micrometer or caliper with a flat anvil interface shall be used for measuring length, width, and other machined surface dimensions. The use of alternative measurement devices is permitted if specified (or agreed to) by the test requestor and reported by the testing laboratory. The accuracy of the instrumentsinstrument(s) shall be suitable for reading to within $\pm 1\%-1\%$ of the sample width and thickness. specimen dimensions. For typical specimen geometries, an instrument with an accuracy of $\pm 2.5 \ \mum [0.0001 \ in.]$ is desirable $\pm 0.0025 \ mm [\pm 0.001 \ in.]$ is adequate for thickness measurements, while an instrument with an accuracy of $\pm 25 \ \mum [0.001 \ in.]$ is desirable for width measurements. $\pm 0.025 \ mm [\pm 0.001 \ in.]$ is adequate for measurement of length, width, and other machined surface for measurements.

8. Sampling and Test Specimens

8.1 *Sampling*—Test at least five specimens per test condition unless valid results can be gained through the use of fewer specimens, such as the case of a designed experiment. For statistically significant data, the procedures outlined in Practice E122 should be consulted. The method of sampling shall be reported.



8.2 Specimen and Test Configuration—Test laminates must contain an even number of plies and must be unidirectional, with delamination growth occurring in the 0° (zero degree) direction. Specimen dimensions shall conform to those presented in Fig. 3 and Fig. 4, which are chosen such that placement of the specimen within the fixture will be as defined in Table 1 and Fig. 2.

8.3 Manufacturing:

8.3.1 A flat composite plate shall be manufactured with a preimplanted non-adhesive film insert. Specimens are to be cut from these plates as shown in Fig. 3 and Fig. 4. Fabrication and machining are to be performed following Guide D5687/D5687M.

8.3.2 A non-adhesive film insert shall be implanted at the midplane of the laminate during layup to form an initiation site for the delamination (Fig. 3 and Fig. 4). The film thickness shall be no greater than 13 μ m [0.0005 in.]. A polymer film is recommended for the insert to avoid problems with folding or crimping at the cut end of the insert. For epoxy matrix composites cured at or below $177^{\circ}C$ [350°F], 177 °C [350 °F], a thin film made of polytetrafluoroethylene (PTFE) is recommended. For composites with polyimide, bismaleimide, or thermoplastic matrices that are manufactured at relatively high temperatures, i.e., that is, greater than $177^{\circ}C$ [350°F], 177 °C [350 °F], a thin polyimide film is recommended. If a polyimide film is used, the film shall be painted or sprayed with a mold release agent before it is inserted in the laminate. Caution should be used, as mold release agents containing silicone may contaminate the laminate by migration through the individual layers. It is often helpful to coat the film at least once and then bake the film before placing the film on the composite. This will help to prevent silicone migration within the composite. It also is often necessary to decohere the light bond that might form between the insert and the composite (2). For materials outside the scope of this standard, different film materials and procedures may be required.

8.3.3 The plate shall be made in such a way that the specimen dimensions presented in Fig. 3 and Fig. 4 may be achieved. Manufacturing large panels with a full-width insert in the center of the length direction is recommended to prevent thickness variations in the test specimens. After manufacture, these panels are cut width-wise along the centerline of the insert to create two plates, each with an edge view as shown in Fig. 3 and Fig. 4. A typical panel would be 400 mm [16 in.] long in the 0° direction with a 100 mm [4 in.] insert. Depending on the saw blade and amount trimmed at the edges, this will yield two plates that are approximately 200 mm [8 in.] long with an initial insert length (a_i) of approximately 50 mm [2 in.].

8.3.4 Prior to cutting the plate into specimens, the end of the insert should be accurately located and marked, and markings should be placed on the plate such that location of each specimen relative to the original plate geometry will be identifiable subsequent to cutting.

8.3.5 Individual specimens are to be cut such that they fall within the range of allowable lengths and widths specified in Fig. 3 and Fig. 4.

