



Designation: E131 – 10 (Reapproved 2015)

Standard Terminology Relating to Molecular Spectroscopy^{1, 2}

This standard is issued under the fixed designation E131; 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 The values stated in SI units are to be regarded as standard. No other units of measurement are included in this standard.

2. Referenced Documents

2.1 ASTM Standards:³

E135 Terminology Relating to Analytical Chemistry for Metals, Ores, and Related Materials

E168 Practices for General Techniques of Infrared Quantitative Analysis (Withdrawn 2015)⁴

E204 Practices for Identification of Material by Infrared Absorption Spectroscopy, Using the ASTM Coded Band and Chemical Classification Index (Withdrawn 2014)⁴

E284 Terminology of Appearance

E386 Practice for Data Presentation Relating to High-Resolution Nuclear Magnetic Resonance (NMR) Spectroscopy

E456 Terminology Relating to Quality and Statistics

2.2 Other Documents:⁵

ISO Guide 30–1981 (E) Terms and definitions used in connections with reference materials

3. Terminology

absorbance, A —the logarithm to the base 10 of the reciprocal of the transmittance, (T).

$$A = \log_{10}(1/T) = -\log_{10}T \quad (1)$$

DISCUSSION—In practice the observed transmittance must be substituted for T . Absorbance expresses the excess absorption over that of a specified reference or standard. It is implied that compensation has

been effected for reflectance losses, solvent absorption losses, and refractive effects, if present, and that attenuation by scattering is small compared with attenuation by absorption. Apparent deviations from the absorption laws (see **absorptivity**) are due to inability to measure exactly the true transmittance or to know the exact concentration of an absorbing substance.

absorption band—a region of the absorption spectrum in which the absorbance passes through a maximum.

absorption coefficient, α —a measure of absorption of radiant energy from an incident beam as it traverses an absorbing medium according to Bouguer's law, $P/P_o = e^{-\alpha b}$.

DISCUSSION—In IRS, α is a measure of the rate of absorption of energy from the evanescent wave.

absorption parameter, a —the relative reflection loss per reflection that results from the absorption of radiant energy at a reflecting surface: $a = 1 - R$, and R = the reflected fraction of incident radiant power.

absorption spectrum—a plot, or other representation, of absorbance, or any function of absorbance, against wavelength, or any function of wavelength.

absorptivity, a —the absorbance divided by the product of the concentration of the substance and the sample pathlength, $a = A/bc$. The units of b and c shall be specified.

DISCUSSION—1—The recommended unit for b is the centimetre. The recommended unit for c is kilogram per cubic metre. Equivalent units are g/dm^3 , g/L , or mg/cm^3 .

DISCUSSION—2—The equivalent IUPAC term is “specific absorption coefficient.”

absorptivity, molar, ϵ —the product of the absorptivity, a , and the molecular weight of the substance.

DISCUSSION—The equivalent IUPAC term is “molar absorption coefficient.”

acceptance angle, n —for an optical fiber, the maximum angle, measured from the longitudinal axis or centerline of the fiber to an incident ray, within which the ray will be accepted for transmission along the fiber by total internal reflection.

DISCUSSION—If the incidence angle exceeds the acceptance angle, optical power in the incident ray will be coupled into leaky modes or rays, or lost by scattering, diffusion, or absorption in the cladding. For a cladded step-index fiber in the air, the sine of the acceptance angle is given by the square root of the difference of the squares of the refractive indexes of the fiber core and the cladding, that is, by the relation as follows:

¹ This terminology is under the jurisdiction of ASTM Committee E13 on Molecular Spectroscopy and Separation Science and is the direct responsibility of Subcommittee E13.94 on Terminology.

Current edition approved May 1, 2015. Published June 2015. Originally approved in 1957. Last previous edition approved in 2010 as E131 – 10. DOI: 10.1520/E0131-10R15.

² For other definitions relating to nuclear magnetic resonance, see Practice E386.

³ 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.

⁴ The last approved version of this historical standard is referenced on www.astm.org.

