



# Standard Test Method for Photoelastic Measurements of Birefringence and Residual Strains in Transparent or Translucent Plastic Materials<sup>1</sup>

This standard is issued under the fixed designation D4093; 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.

## INTRODUCTION

Light propagates in transparent materials at a speed,  $v$ , that is lower than its speed in vacuum,  $c$ . In isotropic unstrained materials the index of refraction,  $n = c/v$ , is independent of the orientation of the plane of vibration of light. Transparent materials, when strained, become optically anisotropic and the index of refraction becomes directional. The change in index of refraction is related to strains. If  $n_o$  is the refractive index of unstrained material, the three principal indices of refraction,  $n_i$ , become linear functions of strain:

$$n_i - n_o = \sum A_{ij} \epsilon_j$$

Using photoelastic techniques (initially developed to measure stresses in transparent models) strains in plastics can be assessed. In isotropic materials, two material constants,  $A$  and  $B$ , are sufficient to describe their optomechanical behavior:

$$\begin{aligned} A_{ij} &= A \text{ when } i = j, \text{ and} \\ A_{ij} &= B \text{ when } i \neq j. \end{aligned}$$

When light propagates through a region (where principal strains  $\epsilon_1$  and  $\epsilon_2$  are contained in the plane perpendicular to the direction of light propagation (see Fig. 1), the incoming vibration splits into two waves vibrating in planes of  $\epsilon_1$  and  $\epsilon_2$ . The difference between the indexes of refraction  $n_1 = c/v_1$  and  $n_2 = c/v_2$  (or birefringence) is:

$$n_1 - n_2 = (A - B)(\epsilon_1 - \epsilon_2) = k(\epsilon_1 - \epsilon_2)$$

where  $k$  is a material property called the strain-optical constant. As a result of their velocity difference, the waves vibrating along the two principal planes will emerge out of phase, their relative distance, or retardation,  $\delta$ , given by:

$$\delta = (n_1 - n_2)t = kt(\epsilon_1 - \epsilon_2)$$

where  $t$  is the thickness of material crossed by the light. A similar equation, relating  $\delta$  to the difference of principal stresses,  $\sigma_1$  and  $\sigma_2$ , can be written:

$$\delta = (n_1 - n_2)t = Ct(\sigma_1 - \sigma_2)$$

The objective of photoelastic investigation is to measure: (a) the azimuth, or direction of principal strains,  $\epsilon_1$  and  $\epsilon_2$  (or stresses  $\sigma_1$  and  $\sigma_2$ ), and (b) the retardation,  $\delta$ , used to determine the magnitude of strains. A complete theory of photoelastic effect can be found in the abundant literature on the subject (an extensive bibliography is provided in Appendix X2).

## 1. Scope

1.1 This test method covers measurements of direction of principal strains,  $\epsilon_1$  and  $\epsilon_2$ , and the photoelastic retardation,  $\delta$ , using a compensator, for the purpose of analyzing strains in transparent or translucent plastic materials. This test method

can be used to measure birefringence and to determine the difference of principal strains or normal strains when the principal directions do not change substantially within the light path.

1.2 In addition to the method using a compensator described in this test method, other methods are in use, such as the goniometric method (using rotation of the analyzer) mostly applied for measuring small retardation, and expressing it as a fraction of a wavelength. Nonvisual methods employing spectrophotometric measurements and eliminating the human judgment factor are also possible.

<sup>1</sup> This test method is under the jurisdiction of ASTM Committee D20 on Plastics and is the direct responsibility of Subcommittee D20.10 on Mechanical Properties.

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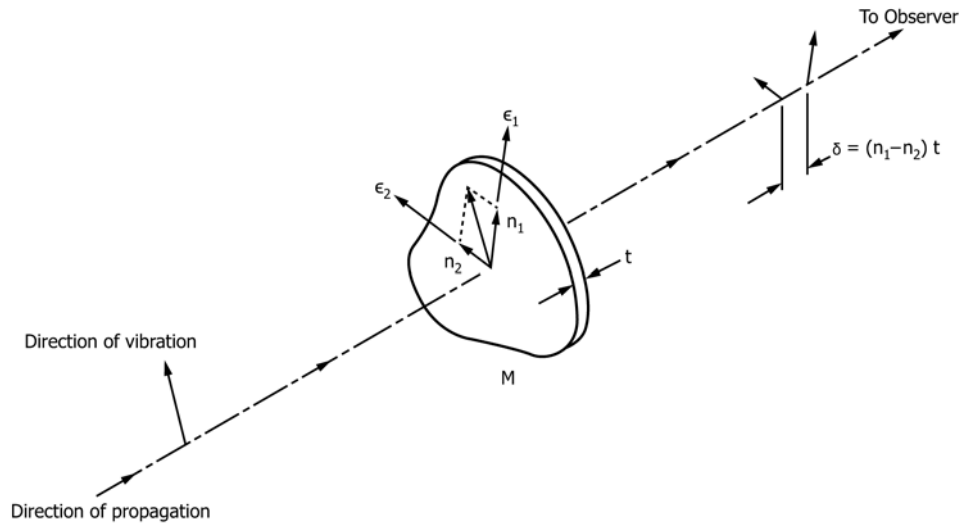


FIG. 1 Propagation of Light in a Strained Transparent Material

1.3 Test data obtained by this test method is relevant and appropriate for use in engineering design.

1.4 The values stated in either SI units or inch-pound units are to be regarded as standard. The values stated in each system may not be exact equivalents; therefore, each system shall be used independently of the other. Combining values from the two systems may result in nonconformance with the standard.

1.5 *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 and health practices and determine the applicability of regulatory limitations prior to use.*

NOTE 1—There is no known ISO equivalent to this test method.

