



(12) **EUROPEAN PATENT SPECIFICATION**

(45) Date of publication and mention
of the grant of the patent:

12.10.2005 Bulletin 2005/41

(21) Application number: **01962218.2**

(22) Date of filing: **16.08.2001**

(51) Int Cl.⁷: **G01N 21/27**

(86) International application number:
PCT/US2001/025706

(87) International publication number:
WO 2002/025233 (28.03.2002 Gazette 2002/12)

(54) **METHOD OF CHARACTERIZING SPECTROMETERS**

VERFAHREN ZUR CHARAKTERISIERUNG VON SPEKTROMETERN

PROCEDE DE CHARACTERISATION D'INSTRUMENTS SPECTROMETRIQUES

(84) Designated Contracting States:
**AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE TR**

(30) Priority: **18.09.2000 US 664973**

(43) Date of publication of application:
18.06.2003 Bulletin 2003/25

(60) Divisional application:
04010284.0 / 1 442 699

(73) Proprietor: **Sensys Medical, Inc.**
Chandler, AZ 85226 (US)

(72) Inventors:
• **HAZEN, Kevin, H.**
Gilbert, AZ 85233 (US)
• **BLANK, Thomas, B.**
Chandler, AZ 85225 (US)
• **MONFRE, Stephen, L.**
Gilbert, AZ 85296 (US)

• **RUCHTI, Timothy, L.**
Gilbert, AZ 85233 (US)

(74) Representative: **Leeming, John Gerard et al**
J.A. Kemp & Co.,
14 South Square,
Gray's Inn
London WC1R 5JJ (GB)

(56) References cited:
EP-A- 0 768 522 **WO-A-00/42907**
US-A- 5 204 532 **US-A- 5 708 593**
US-A- 5 710 713

• **BOUVERESSE E, HARTMANN C, MASSART D L,**
LAST I R, PREBBLE K A: "Standardization of
Near-Infrared Spectrometric Instruments"
ANALYTICAL CHEMISTRY, vol. 68, no. 6, 15
March 1996 (1996-03-15), pages 982-990,
XP002193004 cited in the application

Note: Within nine months from the publication of the mention of the grant of the European patent, any person may give notice to the European Patent Office of opposition to the European patent granted. Notice of opposition shall be filed in a written reasoned statement. It shall not be deemed to have been filed until the opposition fee has been paid. (Art. 99(1) European Patent Convention).

Description

BACKGROUND OF THE INVENTION

FIELD OF THE INVENTION

[0001] The invention relates to variation in spectrometer instruments. More particularly the invention relates to characterizing spectrometer instruments by classifying their spectral responses into a limited number of clusters and developing calibration transfer models between clusters that compensate for instrument variations.

DESCRIPTION OF THE PRIOR ART

[0002] Many of the analytical applications for spectrometers require calibration data sets that are time-consuming and expensive to create. Typically, these calibrations are highly specific. For example, apparently identical instruments produced by the same manufacturer may exhibit minor instrument variations; such variations may be seen when one instrument is built with a component that varies slightly from the same component in another instrument. In addition, a calibration set for an instrument produced by one manufacturer is generally not suitable for a similar instrument produced by another manufacturer. Furthermore, repairs to a single instrument can cause the instrument's spectral response to vary. As an instrument ages, its spectral response may change. An instrument's spectral response may vary according to fluctuations in the operating environment. In applications requiring analysis of very low concentration analytes, non-invasive blood glucose prediction, for example, even minor instrument variation can introduce an unacceptable degree of error into the analysis. Providing another calibration model that takes the instrument's current spectral response into account can compensate for instrument variation. However, development of new calibration models is time-consuming, labor-intensive and costly.

[0003] In the development of spectroscopy-based analyzers for biomedical applications, there is a need for production of thousands to as many as millions of analyzers for a specific application. No methodology exists for providing calibrations for large numbers of instruments quickly and inexpensively.

[0004] Therefore, efforts have been directed at transferring calibrations from one analyzer to another. See, for example, E. Bouveresse, C. Hartmann, D. Massart, I. Last, K. Prebble, *Standardization of near-infrared spectrometric instruments*, *Anal. Chem.*, vol. 68, pp. 982-990 (1996) and M. Defernez, R. Wilson, *Infrared spectroscopy: instrumental factors affecting the long-term validity of chemometric models*, *Anal. Chem.*, vol. 69, pp. 1288-1294 (1997), and E. Bouveresse, D. Massart, P. Dardenne, *Calibration transfer across near-infrared spectrometric instruments using Shenk's algo-*

rithm: effects of different standardization samples, *Analytica Chimica Acta*, vol. 297, pp. 405-416, (1994) and Y. Wang, D. Veltkamp, B. Kowalski, *Multivariate instrument calibration*, *Anal. Chem.*, vol. 63, pp. 2750-2756 (1991).

[0005] Most of the reported methods of calibration transfer have been applied in situations involving high-concentration analytes, wherein the signal-to-noise ratio is high. Because these currently known methods act as a smoothing function when transferring calibrations, they degrade the signal to noise that can be observed, thus hindering analysis of low concentration analytes. Additional problems of changes in resolution or bandwidth across time or between instruments have not been addressed.

[0006] Furthermore, the currently known methods have only been successfully applied in situations involving a small number of instruments. The reported methods are not capable of modeling the complexity encountered when large numbers of instruments are involved.

[0007] R. Rosenthal, Method for providing a general calibration for near infrared instruments for measurement of blood glucose, United States Patent No. 5,204,532 (20 April 1993) describes a method for calibrating a near infrared analysis instrument to an individual user by selecting a spectral cluster having the calibration constants most closely associated with the user's spectrum.

[0008] S. Malin, T. Ruchti, System and method for noninvasive blood analyte measurements, PCT Application No. PCT/US00/01378 (WO 00/42907) (19 January 2000) describes an intelligent system for blood analyte determination wherein the sample to be analyzed is classified into one or more predefined clusters based on extracted spectral features, and a suitable calibration model applied.

[0009] A need exists for the calibration of large numbers of analyzers. It would be desirable to provide a means of reducing the complexity inherent in the transfer of calibrations to large numbers of analyzers. It would also be advantageous to provide a means of transferring calibrations without significant degradation of the signal-to-noise ratio, rendering calibration transfer practical in analysis of low concentration analytes.

[0010] The invention provides a method, defined in the appended claims, of characterizing large numbers of spectrometer instruments by classifying their spectra into a limited number of previously defined clusters according to similarity of spectral features and performance characteristics. The method may also be used to track variation over time within a single instrument. The spectral features used for classification may be related to known instrument parameters, or they may be abstract features derived using a variety of computational methods. The clusters are defined in advance based on an exemplar data set, using either supervised or unsupervised methods. Calibration models for each cluster compensate for instrument variation, either

across instruments or across time within a single instrument.

[0011] In a preferred embodiment of the invention, calibration models are provided using a method of calibration transfer wherein the clusters are mapped to each other, so that a calibration transferred from one cluster to another need only model the difference between the two clusters. In an alternate embodiment, a different calibration is separately calculated for each cluster. In either embodiment, the number of calibration transfers is significantly reduced since each cluster represents a multitude of instruments.

BRIEF DESCRIPTION OF THE DRAWINGS

[0012]

Figure 1 provides a block schematic diagram of a method of characterizing spectrometer instruments by clustering according to spectral features, according to the invention;

Figure 2 provides a flow diagram of a method of generating calibration models by transferring a master calibration to slave calibration models, according to the invention;

Figures 3A and 3B illustrate decreases in spectral cutoff related to decreases in detector temperature in a spectrometer instrument; and

Figure 4 illustrates changes in light throughput related to changes in source temperature in a spectrometer instrument.