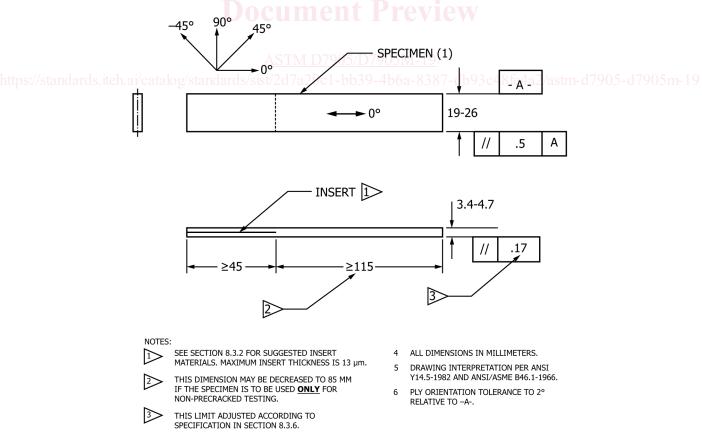
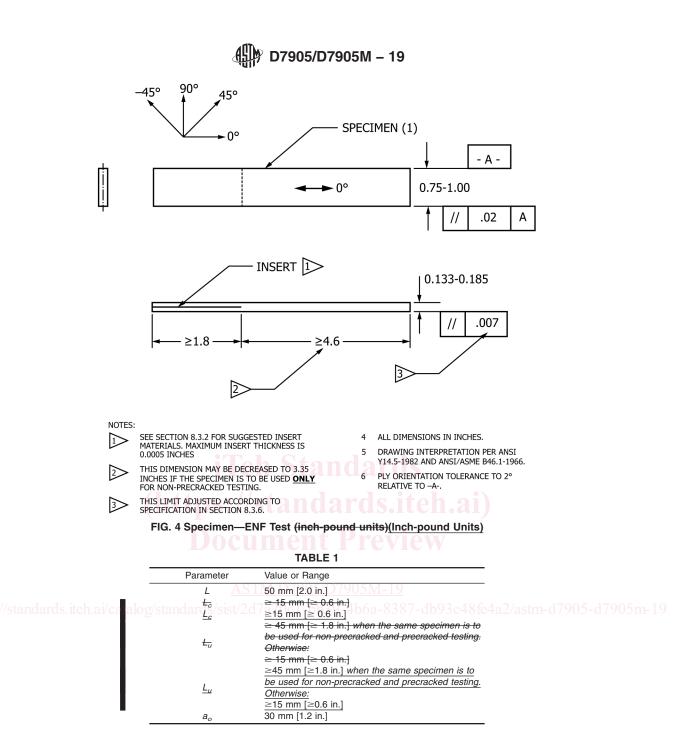


FIG. 3 Specimen—ENF Test (SI units)Units)



8.3.6 Subsequent to cutting, measure the width, *B*, at the three points of each specimen that will correspond to the contact locations of the three rollers when the specimen is tested in the non-precracked configuration. Measure the thickness, 2h, of each specimen at six points, with two thickness measurements at each of the points where the width was measured; one on the left side and one on the right side. The individual and average values of the three width measurements and the six thickness measurements shall be recorded. The variation in specimen width among all measurements shall not exceed 0.5 mm [0.02 in.]in.], and the variation in specimen thickness shall not exceed 5%5 % of the mean value.

8.4 *Labeling*—Label the specimens so that they will be distinct from each other and traceable back to the raw material, and in a manner that will both be unaffected by the test and not influence the test.

8.5 *Void Content*—It is recommended that void content and fiber volume be reported. Void content may be determined using Test <u>MethodMethods</u> D2734 and fiber volume fraction may be determined using Test <u>MethodMethods</u> D3171.

9. Calibration

9.1 The accuracy of all measuring equipment shall have certified calibrations that are current at the time of use of the equipment.

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10. Conditioning

10.1 The recommended pre-test condition is effective moisture equilibrium at a specific relative humidity as established by Test Method D5229/D5229M₅; however, if the test requestor does not explicitly specify a pre-test conditioning environment, no conditioning is required and the test specimens may be tested as prepared.

10.2 The pre-test specimen conditioning process, to include specified environmental exposure levels and resulting moisture content, shall be reported with the test data.

NOTE 2—The term "moisture," as used in Test Method D5229/D5229M, includes not only the vapor of a liquid and its condensate, but the liquid itself in large quantities, as for immersion.

10.3 If no explicit conditioning process is performed the specimen conditioning process shall be reported as "unconditioned" and the moisture content as "unknown."

11. Procedure

11.1 Parameters to be Specified Prior to Test:

11.1.1 The specimen sampling method, specimen geometry, and conditioning travelers (if required); required);

11.1.2 The properties and data reporting format desired; desired;

11.1.3 The environmental conditioning test parameters; parameters; and

11.1.4 If performed, the sampling method, specimen geometry, and test parameters used to determine density and constituent volumes.