2. Referenced Documents

- 2.1 *ASTM Standards:*<sup>2</sup>
  - D618 Practice for Conditioning Plastics for Testing
  - D638 Test Method for Tensile Properties of Plastics
  - D882 Test Method for Tensile Properties of Thin Plastic Sheeting
  - D4000 Classification System for Specifying Plastic Materials
  - E691 Practice for Conducting an Interlaboratory Study to Determine the Precision of a Test Method

3. Terminology

3.1 *Definitions:*

3.1.1 *compensator*—an optical device used to measure retardation in transparent birefringent materials.

3.1.2 *polarizer*—polarizing element transmitting light vibrating in one plane only.

<sup>2</sup> 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.

3.1.3 *quarter-wave plate*—a transparent filter providing a relative retardation of ¼ wavelength throughout the transmitting area.

3.2 *Definitions of Terms Specific to This Standard:*

3.2.1 *birefringence*—retardation per unit thickness,  $\delta/t$ .

3.2.2 *retardation,  $\delta$* —distance (nm) between two wave fronts resulting from passage of light through a birefringent material. (Also called “relative retardations.”)

3.2.3 *strain,  $\epsilon$ -strain (or deformation per unit length)*—could be permanent, plastic strain introduced in manufacturing process, or elastic strain related to the existing state of stress. Both types of strains will produce strain-birefringence in most polymers. Birefringence can also result from optical anisotropy due to crystalline orientation.

3.2.4 *strain-optical constant,  $k$* —material property, relating the strains to changes of index of refraction (dimensionless).

$$k = (n_1 - n_2)/(\epsilon_1 - \epsilon_2)$$

3.2.5 *stress-optical constant,  $C$* —material property relating the stresses to change in index of refraction.  $C$  is expressed in  $m^2/N$  or Brewsters ( $10^{-12} m^2/N$ ).  $C$  is usually temperature-dependent.

$$C = (n_1 - n_2)/(\sigma_1 - \sigma_2)$$

4. Summary of Test Method

4.1 To analyze strains photoelastically, two quantities are measured: (a) the directions of principal strains and (b) the retardation,  $\delta$ , using light paths crossing the investigated material in normal or angular incidence.

4.2 The investigated specimen or sample is introduced between the polarizers (see Fig. 2 and Fig. 3). A synchronous rotation of polarizers follows until light intensity becomes zero at the observed location. The axes of the polarizers are then parallel to direction of strains, revealing these directions.

4.3 To suppress the directional sensitivity of the apparatus, the setup is changed, introducing additional filters. A calibrated

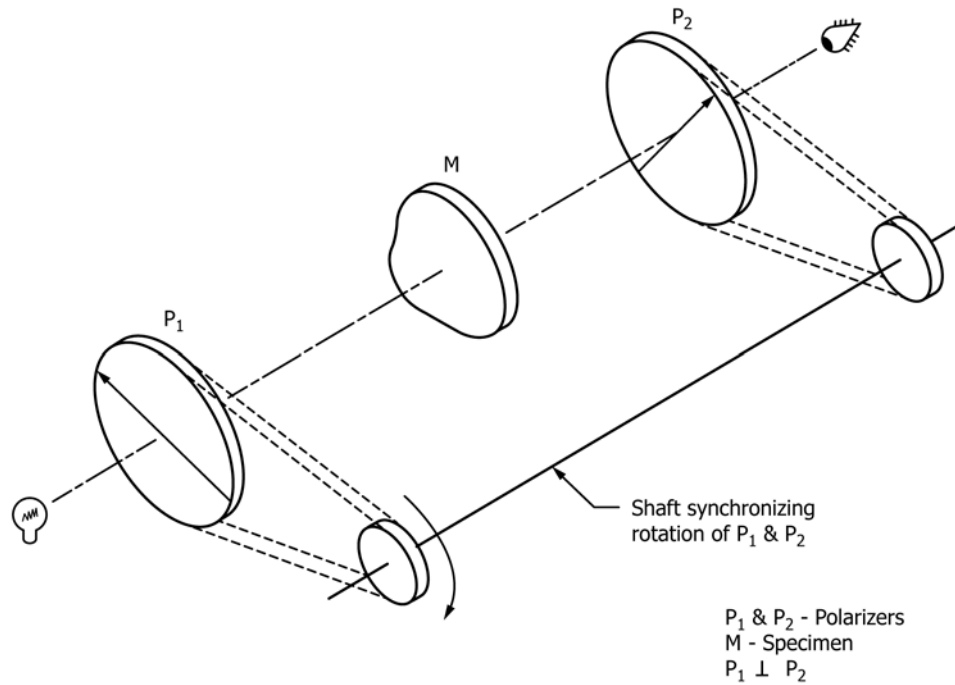


FIG. 2 Transmission Set-up of Polariscope

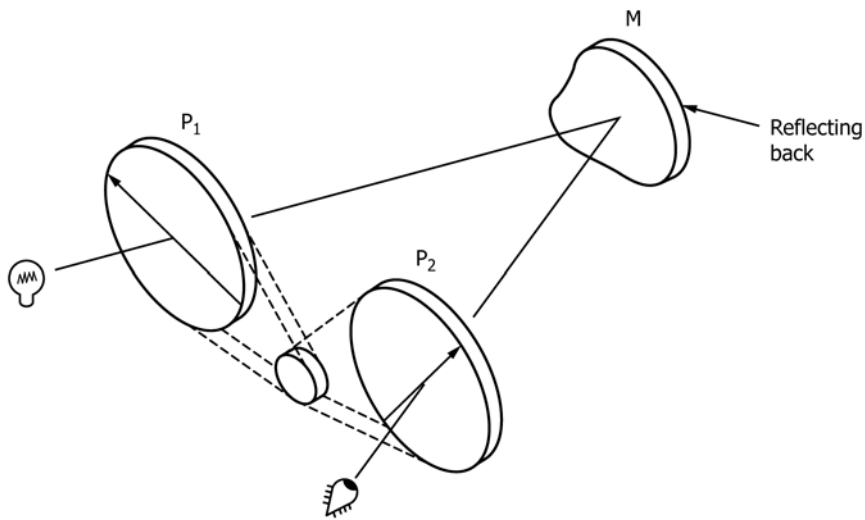


FIG. 3 Reflection Set-up of Polariscope

compensator is introduced and its setting adjusted until light intensity becomes zero at the observed location. The retardation in the calibrated compensator is then equal and opposite in sign to the retardation in the investigated specimen (see Fig. 4).