DETAILED DESCRIPTION

[0013] The invention provides a method of characterizing spectrometer instruments by classifying spectral responses from a large number of spectrometers (analyzers) into a limited number of previously defined clusters. The clusters constitute groups that are defined based on the similarity of specific features. Grouping of instruments according to spectral features and performance characteristics reduces the spectral variation between instruments in a given group. Therefore, spectral measurements corresponding to an individual cluster are more homogeneous than those from the entire group of analyzers. Calibrations created for each cluster may then be utilized for subsequent analysis. The calibration created for a cluster will have instrument variations included in the model that mirror variations resulting from the particular analyzer employed for the subsequent analysis. Calibration models specific to the clusters are expected to be less complex and have an improved level of accuracy. Therefore, multivariate analysis will require fewer factors to model instrument variation resulting in earlier factors modeling the sample

and earlier factors modeling small absorbing analytes. The net result is that fewer factors are required for sample analysis and a more robust algorithm is created.

[0014] In one embodiment of the invention, a separate calibration data set is required for every cluster, requiring a large number of samples in a calibration set for every cluster. In the preferred embodiment, sufficient samples for a calibration model are only required in a single cluster. Principal features defining that cluster and other clusters are determined, and the clusters are mapped to each other, revealing the specific differences from one cluster to another. Subsequently, the calibration from the initial cluster is transferred to another cluster based upon the specific differences between clusters.

[0015] The parent application to the current application, S. Malin and T. Ruchti, *An intelligent system for noninvasive blood analyte prediction*, U.S. Patent Application Ser. No. 09/359,191 (July 22, 1999), provides a detailed description of a method of classifying spectral measurements into previously defined clusters through similarities observed in absorbance spectra. The classification system of the sampled tissue volume of the subjects is herein expanded to include classification of instrumentation variations.

[0016] Referring now to Figure 1, a general method of classifying spectral measurements into previously defined clusters is represented. Briefly the steps of the method are:

1. Measurement. (11)
2. Classification (12), in which the measured spectrum is assigned membership in any one of a number of predefined clusters 13.
3. Calibration (14), in which calibration models suited to each cluster are provided.
4. Identifying outliers (15)

MEASUREMENT

[0017] In general, instrumental variations may affect spectral response by producing either signal intensity changes, bandwidth changes, wavelength changes, or combinations thereof. These instrumental variations may include:

1. wavelength shifts;
2. nonlinear wavelength shifts;
3. wavelength expansions;
4. wavelength contractions;
5. nonlinear wavelength expansions;
6. source intensity drifts;
7. blackbody profile changes;
8. bandwidth changes;
9. resolution changes;

10. baseline deviations;
11. change with time;
12. temperature effects;
13. detector response;
14. differences in optical components (e.g. long-pass filters or fiber optics);
15. variation related to mounting of references;
16. differences in the optical interface to the sample (fiber spacing);
17. linearity;
18. detector cut-off;

and many others, which will be apparent to those skilled in the art. Spectra used for classification will typically be those of commonly known standards. Standards particularly useful for classifying shifts observed in the wavelength axis include polystyrene, rare earth oxides: holmium oxide, erbium oxide or dysprosium oxide, for example; or combinations thereof. Standards such as the diffuse reflectance standards supplied by Labsphere, Inc. (North Sutton NH) may be used to classify shifts in the intensity axis. Spectra of samples may also be used for classification. Additional intensity and wavelength standards will be known to those skilled in the art.

[0018] In the specific case of near-IR noninvasive glucose determination these standards cover the near-IR spectral region. Additional spectroscopy and chromatography-spectroscopy hyphenated techniques such as AA (atomic absorption spectroscopy) or GC-MS (gas chromatography mass spectroscopy) will each require their own standards which are known by those skilled in the art. Additionally, spectra of tissue phantoms collected on an instrument may also be used to characterize that instrument. Tissue phantoms are helpful in characterizing an instrument because spectra of these phantoms simulate noninvasive spectra of living tissue. There are tissue phantoms that simulate the absorption coefficients of various skin tissues; while others simulate the scattering coefficients of the body. Some common simulants include:

1. milk and milk products
2. milk products in combination with India ink, used to adjust the absorption coefficient;
3. emulsions of fatty substances in water maintained in solution with an emulsifier such as lecithin. One such commercial product is INTRALIPID, supplied by Kabivitrum AB (Stockholm, Sweden);
4. Intra-serum and Intra-gel. See K. Hazen, J. Welch, S. Malin, T. Ruchti, A. Lorenz, T. Troy, S. Thénadil, T. Blank, *Intra-serum and intra-gel for modeling human skin tissue*, U.S. Patent Application Ser. No. 09/502,877 (February 10, 2000).

[0019] Other scattering and absorbing mediums are known to those skilled in the art. Concentrations of these tissue simulants may be adjusted to match the scattering and absorption coefficients of body tissues such as

skin or internal organs.

[0020] It will also be apparent that a set of exemplar measurements is required for cluster definition and development of calibration models, in addition to the actual spectral measurements utilized for classification.

CLASSIFICATION

[0021] New spectral measurements are passed to a pattern classification system that classifies the measurements into previously defined clusters having a high degree of internal consistency through spectral features related to instrumental variation.

The classification itself includes the following steps:

1. Feature extraction; and
2. Classification of features according to a classification model.

FEATURE EXTRACTION

[0022] The process of feature extraction is more fully described in the parent application to the current application, U.S. Patent Application Ser. No. 09/359,191, *supra*. Feature extraction is any mathematical transformation that enhances a particular aspect or quality of the data that is useful for interpretation. Features may be of two categories:

1. Abstract, and
2. Simple.

[0023] Instruments may be classified by abstract features, meaning that they may be classified using computational methods. These methods may be supervised or unsupervised. Examples include plotting primary principal components versus one another and identifying clusters of results; discriminant analysis, such as measurement of the Mahalanobis distance, and k-means clustering. Additional methods will be readily apparent to those skilled in the art.

[0024] It is important to note that the clustering techniques listed above are not mutually exclusive. Clustering of raw spectra into one or more groups may be achieved through one or more of these methods and with combinations of these approaches.

[0025] Simple features are derived from an *a priori* understanding of the system, and can be related directly to an instrument parameter or component (or parameters or components). For example, the measured bandwidth, noise characteristics, or linearity and detector cut-offs.

CLUSTER DEFINITION

[0026] As indicated above, clusters must have been previously defined, using a data set of exemplar spectral measurements. Cluster definition is the assignment of

the measurements in the exploratory data set to clusters. After cluster definition, the measurements and class assignments are used to determine the mapping from the features to cluster assignment.

[0027] Cluster definition is performed using either supervised or unsupervised methods. In the supervised case, clusters may be defined using specific knowledge of the system. For example, source intensity and detector temperatures have specific effects on the spectra, as demonstrated further below. The use of *a priori* information in this manner is the first step in supervised pattern recognition, which develops classification models when the class assignment is known.

[0028] Alternatively, clusters may be defined in an unsupervised manner using abstract features such as clustering within plots of principal component scores 'x' versus spectral loading 'y'. The result is that within a given cluster, all of the spectra have the same characteristics (interferences, instrument variations or sample issues). Clusters formed from features with physical meaning can be interpreted based on the known underlying phenomenon causing variation in the feature space.

CLASSIFICATION

[0029] Subsequent to class definition, a classifier is designed through supervised pattern recognition. A model is created, based on cluster definitions, that transforms a measured set of features to an estimated classification. The classification model is a method for determining a set of similarity measures with the predefined clusters. A decision rule assigns membership on the basis of a set of measures calculated by a decision engine.

CALIBRATION

[0030] Once the spectra have been classified into clusters, calibration models 14 are required for each cluster. Provision of the clusters is by one of two methods. In the preferred embodiment, a master calibration is developed for a first cluster, subsequently the master calibration is transferred to slave calibrations, one for each remaining cluster. An alternate embodiment, described further below, individually calculates calibrations for each cluster.

[0031] In the case of spectra collected utilizing a master and slave instrument, the term "calibration transfer," as commonly used in the art, may have different meanings. Calibration transfer may refer to transforming the slave spectral to look like the master spectra or *vice versa*. Additionally, spectra from both the master and the slave can be transferred to a common standard spectrum not present in the master or slave data sets. Furthermore, calibration transfer can refer to preprocessing steps, multivariate adjustments of the sample spectra, or adjustments to the predicted analyte concentrations

based upon standards.