⁵ Available from American National Standards Institute (ANSI), 25 W. 43rd St., 4th Floor, New York, NY 10036, http://www.ansi.org.

$$\sin A = \sqrt{n_1^2 - n_2^2} \quad (2)$$

where A is the acceptance angle and n_1 and n_2 are the refractive indexes of the core and cladding, respectively. If the refractive index is a function of distance from the center of the core, as in the case of graded index fibers, then the acceptance angle depends on the distance from the core center. The acceptance angle is maximum at the center, and zero at the core-cladding boundary. At any radius, r , the sine of the acceptance angle of a graded index fiber is defined in compliance with that of a step-index fiber as follows:

$$\sin A_r = \sqrt{n_1^2 - n_2^2} \quad (3)$$

where A_r is the acceptance angle at a point on the entrance face at a distance, r , from the center, n_r is the refractive index of the core at a radius, r , and n_2 is the refractive index of the cladding. In air, $\sin A$ and $\sin A_r$ are the numerical apertures. Unless otherwise stated, acceptance angles and numerical apertures for fiber optics are those for the center of the endface of the fiber, that is, where the refractive index, and hence the numerical aperture, is the highest.

accuracy—the closeness of agreement between an observed value and an accepted reference value (see Terminology E456).

DISCUSSION—The term accuracy, when applied to a set of observed values, will be a combination of a random component and a common systematic error or bias component. Since in routine use, random components and bias components cannot be completely separated, the reported “accuracy” must be interpreted as a combination of these two components.

active fiber optic chemical sensor, n —a fiber optic chemical sensor in which a transduction mechanism other than the intrinsic spectroscopic properties of the analyte is used to modulate the optical signal.

DISCUSSION—Examples include a pH sensor composed of a chemical indicator substance whose color changes with pH, and an oxygen sensor coupled to an optical fiber bearing a chemical indicator whose fluorescence intensity depends on oxygen concentration.

aliasing—the appearance of features at wavenumbers other than their true value caused by using a sampling frequency less than twice the highest modulation frequency in the interferogram; also known as “folding.”

analytical curve—the graphical representation of a relation between some function of radiant power and the concentration or mass of the substance emitting or absorbing it.

analytical wavelength—any wavelength at which an absorbance measurement is made for the purpose of the determination of a constituent of a sample.

angle of incidence, θ —the angle between an incident radiant beam and a perpendicular to the interface between two media.

anti-Stokes line (band)—a Raman line (band) that has a frequency higher than that of the incident monochromatic beam.

aperture of an IRE, A' —that portion of the IRE surface that can be utilized to conduct light into the IRE at the desired angle of incidence.

apodization—modification of the ILS function by multiplying the interferogram by a weighting function the magnitude of which varies with retardation.

DISCUSSION—This term should strictly be used with reference to a weighting function whose magnitude is greatest at the centerburst and decreases with retardation.

attenuated total reflection (ATR)—reflection that occurs when an absorbing coupling mechanism acts in the process of total internal reflection to make the reflectance less than unity.

DISCUSSION—In this process, if an absorbing sample is placed in contact with the reflecting surface, the reflectance for total internal reflection will be attenuated to some value between zero and unity ($0 < R < 1$) in regions of the spectrum where absorption of the radiant power can take place.

attenuation index, κ —a measure of the absorption of radiant energy by an absorbing material. κ is related to the absorption coefficient by: $n\kappa = ac_o/4\pi v$, where c_o = the speed of light in vacuo, v = the frequency of radiant energy, and n = the refractive index of the absorbing medium.

background—apparent absorption caused by anything other than the substance for which the analysis is being made.

baseline—any line drawn on an absorption spectrum to establish a reference point representing a function of the radiant power incident on a sample at a given wavelength.

basic NMR frequency, ν_o —the frequency, measured in hertz (Hz), of the oscillating magnetic field applied to induce transitions between nuclear magnetic energy levels.

bathochromic shift, n —change of a spectral band to longer wavelength (lower frequency) because of structural modifications or environmental influence; also known as “red shift.”

beamsplitter—a semireflecting device used to create, and often to recombine, spatially separate beams.

