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A simple spectrogram model for high-accuracy spectral calibration of VIPA spectrometers

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The virtually imaged phased array (VIPA) spectrometer uses the orthogonal dispersion method and has the advantages of compact structure, high spectral resolution, and wide wavelength coverage. It has been widely used in different fields. However, due to the non-linear dispersion of the VIPA etalon and the cross-dispersion structure of the VIPA spectrometer, simple and high-accuracy wavelength calibration remains a challenge. In this paper, a new and simple five-parameter spectrogram model is developed by simplifying the phase-matching equation of the VIPA etalon and considering the angle between the camera and dispersion direction, which can achieve a frequency accuracy better than one pixel. The performance of the model is demonstrated by measuring the CO₂ absorption spectrum in the range of 1.42 to 1.45 μm using a self-designed near-infrared VIPA spectrometer ($\mathcal{R} \sim 3 \times 10^5$, 4.5 pm). The reported method is simple and easy to solve with high accuracy, which is conducive to promoting the application of VIPA spectrometers in precision measurement.

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Introduction

Ultra-high resolution ($\mathcal{R} \geq 10^5$) spectroscopy has important applications in many fields, such as multispecies trace gas detection,^{1,2} molecular fingerprinting,³ precision measurement,⁴ spectral-domain optical coherence tomography,^{5,6} astronomical observation,⁷ and basic physics and chemistry.^{8,9} Over more than a decade, a new ultra-high-resolution spectrometer based on the virtually imaged phased array (VIPA) etalon has become an exciting topic in the field of broadband spectroscopy research.³ The VIPA spectrometer uses the orthogonal dispersion structure composed of the VIPA etalon and grating to achieve two-dimensional light dispersion, and uses an area-array camera to collect the light, achieving high-resolution ($\mathcal{R} = 8 \times 10^4 - 9 \times 10^5$) broadband spectral measurements (from about 9 nm in the visible band to 90 nm in the mid-infrared band in a single frame);^{3,7,8} using a high-speed camera, microsecond time-resolved (or even faster) broadband spectral measurements can be achieved,^{8,9} which

has great application potential. However, due to the non-linear dispersion of the VIPA etalon and the cross-dispersion structure of the spectrometer, high accuracy wavelength calibration of VIPA spectrometers using a simple calibration setup remains a challenge.¹⁰

Currently, wavelength calibration of the VIPA spectrometer is usually done by two methods: one is using a high-precision optical frequency reference (HPOFR), such as a tunable laser,¹¹ a frequency comb,^{3,12,13} an F-P etalon,^{5,6} or an optical spectrum analyzer (OSA)^{10,14} to calibrate the frequency. For example, Diddams *et al.* used a single-longitudinal mode laser to provide an absolute frequency reference;³ Zhu *et al.* used a frequency comb to provide a relative frequency reference, achieving an extremely high accuracy of about 0.03 pixels.¹⁵ The other method is using the absorption spectrum of molecules or the absorption or emission spectrum of atoms to fit the parameters of the dispersion model, such as the orthogonal dispersive model,¹⁴ proportional model¹⁶ and linear model.¹⁶ The dispersion model can establish the relation between the pixel coordinates and the wavelengths of the entire spectral image. For example, Thrope *et al.* used the gas absorption data (CO₂, H₂O, CH₄ and NH₃) and the orthogonal dispersive model¹⁴ of the VIPA spectrometer to obtain a high accuracy mapping of pixel to wavelength in the range of 1.5 to 1.7 μm .¹ Klose *et al.* used a third-order polynomial fit of the experimental peak position to the theoretical peak from the HITRAN database to achieve wavelength cali-

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bration.¹⁷ These kinds of methods are easy to integrate into the detection system and realize real-time calibration, and the setup is simpler than that of the HPOFR method. However, due to the deviation between the ideal model and the actual structure, the accurate solution of the parameters is challenging.

In this work, we report the development of a new spectrogram model to achieve high accuracy wavelength calibration. By replacing the optical path difference (OPD) in the phase-matching equation of the VIPA etalon¹⁰ with a low-order polynomial function, the number of parameters of the model is reduced and the conversion between pixel coordinates and spatial positions is elided, making the solving of parameters easier. By adding the angle between the dispersion direction and the camera, the model is more applicable. Similar approaches have been used in some HPOFR applications^{16,18} and in the calibration of an échelle spectrometer,^{19–21} where sub-pixel calibration accuracy can be achieved. Here, a more general approach than the orthogonal dispersive model has been developed, which enables high accuracy calibration in absorption applications without any additional calibration setup like a single-mode laser, an F-P etalon or others. The performance of the technique is demonstrated on a self-designed near-infrared VIPA spectrometer ($\mathcal{R} \sim 3 \times 10^5$, 4.5 pm) with a CO₂ absorption range of 1.42–1.45 μm . By comparing the theoretical coordinates calculated using the spectrogram model and the centroid coordinates of the measured image, an average accuracy of better than 1 pixel is achieved.