[0032] Referring now to Figure 2, the process of calibration transfer is shown. Calibrations 14 are required for each of n clusters. A master calibration 21 is calculated for a first cluster; then, to provide calibrations for each of the remaining n clusters, the master calibration is transferred to slave calibrations. In this case, Calibration transfer refers to the process of transforming spectra on a slave instrument to match the characteristics of the master such that the master calibration can be applied to the slave spectra. For clusters, calibration transfer refers to the process of transforming spectra on a slave instrument to match the characteristics of the master such that a new calibration is generated that satisfies the specification of a cluster other than the one for which the master cluster was developed. The clusters may also be organized into groups of clusters, so that the master calibration is transferred to slave calibrations 23, which in turn are transferred to the slave calibration for the various clusters within each group.

[0033] Prior art methods of calibration transfer have been unsuccessful at modeling the complexity involved in providing calibration models for large numbers of instruments. Classifying the spectral measurements in to clusters having a high degree of internal consistency reduces the complexity of the problem to a level that makes it possible to apply calibration transfer to large numbers of instruments. Clustering the acquired spectra into a limited number of subgroups allows the calibration transfer issues to be broken up into subsets so that only a subset of the overall issues needs to be addressed between any two clusters. The result is that within a given cluster, all of the spectra have the same characteristics. A calibration is generated for a given cluster using spectra of samples containing the limited number of characteristics of that cluster. This eliminates having to deal with all of the variations possible in the raw measurements, significantly reducing the complexity that needs to be modeled by the calibration. Thus, the calibration utilized for a given cluster will have instrument variations included in the model that are similar to variations resulting from the analyzer employed. As each cluster has its own specification, the calibration transfer technique need only deal with the differences between those two clusters. For example, if the only difference between the clusters is a linear x-axis shift, then the calibration transfer technique need only deal with that parameter. This allows a more specific calibration transfer technique to be employed that is more robust, resulting in fewer factors, since fewer instrument variations need to be modeled. This will result in analysis of lower concentration analytes due to retention of signal to noise in the calibration transfer step.

[0034] Typically, calibration transfer techniques have to address instrument-to-instrument variations, such as wavelength shifts, nonlinear wavelength shifts, wavelength expansions (contraction), nonlinear wavelength expansions, source intensity drifts, blackbody profile

changes, bandwidth (resolution) changes, baseline deviations, change with time, temperature effects and others known to those skilled in the art.

[0035] However, in addition to instrumentation issues in calibration transfer, sample composition and sample handling considerations are very important in calibration transfer. Therefore, every sample analyzed for prediction of results should be an interpolation within the matrix space of the calibration data set; otherwise the calibration may not calculate an accurate prediction of analyte concentration. For example, if the calibration data set contains glucose samples ranging from 40 to 400 mg/dL, then predictions on samples with glucose outside of this range are suspect.

[0036] Thus, an important additional benefit of the invented classification algorithm is that outlier spectra may be identified, as shown in Figure 1. Each cluster has its own set of classification requirements. If a spectrum does not fall within the parameters of a given cluster, another cluster must be found which has parameters allowing analysis of that spectrum. If no cluster is found, then the sample is reported as an outlier rather than having a value reported for the analyte that may not be correct. In this manner, analysis on samples or instruments for which no satisfactory calibration model has been developed is prevented.

[0037] Spectra classified as outliers may assume two forms. In some instances, the outlier spectrum is close to a given cluster. In such a case, conventional calibration transfer techniques may be applied to this spectrum to transform the spectrum such that it falls within one of the clusters for which calibrations exist. In case of failure, the spectrum remains classified as an outlier, as will spectra that are characterized as gross outliers.

[0038] Classification and calibration transfer need not be limited to differences in instrumentation; it may also be applied to sample spectra for analyte prediction. For example, a cluster may be defined for healthy 18 to 30 year old men. A noninvasive glucose model may be built for this narrow demographic. A separate cluster may be for 18 to 30 year old non-pregnant women. Differences between the two clusters such as body fat and sampled volume may be addressed and a calibration transfer technique may be optimized for those differences based on relatively few spectra, thus allowing a glucose model initially generated for a narrow demographic to be gradually expanded.

[0039] The foregoing embodiment does not require calibrations to be built for every cluster, and thereby provides the important advantage of sparing the considerable time, money and effort required for building a separate calibration for every cluster. The actual calibration transfer technique employed may be selected from the many algorithms known to those skilled in the art.

[0040] An alternative embodiment classifies spectra into clusters, as in the preferred embodiment, and builds individual calibrations for each cluster. The requirement for calibration transfer is eliminated at the expense of

the requirement for more spectra. However, an additional measure of reliability is gained by having clusters with well-defined and narrow parameters. As in the previous embodiment, fewer instrument variations need to be modeled so that early factors can focus on analyte information, resulting in a more robust model.

[0041] While the foregoing description of the invention has been directed to characterizing different instruments, the invented method also finds application in classifying spectra from a single instrument according to known variations in instrument components or known environmental variations. Two examples are described.

1. The cutoff of the 2.6 μ m InGaAs detector employed in many near-IR analyzers blue shifts as the temperature of the detector decreases. Air spectra collected on a NICOLET 860, supplied by the Nicolet Instrument Corporation (Madison WI) with the InGaAs detector ranging from 0 to -20°C are presented in Figure 3A. The spectral cutoff region from 2400 to 2600 nm is expanded in Figure 3B. The cut-off decreases as the detector temperature decreases. Using reference or air spectra, a simple analysis, such as determining the wavelength at which 10% of the peak intensity is observed, allows the instrument to be classified as to whether the detector is properly cooled and to the extent that it was cooled.

2. As a tungsten halogen source increases in temperature, the blackbody emission increases in magnitude and additional light throughput is observed. For example, in diffuse reflectance spectra of a 5% Labsphere diffuse reflectance standard, the overall intensity is observed to increase as the source temperature increases, as shown in Figure 4. This effect can be used to classify the instrument in terms of the source temperature and total light throughput.

[0042] It is a relatively simple task to characterize the instrument in terms of additional components. Examples are slit width, affecting bandwidth or room temperature, affecting total light throughput. There are, however, a limited number of components to a spectrometer; it has been observed that instruments group into a relatively small number of clusters.

[0043] Once an instrument, or a given spectrum, is classified into a cluster, the appropriate calibration routine can be employed. A single analyzer may be loaded with multiple calibration routines, as the instrument may vary during its lifetime. In fact, environmental factors may cause the appropriate calibration to change with every spectrum collected.

[0044] This instrument classification approach was designed for use on noninvasive glucose analyzers where glucose is measured using diffuse reflectance spectra of skin on the human body. However, the same technology can be applied to any form of noninvasive

analysis including but not limited to: noninvasive analysis of albumin, globulin, urea, creatinine, oxygen, hemoglobin A₁C and electrolytes such as Na⁺, K⁺, Cl⁻. The technology may also be utilized in biomedical applications.

[0045] The classification approach disclosed here finds application in various fields of endeavor, the agricultural field, for example. Exemplary agricultural applications are: analysis of fat in milk, protein or moisture analysis in wheat, or analysis of sugars in fruits. The classification approach may also benefit pharmaceutical companies in the analysis of intact tablets or in the characterization of raw materials. Finally, petrochemical companies may utilize the method in classification of fuels, fuel by-products and in raw fuel materials. In general, this technique is useful wherever a large number of analyzers are utilized to quantify analytes in samples.

[0046] Although the invention is described herein with reference to the preferred embodiment, one skilled in the art will readily appreciate that other applications may be substituted for those set forth herein without departing from the scope of the present invention. Accordingly, the invention should only be limited by the claims included below.

Claims

1. A method of **characterizing** spectrometer instruments according to instrumental variation, that is between instruments and/or variation over time within the same instrument, comprising the steps of:

providing a plurality of spectra (11) of known standards from at least one spectrometer instrument; and

classifying (12) said at least one spectrometer instrument into at least one of a plurality of predefined clusters on the basis of spectral features extracted from said at least one spectrum; and

providing (14) at least one calibration model for each of said predefined clusters, wherein each calibration model compensates for instrumental variation of instruments classified to the respective cluster.