## 5. Significance and Use

5.1 The observation and measurement of strains in transparent or translucent materials is extensively used in various modeling techniques of experimental stress analysis.

5.2 Internal strains induced in manufacturing processes such as casting, molding, welding, extrusion, and polymer stretching can be assessed and part exhibiting excessive strains identified. Such measurements can lead to elimination of defective parts, process improvement, control of annealing operation, etc.

5.3 When testing for physical properties, polariscopic examination of specimens is required, to eliminate those specimens exhibiting abnormal internal strain level (or defects). For

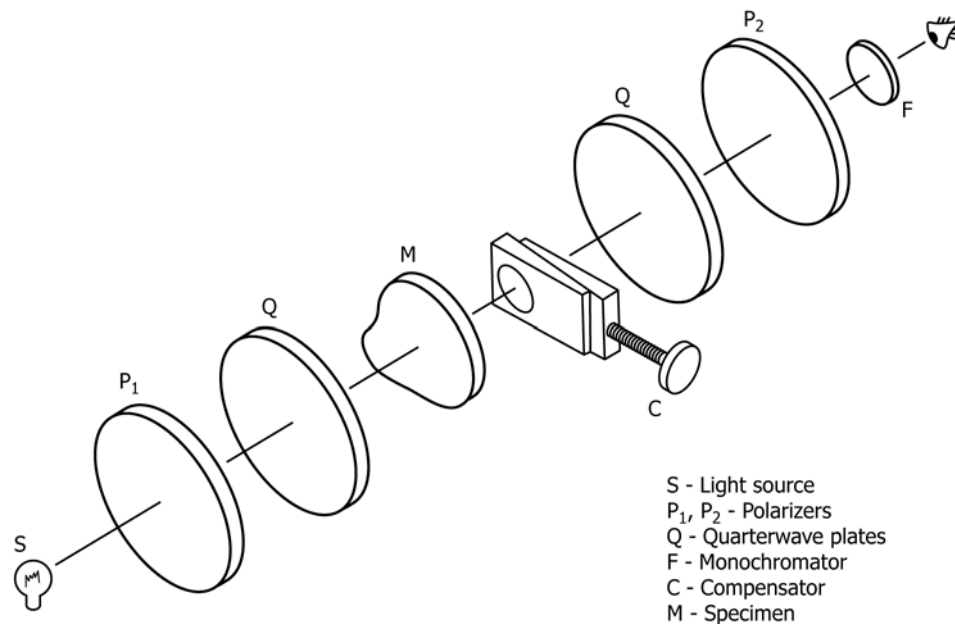


FIG. 4 Apparatus

example: Test Methods **D638** (Note 8) and **D882** (Note 11) recommend a polariscopic examination.

5.4 The birefringence of oriented polymers can be related to orientation, shrinkage, etc. The measurements of birefringence aid in characterization of these polymers.

5.5 For many materials, there may be a specification that requires the use of this test method, but with some procedural modifications that take precedence when adhering to the specification. Therefore, it is advisable to refer to that material specification before using this test method. Table 1 of Classification System **D4000** lists the ASTM materials standards that currently exist.

## 6. Apparatus

6.1 The apparatus used to measure strains is shown schematically in Fig. 4. It consists of the following items:

### 6.1.1 Light Source:

6.1.1.1 *Transmitted-Light Set-Up*—An incandescent lamp or properly spaced fluorescent tubes covered with a diffuser should provide a uniformly diffused light. To ensure adequate brightness, minimum illumination required is 0.3 W/in.<sup>2</sup> (0.0465 W/cm<sup>2</sup>). Maximum light source power is limited to ensure that the specimen temperature will not change more than 2°C during the test. The incandescent lamp must be selected to provide a color temperature no lower than 3150 K. There should be no visible nonuniformity, dark or bright spots on the diffuser surface, when no specimen is inserted in the apparatus.

6.1.1.2 *Reflection-Light Source*—For the reflection set-up an incandescent, reflector-equipped projection lamp is required. The lamp shall be equipped with proper lenses to ensure uniform illumination of the investigated object. At a distance of 2 ft (610 mm) from the lamp an area of 1 ft<sup>2</sup> (0.093 m<sup>2</sup>) should be illuminated, with no visible dark or bright spots. The lamp power should be at least 150 W.

6.1.2 *Polarizer*—The polarizing element shall be kept clean. The ratio of the transmittance of polarizers with their axes parallel, to the transmittance of the polarizers with their axes perpendicular to each other (or in crossed position), should not be less than 500. A glass-laminated construction of polarizers is recommended. The polarizers must be mechanically or electrically coupled to insure their mutually perpendicular setting while rotated together to measure directions. A graduated scale must be incorporated to indicate the common rotation of polarizers to a fixed reference mark.

6.1.3 *Quarter-Wave Plates*—Two quarter-wave plates are required in the procedure described below (see 9.2):

6.1.3.1 The retardation of each quarter-wave plate shall be  $142 \pm 15$  nm, uniform throughout its transmission area. The difference in retardation between the two quarter-wave plates should not exceed  $\pm 5$  nm.