DISCUSSION—Beamsplitters are often made by depositing a film of a high refractive index material onto a flat transmitting substrate with an identical compensator plate being held on the other side of the film.

beamsplitter efficiency—the product $4RT$, where R is the reflectance and T is the transmittance of the beamsplitter.

Beer’s law—the absorbance of a homogeneous sample containing an absorbing substance is directly proportional to the concentration of the absorbing substance (see also **absorbivity**)

bias—a systematic error that contributes to the difference between a population mean of the measurements or test results and an accepted or reference value (see Terminology E456).

DISCUSSION—Bias is determined by the following equation:

$$\text{bias} = \bar{e} = \frac{1}{n} \sum_{i=1}^n e_i \quad (4)$$

where:

n = the number of observations for which the accuracy is determined,

e_i = the difference between a measured value of a property and its accepted reference value, and

\bar{e} = the mean value of all the e_i .

Bouguer's law—the absorbance of a homogeneous sample is directly proportional to the thickness of the sample in the optical path.

DISCUSSION—Bouguer's law is sometimes also known as Lambert's law.

boxcar truncation—identical effective weighting of all points in the measured interferogram prior to the Fourier transform; all points outside of the range of the measured interferogram take a value of zero.

buffer—*in fiber optics*, see **fiber optic buffer**.

bulk reflection—reflection in which radiant energy is returned exclusively from within the specimen.

DISCUSSION—Bulk reflection may be diffuse or specular.

centerburst—the region of greatest amplitude in an interferogram.

DISCUSSION—For unchirped or only slightly chirped interferograms, this region includes the “zero path difference point” and the “zero retardation point.”

certified reference material, n —a reference material, the composition or properties of which are certified by a recognized standardizing agency or group.

DISCUSSION—A certified reference material produced by the National Institute of Standards and Technology (NIST) is designated a Standard Reference Material (SRM).

chemical shift (NMR), δ —the defining equation for δ is the following:

$$\delta = \frac{\Delta\nu}{\nu_R} \times 10^6 \quad (5)$$

where ν_R is the frequency with which the reference substance is in resonance at the magnetic field used in the experiment and $\Delta\nu$ is the frequency difference between the reference substance and the substance whose chemical shift is being determined, at constant field. The sign of $\Delta\nu$ is to be chosen such that shifts to the high frequency side of the reference shall be positive.

DISCUSSION—If the experiment is done at constant frequency (field sweep) the defining equation becomes

$$\delta = \frac{\Delta\nu}{\nu_R} \times \left(1 - \frac{\Delta\nu}{\nu_R} \right) \times 10^6 \quad (6)$$

chirping—the process of dispersing the zero phase difference points for different wavelengths across the interferogram, so that the magnitude of the signal is reduced in the short region of the interferogram where all wavelengths would otherwise constructively interfere.

clad—see **cladding**.

cladding, n —*of an optical fiber*, a layer of a optically transparent lower refractive index material in intimate contact with a core of higher refractive index material used to achieve total internal reflection.

DISCUSSION—The cladding confines electromagnetic waves to the core, provides some protection to the core, and also transmits evanescent waves that usually are bound to waves in the core.

concentration, c —the quantity of the substance contained in a unit quantity of sample.

DISCUSSION—For solution work, the recommended unit of concentration is grams of solute per litre of solution.

core, n —*of an optical fiber*, the center region of an optical waveguide through which radiant energy is transmitted.

DISCUSSION—In a dielectric waveguide such as an optical fiber, the refractive index of the core must be higher than that of the cladding. Most of the radiant energy is confined to the core.

correlation coefficient (r)—a measure of the strength of the linear relationship between X and Y , calculated by the equation:

$$r_{xy} = \frac{\left(\sum_{i=1}^n X_i Y_i \right)}{\left(\sum_{i=1}^n X_i^2 \right)^{1/2} \left(\sum_{i=1}^n Y_i^2 \right)^{1/2}} \quad (7)$$

where:

n = the number of observations in X and Y .