2. The method of Claim 1 wherein said instrumental variation comprises any of:

wavelength shifts;
nonlinear wavelength shifts;
wavelength expansions;
wavelength contractions;
nonlinear wavelength expansions;
source intensity drifts;
blackbody profile changes;
bandwidth changes;

resolution changes;
baseline deviations;
changes over time;
temperature effects;
detector response;
differences in optical components;
variation related to mounting of references;
differences in the optical interface to the sample;
linearity; and
detector cut-off.

3. The method of Claim 1, wherein said plurality of spectra is measured on a plurality of spectrometer instruments.

4. The method of Claim 1, wherein said plurality of spectra is measured on a single spectrometer instruments at successive time intervals.

5. The method of Claim 1, wherein said classifying step comprises steps of:

extracting features; and
classifying said features according to a classification model and decision rule.

6. The method of Claim 5, wherein said extracting step comprises any mathematical transformation that enhances a particular aspect or quality of data that is useful for interpretation.

7. The method of Claim 5, wherein said classification model comprises means for determining a set of similarity measures with predefined classes.

8. The method of Claim 5, wherein said decision rule comprises means for assigning class membership on the basis of a set of measures calculated by a decision engine.

9. The method of Claim 1, wherein individual features are divided into two categories, said categories comprising:

abstract features wherein said features are extracted using various computational methods ;
and
simple features that are derived from an *a priori* understanding of a system, wherein said feature is directly related to an instrument parameter or component.

10. The method of Claim 9, wherein said abstract features are calculated using any of:

plotting primary principal components versus one another and identifying resulting clusters;

discriminant analysis; and
k-means clustering.

11. The method of Claim 5, wherein said classifying step further comprises a step of employing factor-based methods to build a model capable of representing variation in a measured spectrum related to variations in spectral response;

wherein projection of a measured absorbance spectrum onto said model constitutes a feature that represents spectral variation related to instrument variation.

12. The method of Claim 5, wherein said classifying step further comprises steps of:

measuring the similarity of a feature to predefined clusters; and
assigning membership in a cluster.

13. The method of Claim 5, further comprising a step of:

assigning measurements in an exploratory data set to clusters.

14. The method of Claim 13, further comprising a step of:

using measurements and class assignments to determine a mapping from features to cluster assignments.

15. The method of Claim 13, further comprising steps of:

defining clusters from said features in a supervised manner, wherein each set of features is divided into two or more regions, and wherein classes are defined by combinations of feature divisions;
designing a classifier subsequent to class definition through supervised pattern recognition by determining an optimal mapping or transformation from the feature space to a class estimate which minimizes the number of misclassifications; and
creating a model based on class definitions which transforms a measured set of features to an estimated classification.

16. The method of Claim 1, further comprising the step of applying at least one of said calibration models to analysis of new sample measurements.

17. The method of Claim 16, wherein said calibration models model differences between said predefined clusters.

18. The method of Claim 16, wherein a master calibration model is developed for a first of said clusters from a set of exemplar spectra with reference values and pre-assigned classification definitions.

19. The method of Claim 18, further comprising the step of transferring said master calibration model to a plurality of slave calibration models, wherein a slave calibration model is calculated for each remaining cluster, and wherein a transform modifies said master calibration model to a slave calibration model in accordance with principal features defining each of said classes.

20. The method of Claim 19, wherein said transferring step comprises the steps of:

transferring said master calibration model to a first slave calibration model;
transferring said first slave calibration model to a second slave calibration model;
and repeating said transfer from one slave calibration model to another slave calibration model, until a calibration has been provided for each of said predefined clusters;

wherein a transform modifies said transferred calibration models in accordance with principal features defining each of said clusters.

21. The method of Claim 18, further comprising the step of transferring said master calibration model to a plurality of slave calibration models, wherein a slave calibration model is calculated for each remaining cluster, and wherein a transform modifies said slave calibration model to said master calibration model in accordance with principal features defining each of said classes.

22. The method of Claim 21, wherein said transferring step comprises the steps of:

transferring said master calibration model to a first slave calibration model;
transferring said first slave calibration model to a second slave calibration model;
and repeating said transfer from one slave calibration model to another slave calibration model, until a calibration has been provided for each of said predefined clusters;

wherein a transform modifies said transferred calibration models in accordance with principal features defining each of said clusters.

23. The method of Claim 16, wherein a different calibration model is developed for each class, and wherein said calibration models are developed from

a set of exemplar spectra, with reference values and pre-assigned cluster definitions.

24. The method of Claim 23, wherein a spectrum is assigned to one of many predefined clusters for which a calibration model has been developed.

25. The method of Claim 1, further comprising the steps of:

providing new spectral measurements;
comparing said new spectral measurements to each of said predefined clusters according to extracted spectral features;
reporting those measurements as outliers for which a matching cluster is not found.

Patentansprüche

1. Verfahren zum Charakterisieren von Spektrometer-Instrumenten gemäß der instrumentellen Variation, welche zwischen Instrumenten vorliegt, und/oder der Variation über die Zeit innerhalb des gleichen Instruments, umfassend die Schritte des:

Vorsehens einer Mehrzahl von Spektren (11) bekannter Standards von mindestens einem Spektrometer-Instrument; und
Klassifizierens (12) des mindestens einen Spektrometer-Instruments in mindestens einen von einer Mehrzahl vordefinierter Cluster auf der Basis von spektralen Merkmalen, welche aus dem mindestens einen Spektrum extrahiert wurden; und
Vorsehens (14) von mindestens einem Kalibrationsmodell für jeden der vordefinierten Cluster, wobei jedes Kalibrationsmodell die instrumentelle Variation von Instrumenten, die in den jeweiligen Cluster klassifiziert wurden, kompensiert.

2. Verfahren von Anspruch 1, worin die instrumentelle Variation irgendeines aus dem Folgenden umfasst:

Wellenlängenverschiebungen;
nichtlineare Wellenlängenverschiebungen;
Wellenlängenexpansionen;
Wellenlängenkontraktionen;
nichtlineare Wellenlängenexpansionen;
Quellenintensitäts-Driften;
Schwarzkörper-Profiländerungen;
Bandbreitenänderungen;
Auflösungsänderungen;
Grundlinien-Abweichungen;
Änderungen über die Zeit;
Temperatureffekte;
Detektorantwort;

Abweichungen in optischen Komponenten;
Variation im Zusammenhang mit der Referenzen-Montierung;
Abweichungen in der optischen Grenzfläche zu der Probe;
Linearität; und
Detektor-Cut-Off.

3. Verfahren von Anspruch 1, worin die Mehrzahl von Spektren auf einer Mehrzahl von Spektrometer-Instrumenten gemessen wird.
4. Verfahren von Anspruch 1, worin die Mehrzahl von Spektren auf einem einzelnen Spektrometer-Instrument in sukzessiven Zeitintervallen gemessen wird.
5. Verfahren von Anspruch 1, worin der Klassifizierungsschritt die folgenden Schritte umfasst:

Extrahieren von Merkmalen; und
Klassifizieren der Merkmale gemäß eines Klassifikationsmodells und einer Entscheidungsregel.

6. Verfahren von Anspruch 5, wobei der Extraktionsschritt jedwede mathematische Transformation umfasst, welche einen besonderen Aspekt oder eine besondere Qualität von Daten verstärkt, welche(r) für die Interpretation nützlich ist.
7. Verfahren von Anspruch 5, wobei das Klassifikationsmodell Mittel zum Bestimmen eines Satzes von Ähnlichkeitsmaßen mit vordefinierten Klassen umfasst.
8. Verfahren von Anspruch 5, wobei die Entscheidungsregel Mittel zum Zuweisen von Klassenmitgliedschaft auf der Basis eines Satzes von Maßen umfasst, welche durch eine Entscheidungsmaschine berechnet werden.
9. Verfahren von Anspruch 1, worin individuelle Merkmale in zwei Kategorien eingeteilt werden, wobei die Kategorien folgendes umfassen:

abstrakte Merkmale, wobei die Merkmale unter Anwendung verschiedener Berechnungsverfahren extrahiert werden; und
einfache Merkmale, welche aus einem *a priori* Verständnis eines Systems abgeleitet werden, wobei das Merkmal direkt mit einem/einer Instrumentparameter oder -komponente zusammenhängt.