11.2 Condition the specimens as required. Store the specimens in the conditioned environment until test time, if the test environment is different than the conditioning environment.

11.3 Specimen Preparation:

11.3.1 Measure and record the width and thickness of each specimen as specified in subsection 8.3.6.

11.3.2 A light coating of white or silver spray paint, or equivalent, shall be applied to the specimen edges. This is to assist in the visual detection of the delamination tip and in making compliance calibration (CC) markings (Fig. 2). Once the paint is dry, the tip of the insert shall be marked with a thin vertical pencil line. line made with a mechanical pencil containing a 0.5 mm [0.02] in.] diameter lead or smaller. The edges shall then be marked with three vertical compliance calibration markings, within the cracked region, at distances of 20, 30, and $40 \text{ mm} - 40 \text{ mm} [0.8, 1.2, and <math>1.6 \text{ in.} - 1.6 \text{ i$

11.3.3 If specific gravity, density, reinforcement volume, or void volume, or combinations thereof, are to be reported, then obtain these samples from the same panels being tested. Specific gravity and density may be evaluated by means of Test MethodMethods D792. Volume percent of the constituents may be evaluated by one of the matrix digestion procedures of Test MethodMethods D3171, or, for certain reinforcement materials such as glass and ceramics, by the matrix burn-off technique of Test Method D2584. The void content equations of Test Method D2734 are applicable to both Test Method D2584 and the matrix digestion procedures.

11.4 *Test Environment*—If possible, test the specimen under the same fluid exposure level used for conditioning. However, cases such as elevated temperature testing of a moist specimen place unrealistic requirements on the capabilities of common testing machine environmental chambers. In such cases, the mechanical test environment may need to be modified, for example, by testing at elevated temperature with no fluid exposure control, but with a specified limit on time to failure from withdrawal from the conditioning chamber. Record any modifications to the test environment.

NOTE 3—When testing a conditioned specimen at elevated temperature with no fluid exposure control, the percentage moisture loss of the specimen prior to test completion may be estimated by placing a conditioned traveler of known weight within the test chamber at the same time the specimen is placed in the chamber. The traveler should be configured to mimic the specimen, such that moisture evaporation is comparable to that of the test specimen. Upon completion of the test, the traveler is removed from the chamber, weighed, and the percentage weight calculated and reported.

11.5 The specimen shall be placed in the fixture so that its longitudinal direction is perpendicular to the loading rollers (see subsection 7.2.5).

11.6 Loading for all CC and fracture tests shall be performed in displacement control at a nominal rate of θ .50.5 mm mm/min/min [0.02 in. [0.02 in./min],/min], although rates between 0.10 and θ .800.80 mm mm/min/min [0.004 to θ .0310.031 in. in./min]/min] are acceptable. Unless otherwise specified, unloading shall also be in displacement control at a rate between 0.10 and 1.61.6 mm mm/min/min [0.004 to θ .063-0.063 in. in./min]/min].

11.7 Peak forces during CC are equal to $\frac{50\%50\%}{25\%25\%}$ of the expected value of the critical force (P_c) at that particular crack length; these are chosen to correspond to approximately $\frac{25\%25\%}{25\%}$ of G_{IIc} . That is, the peak CC force varies with crack length. The method of determining the peak force for each crack length during CC is presented in Annex A2.

11.8 Compliance calibration tests are performed by loading the specimen to the peak CC force as defined in subsection 11.7 and then unloading. The force and deflection data are to be recorded continuously or at frequent and regular intervals during the loading portion only; a sampling rate of 5 Hz or greater and a target minimum of 500 data points per test are recommended.

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11.9 Values of G_{IIc} shall be obtained from the insert and from the precrack. All toughness values are obtained using the CC method. Subsection 11.9.1 is to be used if both NPC and PC toughness values are to be obtained from the same specimen and the recommended precracking procedure is to be adopted; otherwise, subsections-11.9.2 and 11.9.3 are to be used. Peak forces expected during these fracture tests may be estimated by the method in Annex A1.

NOTE 4—If desired, a five-point CC may be used. In this case, CC is performed for both the NPC and PC tests with crack lengths, *a*, equal to 20, 25, 35, and 40 mm [0.8, 1.0, 1.4, and 1.6 in.),in.], and the fracture test is still performed with $a = 30 \text{ mm} (\frac{1.2 \text{ in.}}{1.2 \text{ in.}})$. Data from all five crack lengths are then used to obtain the CC coefficients (subsection (13.2).