DISCUSSION— X_i and Y_i are any two mean corrected variables. For the simple linear regression only,

$$r_{xy} = R = (\text{sign of } b_1)(R^2)^{1/2} \quad (8)$$

where:

R^2 = the coefficient of multiple determination.

critical angle, θ_c —the angle whose sine is equal to the relative refractive index for light striking an interface from the greater to the lesser refractive medium: $\theta_c = \sin^{-1} n_{21}$, where n_{21} = the ratio of the refractive indices of the two media.

DISCUSSION—Total reflection occurs when light is reflected in the more refractive of two media from the interface between them at any angle of incidence exceeding the critical angle.

depth of penetration, d_p —in internal reflection spectroscopy, the distance into the less refractive medium at which the amplitude of the evanescent wave is e^{-1} (that is, 36.8 %) of its value at the surface:

$$d_p = \frac{\lambda_1}{2\pi(\sin^2\theta - n_{21}^2)^{1/2}} \quad (9)$$

where: $n_{21} = n_2/n_1$ = refractive index of sample divided by that of the IRE; $\lambda_1 = \lambda/n_1$ = wavelength of radiant energy in the sample; and θ = angle of incidence.

derivative absorption spectrum—a plot of rate of change of absorbance or of any function of absorbance with respect to wavelength or any function of wavelength, against wavelength or any function of wavelength.

difference absorption spectrum—a plot of the difference between two absorbances or between any function of two absorbances, against wavelength or any function of wavelength.

diffuse reflection—reflection in which the flux is scattered in many directions by diffusion at or below the surface (see Terminology E284).

digitization—the conversion of an analog signal to digital values using an analog-to-digital converter “sampling” or “digital sampling.”

digitization noise—the noise generated in an interferogram through the use of an analog-to-digital converter whose least

significant bit represents a value comparable to, or greater than, the peak-to-peak noise level in the analog data.

dilution factor—the ratio of the volume of a diluted solution to the volume of original solution containing the same quantity of solute as the diluted solution.

double modulation, n —a technique in which a modulated signal is further varied by a second means.

DISCUSSION—As an example, a spectrometer could generate a modulated signal while at the same time that signal is further varied by an external higher frequency modulator; on detection, the conventional spectrometric signal is filtered out so that only the high frequency signal is recorded.

double-pass internal reflection element—an internal reflection element in which the radiant power transverses the length of the optical element twice, entering and leaving via the same end.

effective pathlength (or effective thickness), d_e —in *internal reflection spectroscopy*, the analog of the sample thickness in transmission spectroscopy that represents the distance of propagation of the evanescent wave within an absorbing sample in IRS. It is defined from the relationship: $R = 1 - \alpha d_e$, and is related to the absorption parameter by: $a = \alpha d_e$.

evanescent wave—the standing wave that exists in the less refractive medium, normal to the reflecting surface of the IRE during internal reflection.

extrinsic fiber optic chemical sensor, n —a fiber optic chemical sensor in which modulation of the optical signal is not effected through a change in the properties of the fiber itself.

DISCUSSION—Examples include a pH sensor composed of a chemical indicator immobilized at the end of the optical fiber, and a sensor based on Raman, fluorescence, infrared, visible, or other spectral information gathered in the acceptance cone of the fiber.

far-infrared—pertaining to the infrared region of the electromagnetic spectrum with wavelength range from approximately 25 to 1000 μm (wavenumber range 400 to 10 cm^{-1}).

fast Fourier transform (FFT)—a method for speeding up the computation of a discrete FT by factoring the data into sparse matrices containing mostly zeroes.

fiber optic buffer, n —material placed on or around a cladded optical fiber to protect it from mechanical damage.

DISCUSSION—Mechanical damage can be caused by such things as microbends and macrobends formed during manufacture, spooling, subsequent handling, and pressure applied during use. Buffers may be bonded to the cladding and may also serve the purpose of preventing ambient energy from entering the core.

fiber optic chemical sensor, n —a fiber optic sensor that responds to a chemical stimulus.

fiber optic sensor, n —a device that responds to an external stimulus and transmits through an optical fiber a modulated optical signal, indicating one or more characteristics of the stimulus.