10. Verfahren von Anspruch 9, worin die abstrakten Merkmale berechnet werden unter Verwendung von irgendeinem unter:

Auftragen der primären Hauptkomponenten gegeneinander und Identifizieren resultierender Cluster;
Diskrimanten-Analyse; und
k-Mittelwert-Clustering.

11. Verfahren von Anspruch 5, worin der Klassifizierungsschritt ferner einen Schritt der Anwendung von Faktor-basierenden Methoden umfasst, um ein Modell aufzubauen, das zum Repräsentieren von Variation in einem gemessenen Spektrum, im Zusammenhang mit Variationen in der spektralen Antwort, fähig ist;
wobei Projektion eines gemessenen Absorptionsspektrums auf das Modell ein Merkmal darstellt, das spektrale Variation repräsentiert, welche mit Instrumentenvariation zusammenhängt.

12. Verfahren von Anspruch 5, worin der Klassifizierungsschritt ferner folgende Schritte umfasst:

Messen der Ähnlichkeit eines Merkmals zu vordefinierten Clustern; und
Zuweisen von Mitgliedschaft in einem Cluster.

13. Verfahren von Anspruch 5, ferner umfassend einen Schritt des:

Zuweisens von Messungen in einem explorativen Datensatz zu Clustern.

14. Verfahren von Anspruch 13, ferner umfassend einen Schritt des:

Verwendens von Messungen und Klassenzuweisungen, um eine Kartierung von Merkmalen aus zu Cluster-Zuweisungen zu bestimmen.

15. Verfahren von Anspruch 13, ferner umfassend die Schritte des:

Definierens von Clustern aus den Merkmalen in einer überwachten Weise, wobei jeder Satz von Merkmalen in zwei oder mehr Regionen eingeteilt wird, und wobei Klassen durch Kombinationen von Merkmalseinteilungen definiert sind;
Entwerfens eines Klassifikators anschließend an Klassendefinition durch überwachte Mustererkennung mittels Bestimmung einer optimalen Kartierung oder Transformation aus dem Merkmalsraum zu einer Klasseneinschätzung, welche die Anzahl von Fehlklassifikationen minimiert; und
Erzeugens eines Modells auf Basis von Klassendefinitionen, welches einen gemessenen Satz von Merkmalen zu einer geschätzten Klassifikation transformiert.

16. Verfahren von Anspruch 1, ferner umfassend den Schritt des Anwendens von mindestens einem der Kalibrationsmodelle auf die Analyse von neuen Proberunessungen.

17. Verfahren von Anspruch 16, worin die Kalibrationsmodelle Abweichungen zwischen den vordefinierten Clustern modellhaft angeben.

18. Verfahren von Anspruch 16, worin ein Master-Kalibrationsmodell für einen ersten der Cluster aus einem Satz von exemplarischen Spektren, mit Referenzwerten und vorab-zugewiesenen Klassifikations-Definitionen, entwickelt wird.

19. Verfahren von Anspruch 18, ferner umfassend den Schritt des Transferierens des Master-Kalibrationsmodells zu einer Mehrzahl von Slave- bzw. Tochter-Kalibrationsmodellen, wobei ein Slave-Kalibrationsmodell für jeden verbleibenden Cluster berechnet wird, und wobei eine Transformation das Master-Kalibrationsmodell zu einem Slave-Kalibrationsmodell gemäß Haupt-Merkmalen, welche jede der Klassen definieren, modifiziert.

20. Verfahren von Anspruch 19, worin der Transferierungsschritt die folgenden Schritte umfasst:

Transferieren des Master-Kalibrationsmodells zu einem ersten Slave-Kalibrationsmodell;
Transferieren des ersten Slave-Kalibrationsmodells zu einem zweiten Slave-Kalibrationsmodell;
und Wiederholen des Transfers aus einem Slave-Kalibrationsmodell zu einem anderen Slave-Kalibrationsmodell, bis eine Kalibration für jeden der vordefinierten Cluster vorgesehen worden ist;

wobei eine Transformation die transferierten Kalibrationsmodelle gemäß Haupt-Merkmalen, welche jeden der Cluster definieren, modifiziert.

21. Verfahren von Anspruch 18, ferner umfassend den Schritt des Transferierens des Master-Kalibrationsmodells zu einer Mehrzahl von Slave-Kalibrationsmodellen, wobei ein Slave-Kalibrationsmodell für jeden verbleibenden Cluster berechnet wird, und wobei eine Transformation das Slave-Kalibrationsmodell zu dem Master-Kalibrationsmodell gemäß Haupt-Merkmalen, welche jede der Klassen definieren, modifiziert.

22. Verfahren von Anspruch 21, worin der Transferierungsschritt die folgenden Schritte umfasst:

Transferieren des Master-Kalibrationsmodells zu einem ersten Slave-Kalibrationsmodell;

Transferieren des ersten Slave-Kalibrationsmodells zu einem zweiten Slave-Kalibrationsmodell;
und Wiederholen des Transfers von einem Slave-Kalibrationsmodell zu einem anderen Slave-Kalibrationsmodell, bis eine Kalibration für jeden der vordefinierten Cluster vorgesehen worden ist;

wobei eine Transformation die transferierten Kalibrationsmodelle gemäß Haupt-Merkmalen, welche jeden der Cluster definieren, modifiziert.

23. Verfahren von Anspruch 16, worin ein unterschiedliches Kalibrationsmodell für jede Klasse entwickelt wird, und worin die Kalibrationsmodelle aus einem Satz von exemplarischen Spektren, mit Referenzwerten und vorab-zugewiesenen Cluster-Definitionen, entwickelt werden.

24. Verfahren von Anspruch 23, worin ein Spektrum zu einem von vielen vordefinierten Clustern zugewiesen wird, für welche ein Kalibrationsmodell entwickelt worden ist.

25. Verfahren von Anspruch 1, ferner umfassend die Schritte des:

Vorsehens von neuen spektralen Messungen;
Vergleichens der neuen spektralen Messungen mit jedem der vordefinierten Cluster gemäß extrahierten spektralen Merkmalen;
Aufführens derjenigen Messungen als Ausreißer, für welche ein passender Cluster nicht gefunden wird.

Revendications

1. Procédé de caractérisation d'instruments de spectromètre selon une variation instrumentale, c'est-à-dire entre les instruments et/ou la variation dans le temps dans le même instrument, comprenant les étapes consistant à :

fournir une pluralité de spectres (11) de normes connues à partir d'au moins un instrument de spectromètre ; et
classifier (12) ledit au moins un instrument de spectromètre dans au moins un d'une pluralité de groupes prédéfinis sur la base de caractéristiques spectrales extraites dudit au moins un spectre ; et
fournir (14) au moins un modèle d'étalonnage pour chacun desdits groupes prédéfinis,

dans lequel chaque modèle d'étalonnage compense la variation instrumentale des instruments clas-

sifiés dans le groupe respectif.

2. Procédé selon la revendication 1 dans lequel ladite variation instrumentale comprend n'importe lequel des :

décalages de longueur d'onde ;
décalages de longueur d'onde non linéaires ;
expansions de longueur d'onde ;
contractions de longueur d'onde ;
expansions de longueur d'onde non linéaires ;
dérives d'intensité de source ;
changements de profil de corps noir ;
changements de largeur de bande ;
changements de résolution ;
déviations de la ligne de base ;
changements dans le temps ;
effets de la température ;
réaction d'un capteur ;
différences dans les composants optiques ;
variation liée au montage de références ;
différences dans l'interface optique de l'échantillon.
Linéarité, et
Arrêt du capteur.

3. Procédé selon la revendication 1, dans lequel ladite pluralité de spectres est mesurée par une pluralité d'instruments de spectromètres.

4. Procédé selon la revendication 1, dans lequel ladite pluralité de spectres est mesurée par des instruments à spectromètre unique à des intervalles de temps successifs.

5. Procédé selon la revendication 1, dans lequel ladite étape de classification comprend les étapes consistant à :

extraire les caractéristiques ; et
classifier lesdites caractéristiques selon un modèle de classification et une règle de décision.