6.1.3.2 The quarter-wave plates will be indexed, to permit their insertion in the field of the apparatus with their axes at 45° to the polarizers direction. The two quarter-wave plates shall have their axes crossed (that is, their optical axes perpendicular to each other), thus insuring that the field remains at maximum darkness when both quarter-wave plates are inserted (see Fig. 5).

6.1.4 *Compensator*—The compensator is the essential means of measuring retardation. The following types of compensators can be used:

6.1.4.1 *Linear Compensator*<sup>3</sup>—In the linear compensator the retardation in the compensator is linearly variable along its length. A graduated scale shall be attached to the compensator body in such a manner that slippage cannot occur. The calibration characteristic of the compensator shall include the position along its length (as indicated by the scale) of the line where the retardation is zero and the number of divisions  $d$  per

<sup>3</sup> Also known as “Babinet” compensator.

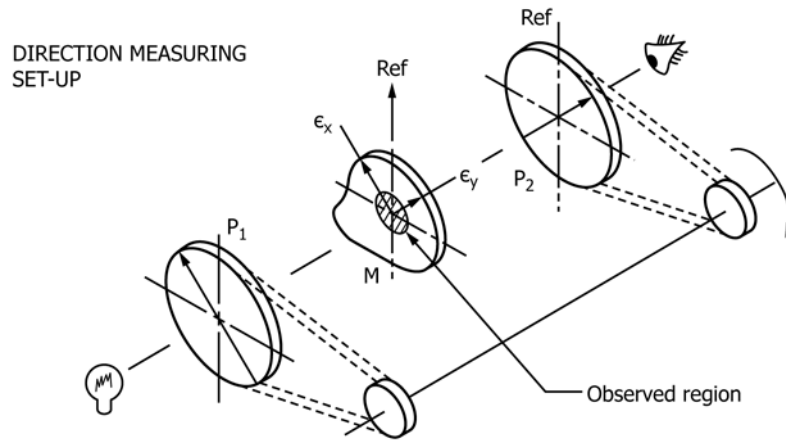


FIG. 5 Direction Measuring Set-up

unit retardation (usually one wavelength). (The retardation per division is  $D = \lambda/d$ .) The scale density shall be sufficient to provide clear visibility for observing 1 % of the useful range of the compensator.

6.1.4.2 *Uniform Field Compensator*<sup>4</sup>—The uniform field compensator is usually constructed from two optical wedges moved by means of a lead screw, the amount of relative motion being linearly related to the total thickness and the retardation. The lead screw motion shall be controlled by a dial drum or counter. Calibration of this compensator shall include the position, as indicated by the drum or counter, where the retardation is zero and the number of division of drum or counter  $d$  per unit of retardation. (The retardation per division is  $D = \lambda/d$ .)

6.1.4.3 Compensators have a limited range of measured retardation. In case the retardation in the sample exceeds the range of the compensator used, insertion of an offset retarder is needed. The offset retarder must be calibrated and positioned along the axes of the compensator, between the analyzer and the sample.

6.1.5 *Filter*—Monochromatic light is required to perform various operations in photoelasticity and some operations cannot be successfully accomplished using white light. In those instances a monochromatic light can be obtained introducing within the light path, a filter transmitting only light of the desired wave length. To best correlate with observation in white light, a narrow band-pass filter with peak transmittance at  $570 \pm 6$  nm and a maximum transmitted band-width (at half-peak point) of 10 nm should be used.

## 7. Test Specimen

7.1 Sheet, film, or more generally, a constant-thickness item can be examined using a transmission set-up. For use in reflection, a reflecting surface must be provided. This can be accomplished by painting one side of the specimen with aluminum paint.<sup>5</sup> Alternatively, it is possible to place the examined sheet specimen against a clean metal surface (preferably aluminum) or an aluminum-painted surface.

<sup>4</sup> Also known as “Babinet-Soleil” compensator.

<sup>5</sup> Krylon aluminum aerosol can spray paint was found satisfactory.

7.2 Examination of complex surfaces or shapes sometimes requires the use of an immersion liquid. The examined item is placed inside a tank containing a liquid selected to exhibit approximately the same index of refraction as the tested item. This technique is commonly used to examine three-dimensional shapes.

7.3 If conditioning is required, Procedure A of Practice D618 shall be used.

## 8. Calibration and Standardization

8.1 A periodic verification (every 6 months) is required to ensure that the apparatus is properly calibrated. The following points require verification:

### 8.1.1 Verification of Polariscopes:

8.1.1.1 Verify that the polarizers remain in “crossed” position. A small deviation of one of the polarizers produces an increase in the light intensity transmitted.

8.1.1.2 Verify that the quarter-wave plates are properly crossed. A small deviation of one quarter-wave plate from its “indexed” position will produce an increase in the light intensity transmitted.

### 8.1.2 Verification of the Compensator:

8.1.2.1 Examine the compensator in the polariscopes and verify that its  $\delta = 0$  point coincides with the calibration reported.

8.1.2.2 Using monochromatic light (filter), verify that the spacing of interference fringes,  $D$ , coincides with the calibration report. If  $\lambda$  is the wavelength of monochromatic light used, it should be verified that  $d = \lambda/D$ .

## 9. Procedure

### 9.1 Measuring Direction of Principal Strains:

9.1.1 Insert the specimen between the polarizers and align a characteristic reference direction of the specimen (for example: edge, axis of symmetry, base) with the reference of the instrument.

9.1.2 Set the polariscopes in the direction measuring set-up. The quarter-wave plates must be removed or their axes aligned with the polarizers (see Fig. 5).