DISCUSSION—Examples include sensors which provide a suitable signal or impulse to a meter. Such sensors might be found as the active elements in pH meters, strain gages, or pressure gages.

fiber optics, n —the branch of science and technology devoted to the transmission of radiant energy through fibers made of transparent materials.

DISCUSSION—Transparent materials include glass, fused silica, and plastic. Optical fibers in fiber optic cables may be used for data transmission, and for sensing, illumination, endoscopic, control, and display purposes, depending on their use in various geometric configurations, modes of excitation, and environmental conditions. The fibers may be wound and bound in various shapes and distributions singly or in bundles. Bundles may be aligned or unaligned. Aligned bundles are often used to transmit and display images.

filter—a substance that attenuates the radiant power reaching the detector in a definite manner with respect to spectral distribution.

filter, neutral—a filter that attenuates the radiant power reaching the detector by the same factor at all wavelengths within a prescribed wavelength region.

fixed-angle internal reflection element—an internal reflection element which is designed to be operated at a fixed angle of incidence.

fluorescence—the emission of radiant energy from an atom, molecule, or ion resulting from absorption of a photon and a subsequent transition to the ground state without a change in total spin quantum number.

DISCUSSION—The initial and final states of the transition are usually both singlet states. The average time interval between absorption and fluorescence is usually less than 10^{-6} s.

folding—see **aliasing**.

Fourier transform (FT)—the mathematical process used to convert an amplitude-time spectrum to an amplitude-frequency spectrum, or *vice versa*.

DISCUSSION—In FT-IR spectrometry, retardation is directly proportional to time; therefore the FT is commonly used to convert an amplitude-retardation spectrum to an amplitude-wavenumber spectrum, and *vice versa*.

Fourier transform infrared (FT-IR) spectrometry—a form of infrared spectrometry in which an interferogram is obtained; this interferogram is then subjected to a Fourier transform to obtain an amplitude-wavenumber (or wavelength) spectrum.

DISCUSSION—1—The abbreviation FTIR is not recommended.

DISCUSSION—2—When FT-IR spectrometers are interfaced with other instruments, a slash should be used to denote the interface; for example, GC/FT-IR; HPLC/FT-IR, and the use of FT-IR should be explicit; that is, FT-IR not IR.

frequency, ν —the number of cycles per unit time.

DISCUSSION—The recommended unit is the hertz (Hz) (one cycle per second).

frustrated total reflection (FTR)—the reflection which occurs when a nonabsorbing coupling mechanism acts in the process of total internal reflection to make the reflectance less than unity.

DISCUSSION—In the process the reflectance can vary continuously between zero and unity if: (1) An optically transparent medium is within a fraction of a wavelength of the reflecting surface and its distance from the reflecting surface is changed, or (2) Both the angle of incidence and the refractive index of one of the media vary in an

appropriate manner.

In these cases part of the radiant power may be transmitted through the interface into the second medium without loss at the reflecting surface such that transmittance plus reflectance equals unity. It is possible, therefore to have this process taking place in some spectral regions even when a sample having absorption bands is placed in contact with the reflecting surface.

high-resolution NMR spectrometer—an NMR apparatus that is capable of producing, for a given isotope, line widths that are less than the majority of the chemical shifts and coupling constants for that isotope.

DISCUSSION—By this definition, a given spectrometer may be classed as a high-resolution instrument for isotopes with large chemical shifts, but may not be classed as a high-resolution instrument for isotopes with smaller chemical shifts.

hole-burning, n —in luminescence, the photo-induced disappearance of a narrow segment within a broader absorption or emission band.

DISCUSSION—Holes are produced by the disappearance of resonantly excited molecules because of photochemical or photophysical processes.

infrared—pertaining to the region of the electromagnetic spectrum with wavelength range from approximately 0.78 to 1000 μm (wavenumber range 12 800 to 10 cm^{-1}).

infrared spectroscopy—pertaining to spectroscopy in the infrared region of the electromagnetic spectrum.

DISCUSSION—1—Spectroscopy and other related terms are defined in Terminology E135.