6. Procédé selon la revendication 5, dans lequel ladite étape d'extraction comprend toute transformation mathématique qui améliore un aspect particulier ou une qualité particulière des données qui sont utiles pour l'interprétation.

7. Procédé selon la revendication 5, dans lequel ledit modèle de classification comprend des moyens pour déterminer un ensemble de mesures de similarité avec des classes prédéfinies.

8. Procédé selon la revendication 5, dans lequel ladite règle de décision comprend un moyen pour attribuer une appartenance à une classe sur la base d'un ensemble de mesures calculées par un moteur

de décision.

9. Procédé selon la revendication 1, dans lequel les caractéristiques individuelles sont divisées en deux catégories, lesdites catégories comprenant :
- des caractéristiques abstraites desquelles lesdites caractéristiques sont extraites en utilisant divers procédés de calcul ; et
- des caractéristiques simples qui découlent d'une compréhension a priori d'un système,
- dans lequel ladite caractéristique est directement liée à un paramètre ou à un composant d'un instrument.
10. Procédé selon la revendication 9, dans lequel lesdites caractéristiques abstraites sont calculées en utilisant l'un de :
- report des composants primaires principaux les uns contre les autres et identification des groupes résultants ;
- analyse discriminante ; et
- classification par des moyens k.
11. Procédé selon la revendication 5, dans lequel ladite étape de classification comprend en outre une étape d'emploi de procédés par facteurs pour construire un modèle capable de représenter la variation dans un spectre mesuré selon les variations de la réponse spectrale ;
- dans lequel la projection d'un spectre d'absorbance mesuré sur ledit modèle constitue une caractéristique qui représente une variation spectrale liée à la variation d'instrument.
12. Procédé selon la revendication 5, dans lequel ladite étape de classification comprend en outre les étapes consistant à :
- mesurer la similarité d'une caractéristique à des groupes prédéfinis ; et
- attribuer une appartenance à un groupe.
13. Procédé selon la revendication 5, comprenant en outre une étape consistant à :
- attribuer des mesures dans un ensemble de données exploratoires à des groupes.
14. Procédé selon la revendication 13, comprenant en outre une étape consistant à :
- utiliser des mesures et des attributions de classe pour déterminer une carte des caractéristiques aux attributions de groupe.

15. Procédé selon la revendication 13, comprenant en outre des étapes consistant à :

définir des groupes à partir desdites caractéristiques de manière supervisée, dans lesquels chaque ensemble de caractéristiques est divisé en deux régions ou plus, et dans lesquels les classes sont définies par des combinaisons de divisions de caractéristiques ;

concevoir un classificateur après la définition de classe via une reconnaissance de modèle supervisée en déterminant une cartographie ou une transformation optimale de l'espace de la caractéristique en une estimation de classe qui minimise le nombre de d'erreurs de classifications ; et

créer un modèle basé sur les définitions de classe qui transforme un ensemble de caractéristiques mesurées en une classification estimée.

16. Procédé selon la revendication 1, comprenant en outre l'étape consistant à appliquer au moins un desdits modèles d'étalonnage à l'analyse des mesures d'un nouvel échantillon.

17. Procédé selon la revendication 16, dans lequel lesdits modèles d'étalonnage modélisent des différences entre lesdits groupes prédéfinis.

18. Procédé selon la revendication 16, dans lequel un modèle d'étalonnage maître est développé pour un premier desdits groupes à partir d'un ensemble de spectres exemplaires avec des valeurs de référence et des définitions de classification pré-attribuées.

19. Procédé selon la revendication 18, comprenant en outre l'étape consistant à transférer ledit modèle d'étalonnage maître à une pluralité de modèles d'étalonnage esclaves, dans lesquels un modèle d'étalonnage esclave est calculé pour chaque groupe restant, et dans lesquels une transformée modifie ledit modèle d'étalonnage maître en un modèle d'étalonnage esclave selon les principales caractéristiques définissant chacune desdites classes.

20. Procédé selon la revendication 19, dans lequel ladite étape de transfert comprend les étapes consistant à :

transférer ledit modèle d'étalonnage maître à un premier modèle d'étalonnage esclave ;

transférer ledit modèle d'étalonnage esclave à un deuxième modèle d'étalonnage esclave ;

et répéter ledit transfert d'un modèle d'étalonnage esclave à un autre modèle d'étalonnage esclave, jusqu'à ce qu'un étalonnage ait été fourni pour chacun desdits groupes prédéfinis ;

dans lequel une transformée modifie lesdits modèles d'étalonnage transférés selon les principales caractéristiques définissant chacun desdits groupes.

21. Procédé selon la revendication 18, comprenant en outre l'étape consistant à transférer ledit modèle d'étalonnage maître à une pluralité de modèles d'étalonnage esclaves, dans lequel un modèle d'étalonnage esclave est calculé pour chaque groupe restant, et dans lequel une transformée modifie ledit modèle d'étalonnage esclave en ledit modèle d'étalonnage maître selon les principales caractéristiques définissant chacune desdites classes.

22. Procédé selon la revendication 21, dans lequel ladite étape de transfert comprend les étapes consistant à :

transférer ledit modèle d'étalonnage maître à un premier modèle d'étalonnage esclave ;
transférer ledit premier modèle d'étalonnage esclave à un deuxième modèle d'étalonnage esclave ;
et répéter ledit transfert d'un modèle d'étalonnage esclave à un autre modèle d'étalonnage esclave, jusqu'à ce qu'un étalonnage ait été fourni pour chacun desdits groupes prédéfinis ;

dans lequel une transformée modifie lesdits modèles d'étalonnage transférés selon les principales caractéristiques définissant chacun desdits groupes.

23. Procédé selon la revendication 16, dans lequel un modèle d'étalonnage différent est développé pour chaque classe, et dans lequel lesdits modèles d'étalonnage sont développés à partir d'un ensemble de spectres exemplaires, avec des valeurs de référence et des définitions de groupes pré-attribuées.

24. Procédé selon la revendication 23, dans lequel un spectre est attribué à l'un des nombreux groupes prédéfinis pour lesquels un modèle d'étalonnage a été développé.

25. Procédé selon la revendication 1, comprenant en outre les étapes consistant à :

fournir de nouvelles mesures spectrales ;
comparer lesdites nouvelles mesures spectrales à chacun desdits groupes prédéfinis selon les caractéristiques spectrales extraites ;
reporter ces mesures comme observations aberrantes pour lesquels un groupe correspondant ne peut être trouvé.