9.1.3 Observe the light intensity at the point ( $s$ ) (or the region) where measuring is to be performed. Rotate polarizers

RETARDATION MEASURING  
SET - UP

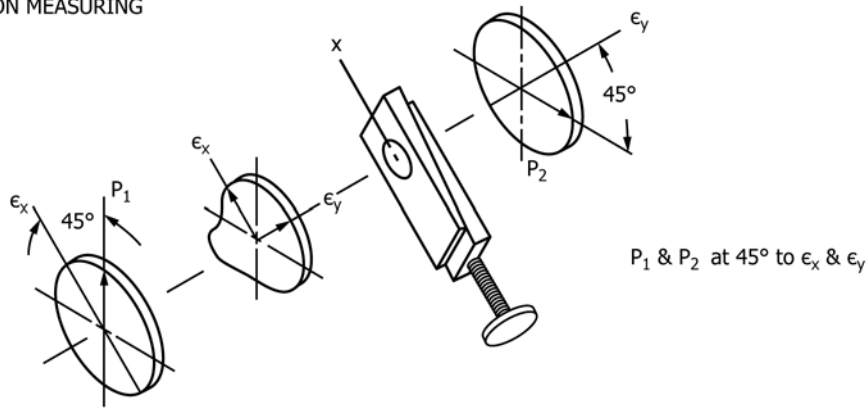


FIG. 6 Retardation Measuring Set-up

(synchronized together) until a minimum of light intensity emerges and the point (*s*) (or the region) appear dark or black.

9.1.4 Read on the dial the angle indicating the directions of the polarizer axes which are also the direction of principal strains at the point with respect to the reference direction.

9.1.5 In polarizing microscopes and all other instruments containing fixed polarizer and analyzer, a rotating stage shall be provided to support the sample and to measure the angle between the polarizer and sample reference direction. The polarizer in this setup must be aligned with the reference of the stage scale. Rotate the stage until a minimum of light intensity is observed and the area or point that is observed is dark or black. Read on stage scale the angle indicating the rotation of the stage to the reference, which is also the direction angle of the strain to the same reference.

NOTE 2—If the field of view appears dark and remains dark as the polarizers are rotated, the specimen is strain-free. If continuous rotation cannot produce total extinction (black), small changes of strain direction within the thickness of the observed region could be present. If no minimum light can be detected, the variations of strain directions are significant and the method described here is not applicable.

9.2 *Measuring Retardation Using a Linear (Babinet-Type) Compensator (Procedure A)*—The measurement of the retardation,  $\delta$ , can be performed after the direction of strains has been established. Two set-ups are possible:

(a) Place polarizers at 45° to the direction of principal strains measured in 9.1 (Fig. 6), or

(b) Insert two quarter-wave plate filters at 45° to the polarizers, with their axes crossed, as shown on Fig. 7. This set-up facilitates the observation of the specimen and selection of points for measurement when directions of strain vary significantly from point to point on the specimen.

9.2.1 After completing the set-up (a) or (b), observe and identify the point of measurement. The color versus retardation table (Appendix X1) provides a simple means to select the point of measurement properly. Uniform color observed over a broad region indicates a uniform strain area. Closely spaced color bands (isochromatics) indicate that strain gradient are substantial and the points must be selected carefully to provide meaningful data.

9.2.2 Introduce the compensator in the field of view, with the axes, *xy*, of the compensator closely aligned with the

direction of principal strains,  $\epsilon_x$  and  $\epsilon_y$ , in the specimen. The retardation produced by the specimen and the compensator are additive, producing the shift of color fringes in the compensator. Two mutually perpendicular positions of the compensator are possible; select the position which produces an upscale (toward larger number) shift of the black fringe.

9.2.3 Determine the shift of the black fringe (*d* division). If the compensator calibration constant is *D* nm per division, the measured retardation is:

$$\delta = Dd \text{ nm}$$

9.3 *Measuring the Retardation Using a “Uniform Field” (Babinet-Soleil)-Type Compensator (Procedure B)*:

9.3.1 Set up the polariscope (as indicated in 9.2).

9.3.2 Introduce the compensator, with its axes aligned with the direction,  $\epsilon_x$  and  $\epsilon_y$ , of measured strains. Observe the light transmitted by the specimen and the compensator and adjust the retardation of the compensator (advancing its lead-screw) until the total retardation observed is zero and a black fringe or area covers the observed point or region.

9.3.2.1 Two positions of the compensator are possible. Compensation can be accomplished in one of these positions. If the calibration of the compensator is *D* nm per division and *d* is the observed lead screw advance (drum or counter reading), the measured retardation is:

$$\delta = Dd \text{ nm}$$

The compensator also indicates which of the two directions, *x* or *y*, coincides with the larger index of refraction ( $n_x > n_y$ , or  $n_y > n_x$ ).

9.4 At every point where measurement of stress is performed, in addition to measuring the retardation, measure the thickness, *t*, using a suitable micrometer.

9.5 In some instances not only the differences of principal strains (shear strains) but also individual (normal) strain values are measured. In addition to the “normal-incidence” measurements of retardation described in 9.2 and 9.3 (rays of light approximately perpendicular to the specimen plane) “oblique-incidence” measurements are then required, with rays oriented at an angle to the normal. To perform these measurements proceed as follows:

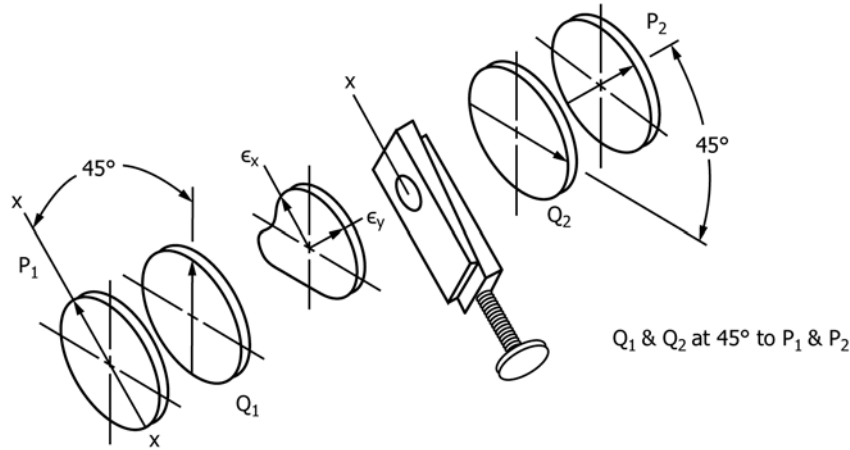


FIG. 7 Retardation Measuring Set-up

9.5.1 After completing the measurements of direction and retardation in normal incidence, place the specimen in the polariscope, using the tilting stage or prism arrangement shown in Fig. 8. Tilt the specimen to produce an angle  $\theta_x$  between light rays and the normal to the specimen. In both cases, the rotation  $\theta_x$  must be accomplished about one of the principal directions of strains  $x$  as measured in 9.1.

9.5.2 Measure retardation,  $\delta_{\theta_x}$ , with a compensator, using the same procedure as described in 9.2 or 9.3.

9.5.3 Establish the angle  $\theta_x$ .

9.5.3.1 If the specimen is immersed in an index-matching liquid, the angle  $\theta_x$  is the same as the tilt angle  $i$  of the specimen (Fig. 8).

9.5.3.2 If the specimen is not immersed, the angle  $\theta_x$  must be computed or established by calibration. The computed value is:

$$\sin \theta_x = \frac{\sin i}{n_o}$$

where  $i$  is the tilt angle and  $n_o$  is the index of refraction of the specimen. The effective angle  $\theta_x$  can also be established by calibration, as shown in Appendix X3.

## 10. Calculation of Birefringence and Strains

10.1 After measuring the direction of strains and retardation, the birefringence, strains, or stresses are calculated using the following relations and formulae:

10.1.1 *Birefringence* (retardation per unit thickness) in the plane  $xy$  of the specimen is as follows:

$$n_x - n_y = \delta/t$$

In the plane perpendicular to the specimen plane:

$$n_z - n_y = \frac{(\delta - \delta_{\theta_x} \cos \theta_x)}{t \sin^2 \theta_x}$$

where:

$\delta$  and  $\delta_{\theta_x}$  = retardations measured in normal and oblique passage of light,

$t$  = the thickness of the specimen (in the reflection technique use  $2t$ ), and

$\theta_x$  = the angle of incidence.

10.1.2 *Strains*—Strains and stresses can be calculated from the measured birefringence when specimen material is optically isotropic in its stress-free state.

10.1.2.1 The difference of principal strains in the plane of the specimen ( $xy$ ) is as follows:

$$\epsilon_x - \epsilon_y = \delta/tk$$

10.1.2.2 In the case of uniaxially stressed material ( $\sigma_x \neq 0$ ,  $\sigma_y = \sigma_z = 0$ ) the principal strains are as follows:

$$\epsilon_x = \delta/(1+\nu)tk$$

$$\epsilon_y = \epsilon_z = -\nu\epsilon_x$$

where:

$\epsilon_x$ ,  $\epsilon_y$ , and  $\epsilon_z$  = the principal strains,

$k$  = the strain optical constant, obtained from references or established by calibration, and

$\nu$  = Poisson's ratio.

10.1.2.3 In the case of biaxially stressed materials ( $\sigma_x \neq 0$  and  $\sigma_y \neq 0$ ) two measurements of retardation are obtained,  $\delta$  and  $\delta_{\theta_x}$  in normal incidence (9.2 and 9.3), and oblique incidence of light (9.5) using an angle  $\theta_x$ :

$$\epsilon_x = \frac{1}{(1+\nu)tk \sin^2 \theta_x} [\delta_{\theta_x}(1-\nu)\cos \theta_x - \delta(\cos^2 \theta_x - \nu)]$$

$$\epsilon_y = \frac{1}{(1+\nu)tk \sin^2 \theta_x} [\delta_{\theta_x}(1-\nu)\cos \theta_x - \delta(1 - \cos^2 \theta_x)]$$

$$\epsilon_z = -\frac{\nu}{1-\nu}(\epsilon_x + \epsilon_y)$$

10.1.2.4 In the case of plastically deformed material and in all instances where (approximately)  $\nu = 0.5$ , the equations in 10.1.2.3 reduce to:

$$\epsilon_x = \frac{1}{1.5tk \sin^2 \theta_x} [0.5\delta_{\theta_x} \cos \theta_x - \delta(\cos^2 \theta_x - 0.5)]$$

$$\epsilon_y = \frac{1}{1.5tk \sin^2 \theta_x} [0.5\delta_{\theta_x} \cos \theta_x - \delta(1 - 0.5 \cos^2 \theta_x)]$$

$$\epsilon_z = -(\epsilon_x + \epsilon_y) = \frac{-1}{1.5tk \sin^2 \theta_x} [\delta_{\theta_x} \cos \theta_x + 0.5\delta \sin^2 \theta_x]$$