DISCUSSION—2—Common applications of infrared spectroscopy are the identification of materials and the quantitative analysis of materials (see, for example, Practices E204 and Practices E168).

instrument line shape (ILS) function—the FT of the function by which an interferogram is weighted.

DISCUSSION—This weighting may be performed optically, due to the finite optical throughput, or digitally, through multiplication by an apodization function, or both. The ILS function is the profile of the spectrum of a monochromatic source producing a beam with the same throughput as the beam in the actual measurement being performed.

instrument response time—the time required for an indicating or detecting device to undergo a defined displacement following an abrupt change in the quantity being measured.

integration period, π —the time, in seconds, required for the pen or other indicator to move 98.6 % of its maximum travel in response to a step function.

DISCUSSION—For instruments with a first-order response, the integration period will be approximately equal to four times the exponential time constant. It is equal to the period, classically defined, for a second order, critically damped response system.

intercorrelation coefficient, (r_{XX}) —a measure of the linear association between values of the same type of variable expressed as a correlation coefficient, (r).

DISCUSSION—The variables X and Y are replaced by X_j and X_k in the equation for the correlation coefficient, r .

interferogram, $I(\delta)$ —record of the modulated component of the interference signal measured as a function of retardation by the detector.

DISCUSSION—1—An alternate symbol is $I(x)$.

DISCUSSION—2—The recommended symbol for the spectrum computed from $I(\delta)$ is $B(v)$. An alternate symbol is $B(\sigma)$.

interferogram, double-sided—interferogram measured with approximately equal retardation on either side of the centerburst.

interferogram, laser reference—sinusoidal interferogram of a laser source measured at the same time as the signal interferogram.

DISCUSSION—The zero crossings of this interferogram are used to control sampling of the signal interferogram. It may also be noted that other effectively monochromatic sources can be used in place of the laser.

interferogram, signal—interferogram of the beam of radiant energy whose spectrum is desired.

interferogram, single-sided—interferogram in which sampling is initiated close to the centerburst and continues through that point to the maximum retardation desired.

interferogram, white light—reference interferogram of a broadband light source measured at the same time as the signal interferogram and used to initiate data acquisition of consecutive scans for signal-averaging.

interferometer—device used to divide a beam of radiant energy into two or more paths, generate an optical path difference between the beams, and recombine them in order to produce repetitive interference maxima and minima as the optical retardation is varied.

interferometer, Genzel—interferometer in which the beam is focused in the plane of the beamsplitter and collimated before the moving mirror(s).

interferometer, lamellar grating—interferometer in which the beam is reflected from two interleaved mirrors, one of which is stationary while the other is movable.

DISCUSSION—This type of interferometer is generally used only for far infrared spectrometry.

interferometer, Michelson—interferometer in which an approximately collimated beam of radiant energy is divided into two paths by a beamsplitter; one beam is reflected from a movable mirror and the other from a stationary mirror, and they are then recombined at the beamsplitter.

interferometer, rapid-scanning—interferometer in which the retardation is varied rapidly enough that the modulation frequencies in the interferogram are sufficiently high that the interferogram signal can be amplified directly without additional modulation by an external chopper.

interferometer, refractively scanned—interferometer in which the retardation between two beams is generated by the movement of a wedged optical element.

interferometer, slow-scanning—interferometer in which the retardation is continuously varied, but so slowly that an external chopper is needed to modulate the beam at a frequency which is high enough for ac signal amplification.

interferometer, stepped-scanning—interferometer in which the movable element is held stationary for the length of time

required for signal integration and digitization of each sample point, and then translated to the next sample point.

internal conversion, n —a transition between electronic states of the same total spin quantum number (multiplicity).

internal, reflection attachment, IRA—the transfer optical system which supports the IRE, directs the energy of the radiant beam into the IRE, and then redirects the energy into the spectrometer or onto the detector. The IRA may be part of an internal reflection spectrometer or it may be placed into the sampling space of a spectrometer.

internal reflection element (IRE)—the transparent optical element used in internal reflection spectroscopy for establishing the conditions necessary to obtain the internal reflection spectra of materials.