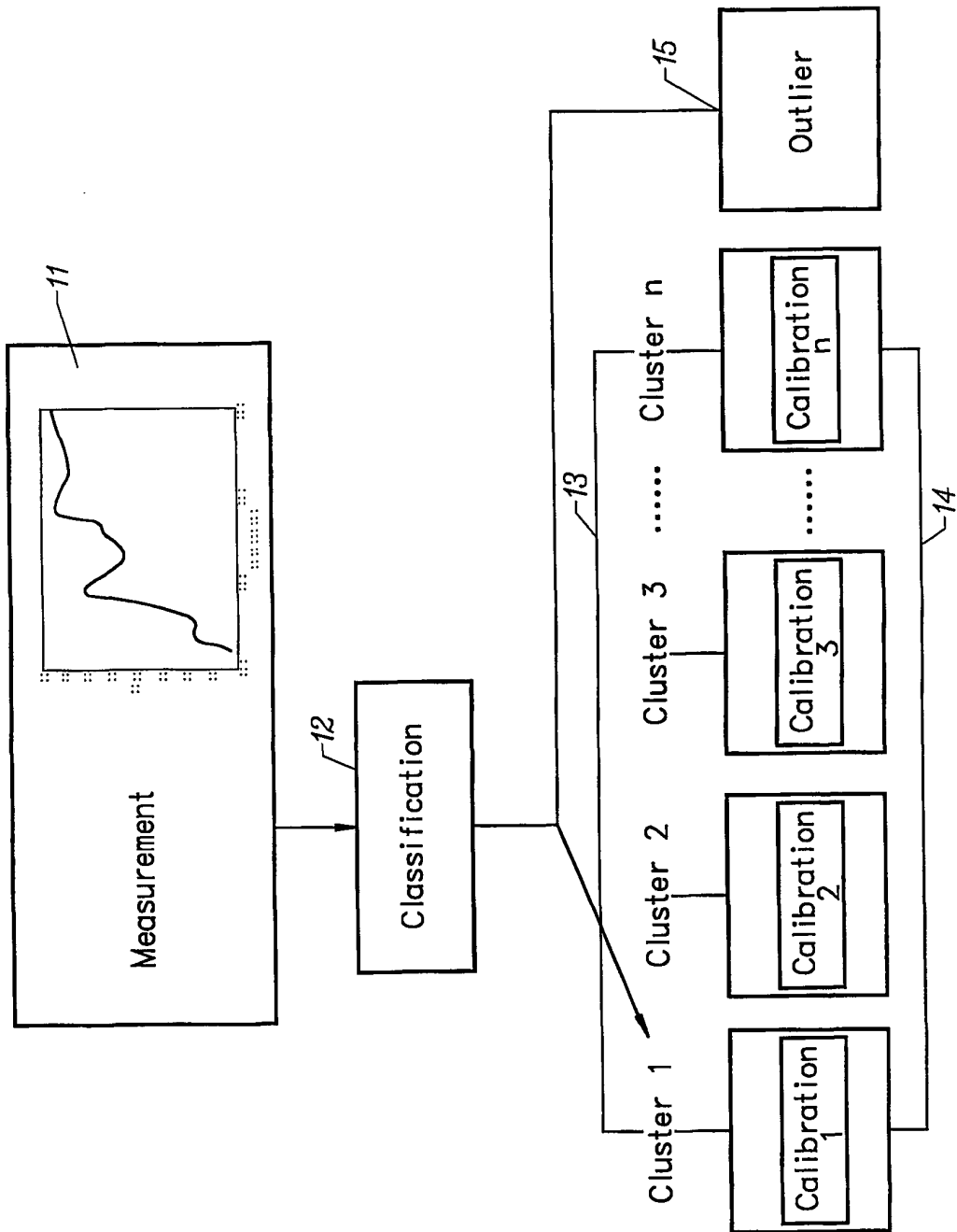


FIG. 1

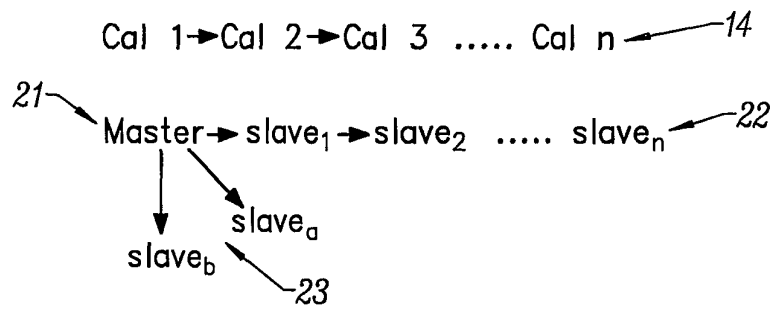


FIG. 2

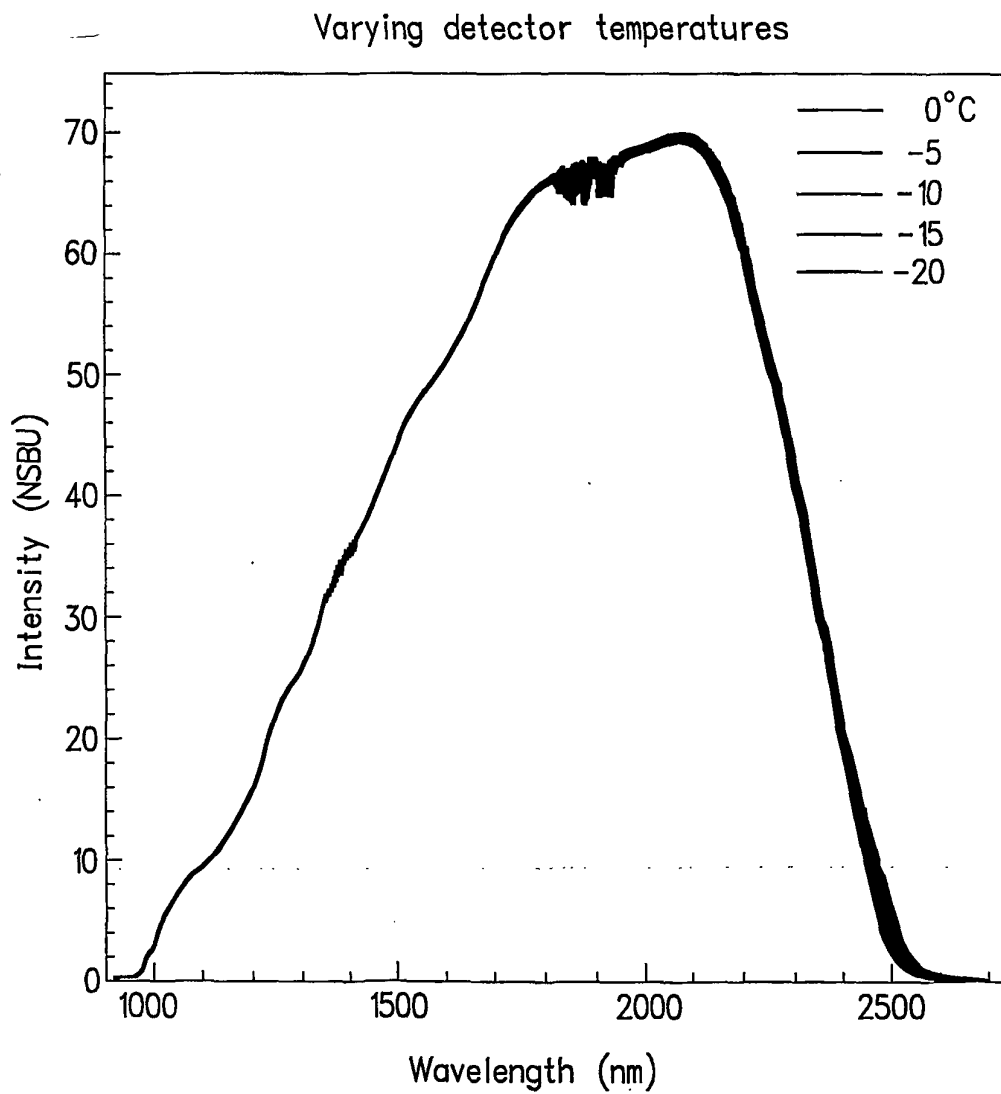
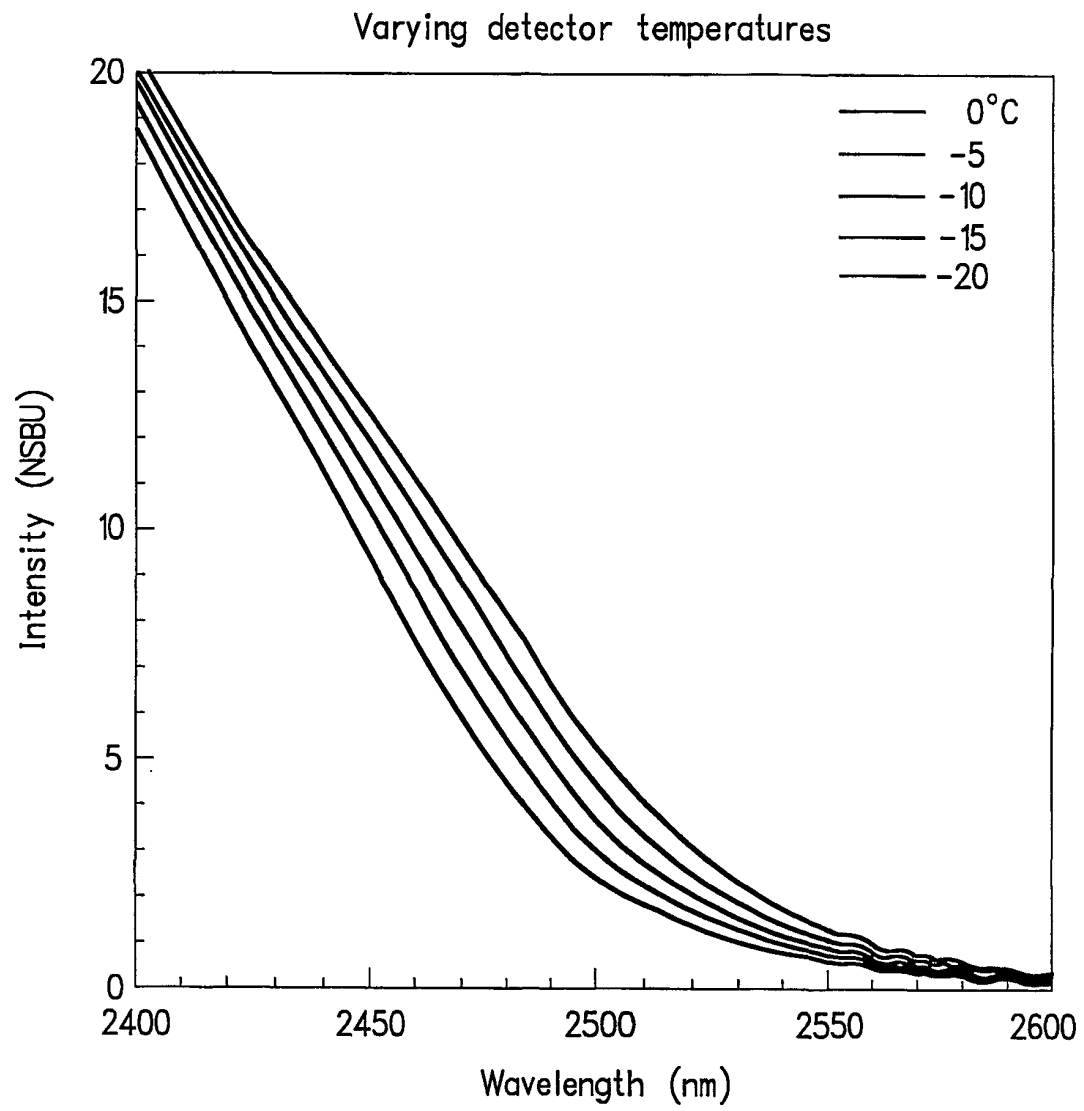