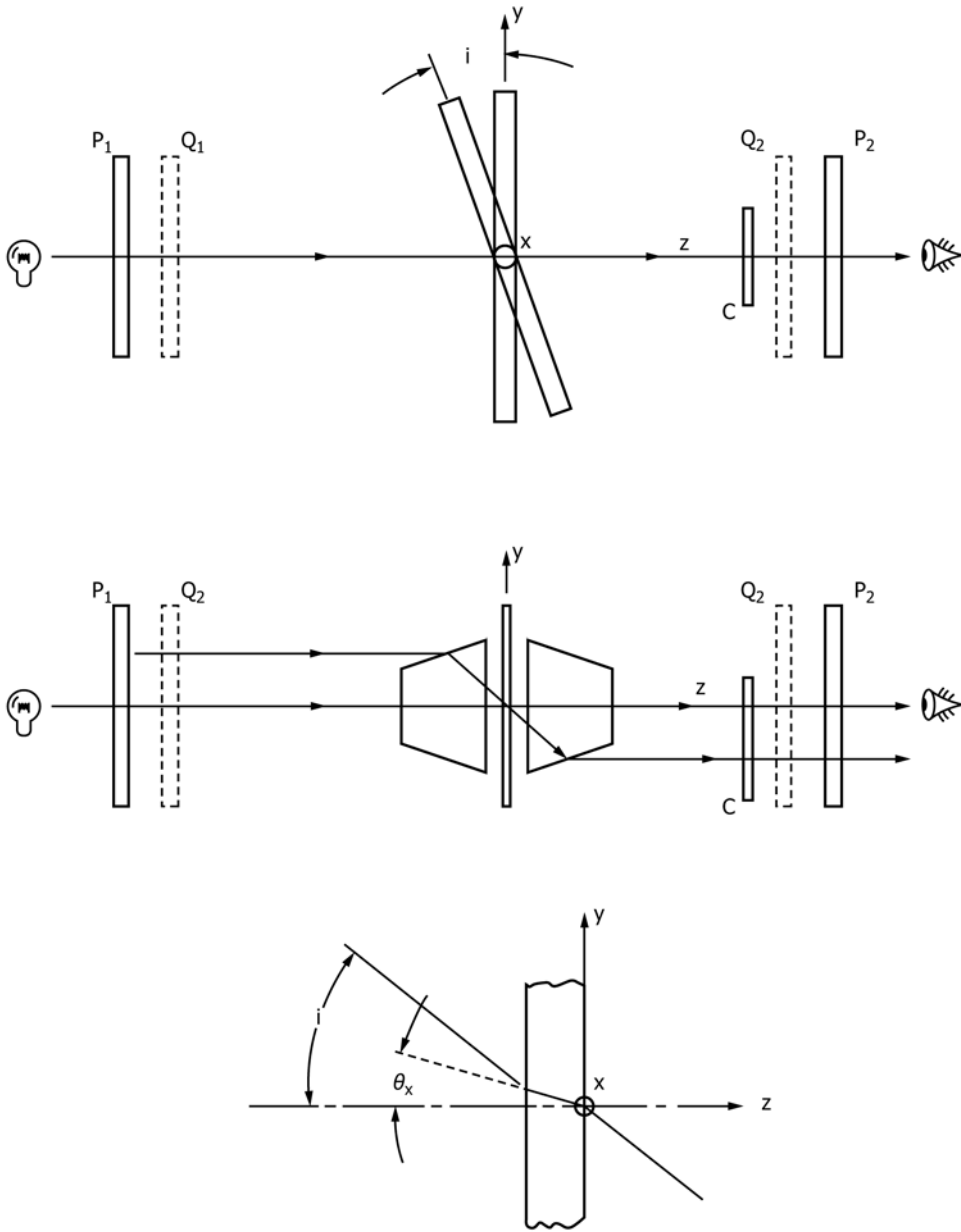


FIG. 8 Oblique Light Passage to a Specimen

10.1.3 *Stresses*—Stresses due to applied forces and elastic residual stresses can be calculated from the measured birefrin-

gence.



**TABLE 1 Precision<sup>A</sup>**

Material	Nominal Thickness, in.	Measured Retardation Mean, nm	$S_r$ , nm	$S_R$ , nm	$I_r$ , nm	$I_R$ , nm
Cellulose triacetate	0.0055 <sup>B</sup>	60	5.8	8.3	16.3	23.6
PETG 6763 copolyester	0.040	353	8.6	31.5	24.3	89.1
Polypropylene	0.00135 <sup>C</sup>	431	10.5	35.2	29.7	99.6
Polycarbonate	0.231	766	14.0	40.0	39.6	113
Cast epoxy	0.084	1091	11.5	33.6	32.5	95.1

<sup>A</sup>  $S_r$  is the within-laboratory standard deviation of the average (median/other function).

$S_R$  is the between-laboratories standard deviation of the average (median/other function).

$I_r = 2.83 S_r$

$I_R = 2.83 S_R$

<sup>B</sup> Stack of five, total thickness 0.0275 in.

<sup>C</sup> Stack of five, total thickness 0.0675 in.

10.1.3.1 When material is optically isotropic and free of birefringence in its stress-free state, the difference of principal stresses in the xy plane is:

$$\sigma_x - \sigma_y = \delta/Ct$$

where  $\sigma_x$ ,  $\sigma_y$  are principal stresses, and  $C$  is Brewster's constant of material, established by calibration.

10.1.3.2 When material exhibits birefringence in its stress-free state (as a result of orientation, crystallinity, plastic deformation, etc.), this initial birefringence (retardation,  $\delta_i$ ) must be subtracted from the measured birefringence (retardation  $\delta_f$ ) before the stresses can be calculated as follows:

$$\delta = \sqrt{\delta_f^2 + \delta_i^2 - 2\delta_f\delta_i \cos 2(\beta_f - \beta_i)}$$

where  $\delta_f$  and  $\delta_i$  are measured retardation and initial retardation measured in stress-free condition, and  $\beta_f$ ,  $\beta_i$  are directions of principal axes, measured, and initial (stress-free condition).