11.9.1 Non-Precracked and Precracked Toughness from the Same Specimen—In the approach that follows, crack advance during the NPC test creates the precrack that is used for the PC test. The approach has been shown to produce accurate NPC and PC toughnesses with a PC G_{IIc} that is within or approaching the "minimum toughness plateau" that some materials evidence, i.e., that is, when G_{IIc} decreases to a minimum plateau value with the amount of dynamic advance that occurs during precracking (2-4). The approach also ensures that any differences between the location of the true and calculated crack tip do not affect the accuracy of G_{IIc} (2).

11.9.1.1 *Non-Precracked CC*—With reference to Fig. 2, the specimen is placed in the fixture so that the CC mark that is farthest from the cracked end is aligned with the center of the support roller at the cracked end. The first CC test is then performed with a crack length, *a*, equal to 20 mm [0.8 in.], following the procedure defined in subsection-11.8. The specimen is then repositioned such that a = 40 mm [1.6 in.], i.e., that is, so that the CC mark that is closest to the cracked end is aligned with the center of the support roller at the cracked end. The second CC test is then performed as defined in subsection-11.8.

11.9.1.2 Non-Precracked Fracture Test—Following NPC CC, the specimen shall be repositioned in the fixture so that a = 30 mm. This shall correspond to placing the center CC mark over the center of the support roller that is at the cracked end. The specimen is then loaded until the delamination advances, as seen by visual assessment on the specimen or by a drop in force on the force versus displacement plot. The specimen shall be unloaded at a nominal rate of 0.5 mm/min [0.02 in./min], although rates between 0.10 and 0.80 mm/min [0.004 to 0.031 in./min] are acceptable (i.e., (that is, in order to decrease the total time required for the test). The force and displacement data are to be recorded continuously or at frequent and regular intervals during the entire test; a sampling rate of 5 Hz or greater and a target minimum of 1000 data points per test are recommended.

11.9.1.3 Determination of Crack Length for the Precracked Test—The unloading data from the non-precracked fracture test of subsection-11.9.1.2 is used to compute a value of a_{calc} using the method of subsection-13.6. This value of a_{calc} is measured from the existing center CC mark. A new "PC crack tip mark" shall be placed at this location. Three new "PC CC markings" shall then be placed at 20, 30, and 40 mm [0.8, 1.2, and 1.6 in.] from the PC crack tip mark as shown in Fig. 5. The center mark, with distance from the crack tip equal to 30 mm [1.2 in.] is for the fracture test, and the other two marks are for CC testing.

If desired, the location of the crack tip may also be determined visually as the average of the locations found on the two edges. If the visually determined crack tip, a_{vis} , is (1) past (to the right of) the loading roller and (2) longer than a_{calc} , then a_{vis} may be used in place of a_{calc} for the placement of the PC crack tip mark.

11.9.1.4 *Precracked CC*—Prior to the PC fracture test, the compliances from two different crack lengths are obtained by appropriate placement of the specimen in the fixture. The first CC test is performed with a = 20 mm [0.8 in.] and the second with a = 40 mm [1.6 in.]. This is performed using the PC CC markings of subsection 11.9.1.3 and following the procedure of subsection 11.9.1.1.

11.9.1.5 *Precracked Fracture Test*—Following PC CC, the specimen shall be repositioned in the fixture so that a = 30 mm [1.2 in.]. This shall correspond to placing the center PC CC mark over the center of the support roller that is at the cracked end. The specimen is then loaded until the delamination advances, as seen by visual assessment on the specimen or by a drop in force on the force versus displacement plot. The specimen is then unloaded. The force and deflection data are to be recorded continuously or at frequent and regular intervals during the loading portion only; a sampling rate of 5 Hz or greater and a target minimum of 750 data points per test are recommended.

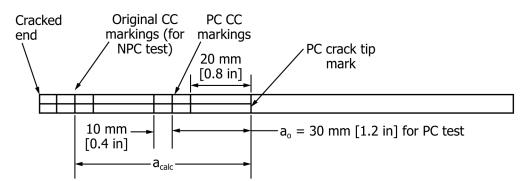


FIG. 5 Configuration of Specimen for Precracked Test When the Same Specimen is Used for NPC and PC Testing (nominal dimensions shown)(Nominal Dimensions Shown)