FIG. 3A

*FIG. 3B*

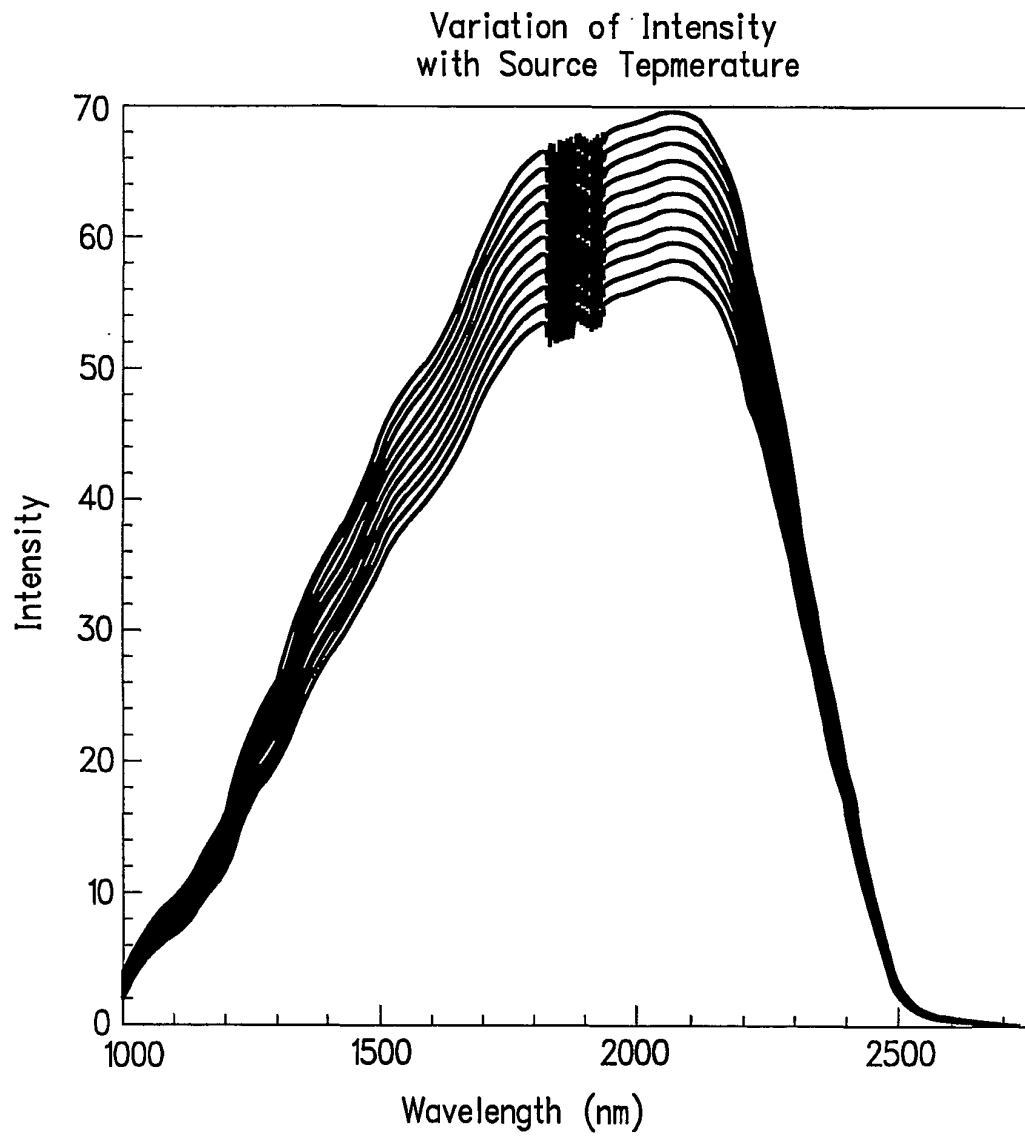


FIG. 4

专利名称(译)	表征光谱仪的方法		
公开(公告)号	EP1319176B1	公开(公告)日	2005-10-12
申请号	EP2001962218	申请日	2001-08-16
[标]申请(专利权)人(译)	INSTR度量		
申请(专利权)人(译)	仪器指标INC.		
当前申请(专利权)人(译)	SENSYS MEDICAL , INC.		
[标]发明人	HAZEN KEVIN H BLANK THOMAS B MONFRE STEPHEN L RUCHTI TIMOTHY L		
发明人	HAZEN, KEVIN, H. BLANK, THOMAS, B. MONFRE, STEPHEN, L. RUCHTI, TIMOTHY, L.		
IPC分类号	G01N21/35 A61B5/00 A61B5/107 A61B5/145 A61B5/1455 A61B5/1495 G01J3/02 G01J3/28 G01N21/27 G01N21/47 G01N21/49 G06F19/00		
CPC分类号	A61B5/0075 A61B5/1075 A61B5/14532 A61B5/1455 A61B5/1495 A61B5/7264 A61B2560/0223 A61B2560/0233 G01J3/28 G01J2003/2866 G01N21/274 G01N21/3563 G01N21/359 G01N21/4785 G01N21/49 G16H50/20		
代理机构(译)	利明，约翰·杰勒德		
优先权	09/664973 2000-09-18 US		
其他公开文献	EP1319176A2		
外部链接	Espacenet		

摘要(译)

光谱仪仪器的特征在于将它们的光谱分类为先前定义的簇。将光谱映射到聚类，并且基于提取的光谱特征与先前定义的聚类之一的相似性来进行分类（12）。提供每个群集的校准模型以补偿仪器变化。通过将主校准（21）传送到从属校准（23）或通过为每个簇计算单独的校准来提供校准模型。校准传递的简化方法将聚类彼此映射，使得在聚类之间传递的校准仅模拟两个聚类之间的差异，从而大大降低了模型的复杂性。