DISCUSSION—Radiant power is propagated through it by means of internal reflection. The sample material is placed in contact with the reflecting surface or it may be the reflecting surface itself. If only a single reflection takes place from the internal reflection element the element is said to be a single reflection element; if more than one reflection takes place, the element is said to be a multiple reflection element. When the element has a recognized shape it is identified according to each shape, for example, internal reflection prism, internal reflection hemicylinder, internal reflection plate, internal reflection rod, internal reflection fiber, etc.

internal reflection spectroscopy (IRS)—the technique of recording optical spectra by placing a sample material in contact with a transparent medium of greater refractive index and measuring the reflectance (single or multiple) from the interface, generally at angles of incidence greater than the critical angle.

intersystem crossing—a transition between electronic states that differ in total spin quantum number (multiplicity).

DISCUSSION—Current experimental evidence indicates this process is nonradiative.

intrinsic fiber optic chemical sensor, n —a fiber optic chemical sensor in which the modulation of the optical signal is effected through a change in the properties of the optical fiber itself, and such modulation occurs while the radiant energy is guided by the optical fiber.

irreversible fiber optic chemical sensor, n —a fiber optic chemical sensor that undergoes a permanent depletion or degradation of the transduction element as a result of the transduction process.

DISCUSSION—An example is a sensor based on an indicator that reacts irreversibly with the target analyte and that cannot be replenished after measurement.

isoabsorptive point—a wavelength at which the absorptivities of two or more substances are equal.

isosbestic point—the wavelength at which the absorptivities of two substances, one of which can be converted into the other, are equal.

isostilbic point, n —in luminescence, the wavelength at which the intensity of emission of a sample does not change during a physical interaction or chemical reaction.

level one (1) test, n —a simple series of measurements designed to provide quantitative data on various aspects of instrument performance and information on which to base the diagnosis of problems.

level zero (0) test, n —a routine check of instrument performance, that can be done in a few minutes, designed to virtually detect significant changes in instrument performance and provide a database to determine instrument function over time.

linear dispersion—the derivative, $dx/d\lambda$, where x is the distance along the spectrum, in the plane of the exit slit, and λ is the wavelength.

linearity—the property of paired (X, Y) data such that when an equation for a straight line is calculated for that data using linear least-square regression mathematics, no statistically significant reduction to the sum-squared difference of the data from that line is achieved by the addition of another function to that equation for the straight line.

lock signal (NMR)—the NMR signal used to control the field-frequency ratio of the spectrometer. It may or may not be the same as the reference signal.

luminescence—the emission of radiant energy during a transition from an excited electronic state of an atom, molecule, or ion to a lower electronic state.

DISCUSSION—1—The recommended unit for “sample pathlength” is centimetres. This distance does not include the thickness of the walls of any absorption cell in which the specimen is contained.

DISCUSSION—2—In strict usage, a more appropriate term would be “specimen pathlength.” This is currently under advisement by Committee E13.

mid-infrared—pertaining to the infrared region of the electromagnetic spectrum with wavelength range from approximately 2.5 to 25 μm (wavenumber range 4000 to 400 cm^{-1}).

modulate, v —to vary a characteristic or parameter of an entity in accordance with a characteristic or parameter of another entity.

modulation frequency, f_v —the frequency, in Hz, at which radiant energy of a given wavenumber is modulated by a rapid-scanning interferometer.

DISCUSSION—1—This is given by the product of the wavenumber (cm^{-1}) and the rate of change of retardation ($\text{cm}\cdot\text{s}^{-1}$).

DISCUSSION—2—An alternate symbol is f_o .

molar absorptivity, ϵ —see **absorptivity, molar**.

monochromator—a device or instrument that, with an appropriate energy source, may be used to provide a continuous calibrated series of electromagnetic energy bands of determinable wavelength or frequency range.

multiple correlation coefficient, (R)—the correlation, r_{yy} , between the accepted reference values, Y_i , and the values determined using the calibration equation, \hat{Y}_i , equal to the square root of the coefficient of multiple determination, R^2 .