10.1.3.3 In the case of uniaxially stressed material the principal stresses are as follows:

$$\begin{aligned} \sigma_x &= \delta/Ct \\ \sigma_y &= \sigma_z = 0 \end{aligned}$$

where  $C$  is Brewster's material constant established by calibration.

10.1.3.4 In the case of biaxially stressed material ( $\sigma_z = 0$ ) two measurements of retardation,  $\delta$  and  $\delta_{\theta_x}$ , are required:

$$\begin{aligned} \sigma_x &= \frac{(\delta_{\theta_x} \cos \theta_x - \delta \cos^2 \theta_x)}{tC \sin^2 \theta_x} \\ \sigma_y &= \frac{(\delta_{\theta_x} \cos \theta_x - \delta)}{tC \sin^2 \theta_x} \\ \sigma_z &= 0 \end{aligned}$$

10.1.4 In all computations above,  $t$  indicates the thickness of material. When reflection technique is used, the light travels twice through the material and therefore  $2t$  must be used throughout "calculation" paragraph.

## 11. Report

11.1 Report the following information:

11.1.1 Test objectives (or purpose),

11.1.2 Description of tested item(s) and materials,

11.1.3 Set-up used (transmission, reflection),

11.1.4 Calibration data (compensator, stress, or strain-optical material constant), and

11.1.5 Tabulation of measurements (directions, retardation, thickness) and results of calculation of strains (or stresses).

## 12. Precision and Bias<sup>6</sup>

12.1 **Table 1** is based on a round-robin test conducted in 1983 in accordance with Practice **E691**, involving five materials tested by five laboratories. For each material, all the samples were prepared at one source. Each test result was the average of three individual determinations. Each laboratory obtained five test results for each material.

12.2 **Warning**—The following explanations of  $I_r$  and  $I_R$  (see **12.3 – 12.3.3**) are intended only to present a meaningful way of considering the approximate precision of this test method. The data in **Table 1** should not be rigorously applied to acceptance or rejection of material, as those data are specific to the round robin and may not be representative of other lots, conditions, materials, or laboratories.

12.2.1 Users of this test method should apply the principles outlined in Practice **E691** to generate data specific to their laboratory and materials, or between specific laboratories. The principles of **12.3 – 12.3.3** would then be valid for such data.

12.3 **Concept of  $I_r$  and  $I_R$** —If  $S_r$  and  $S_R$  have been calculated from a large enough body of data, and for test results that were averages from testing five specimens:

12.3.1 **Repeatability,  $I_r$** —In comparing two test results for the same material, obtained by the same operator using the same equipment on the same day, the two test results should be judged not equivalent if they differ by more than the  $I_r$  value for that material.

12.3.2 **Reproducibility,  $I_R$** —In comparing two test results for the same material, obtained by different operators using different equipment on different days, the two test results should be judged not equivalent if they differ by more than the  $I_R$  value for that material.

12.3.3 Any judgment in accordance with **12.3.1** and **12.3.2** would have an approximate 95 % (0.95) probability of being correct.

12.4 **Bias**—Bias is systematic error which contributes to the difference between a test result and a true (or reference) value.

<sup>6</sup> Supporting data are available from ASTM Headquarters. Request RR:D20-1121.

There are no recognized standards on which to base an estimate of bias for this test method.

### 13. Keywords

13.1 birefringence; photoelastic measurements; photoelastic retardation; strain; strain-optical constant; transparent plastics

## APPENDIXES

### (Nonmandatory Information)

#### X1. SEQUENCE OF COLORS PRODUCED IN A DARK-FIELD WHITE-LIGHT POLARISCOPE

Color	Retardation, nm <sup>A</sup>	Fringe Order, $\delta/\lambda$
Black	0	0
Gray	160	0.28
White	260	0.45
Yellow	350	0.60
Orange	460	0.79
Red	520	0.90
Tint of Passage 1 <sup>B</sup>	577	1.00
Blue	620	1.06
Blue-green	700	1.20
Green-Yellow	800	1.38
Orange	940	1.62
Red	1050	1.81
Tint of Passage 2 <sup>B</sup>	1150	2.00

Green	1350	2.33
Green-Yellow	1450	2.50
Pink	1550	2.67
Tint of Passage 3 <sup>B</sup>	1730	3.00
Green	1800	3.10
Pink	2100	3.60
Tint of Passage 4 <sup>B</sup>	2300	4.00
Green	2400	4.13

<sup>A</sup>The above sequence is typical for a colorless transparent material. A tinted plastic will change the appearance considerably but will not affect the sequence of the basic colors.

<sup>B</sup>The tint of passage is a sharp dividing zone occurring between red and blue in the first-order fringe, red and green in the second-order fringe, and pink and green in the third-, fourth-, and fifth-order fringes. Beyond five fringes, white-light analysis is not adequate.

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### X3. PROCEDURE FOR EXPERIMENTAL CALIBRATION OF OBLIQUE INCIDENCE ANGLE $\theta_x$

X3.1 In those instances where  $n_o$  is not known accurately or when the angle  $i$  is determined by a fixture rather than a graduated scale, the value of  $\theta_x$  can be determined experimentally. To perform the calibration, introduce in the polariscope a specimen subjected to a uniaxial tension. (The direction of tensile stress will be  $x$  direction.) The state of stress is then:

$\sigma_x$  is applied uniaxial stress

$$\sigma_y = \sigma_z = 0$$

X3.2 Measure the retardation  $\delta$  and  $\delta_{ox}$  using the procedure of this test method, following 9.2 or 9.3 and 9.5. From equations in 10.1.3.4 it follows that:

$$\sigma_y = \frac{(\delta_{ox} \cos \theta_x - \delta)}{tc \sin^2 \theta_x} = 0$$

Then

$$\cos \theta_x = \frac{\delta}{\delta_{ox}}$$

establishing the effective value of the angle  $\theta_x$ .

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