Principle of VIPA spectrometers

The traditional dispersion model

In an ideal VIPA-grating spectrometer, the traditional dispersion model (spatial (x, y) and wavelength dependence of the intensity distribution) can be expressed as the product of the grating and the VIPA etalon distributions:^{6,14}

$$I_{\text{out}}(x, y, \lambda) \propto I_{\text{in}}(\lambda) \times \exp\left(-2\frac{f_c^2 y^2}{f^2 W^2}\right) \times \left((1 - r_1 r_2)^2 + 4r_1 r_2 \sin^2\left(\frac{\pi \text{OPD}}{\lambda}\right)\right)^{-1} \times \exp\left(-\left(\frac{x - \lambda f(\lambda_0 - \lambda)}{\frac{d\lambda_0 \cos(\theta_d)}{\cos(\theta_{\text{ig}})f\lambda}}\right)^2\right) \quad (1)$$

where the first product term $I_{\text{in}}(\lambda)$ is the input beam power spectral density, the second term $\exp(-2f_c^2 y^2 / f^2 W^2)$ is the envelope of the light intensity along the y direction, the third term represents the VIPA dispersion law, and the fourth term describes the grating dispersion. f_c and f are the focal lengths of the cylindrical lens and the imaging lens, respectively. r_1 and r_2 are the reflection coefficients of the front and back surfaces of the VIPA etalon, respectively. d , θ_{ig} and θ_d are the grating constant, incident angle and diffraction angle, respectively.

λ_0 and W are the central wavelength and the waist radius of the collimated beam, respectively. $\text{OPD} = m\lambda$ is the optical path difference between adjacent beams of the VIPA etalon.

For different VIPA interference orders (m), the phase-matching equation in the vertical direction can be written as follows:¹⁴

$$m\lambda = 2t \cos(\theta_{\text{in}}) - \frac{2t \tan(\theta_{\text{in}}) \cos(\theta_i) y}{f} - \frac{t \cos(\theta_{\text{in}}) y^2}{n_e f^2} \quad (2)$$

where n_e , t , θ_i and θ_{in} are the refractive index, thickness, incident angle and refraction angle of the VIPA etalon, respectively, and the relationship between n_e , θ_i and θ_{in} is $\theta_{\text{in}} \approx \theta_i / n_e$.

In the horizontal direction, the dispersion is caused only by the grating, and the phase-matching equation can be expressed as follows:^{19–21}

$$m_{\text{grating}} \lambda = d(\sin \theta_{\text{ig}} \pm \sin \theta_d) \quad (3)$$

$$x = \frac{m_{\text{grating}} f (\lambda - \lambda_0)}{d \cos(\theta_d)} \quad (4)$$

where m_{grating} is the diffraction order, which is +1 in most cases in VIPA spectrometers.

The spectrogram model

For the spectrogram model, there is only one spatial position variable (y) in the OPD expression in the VIPA dispersion direction. The phase-matching equation for the VIPA etalon can be modified as follows:

$$\lambda(Y_p) = \frac{\text{OPD}}{m} = a_0 + a_1 Y_p + a_2 Y_p^2 \quad (5)$$

where Y_p is the vertical coordinate of the pixel, m is the VIPA dispersion order at wavelength λ_{Y_p} , and a_0 , a_1 , and a_2 are the coefficients of the second-order polynomial.

In the grating dispersion direction, each fringe represents a VIPA interference order within one VIPA FSR. Therefore, the relationship between the interference order m and a series of coordinates X_p on the fringe can be determined by the fringe recognition algorithm,²² as shown in the following equation:

$$\lambda(X_p) = \frac{(b_0 + b_1 X_p)}{m_{\text{grating}}} \quad (6)$$

where X_p is the horizontal coordinate and b_0 and b_1 are the one-order polynomial coefficients.

Fig. 1(a) shows an ideal schematic spectrogram of a broadband light source and four beams of single longitudinal mode light (wavelengths are λ_0 to λ_3). For a light beam of a specified frequency, such as λ_2 , there are multiple light spots (red, 3 in this figure) along the VIPA dispersion direction, which are distributed on different fringes, representing different interference orders of the VIPA etalon. For two beams with the same dispersion angle in the y direction, such as λ_2 and λ_3 , the light spots are separated into different fringes by the grating.

The above are all ideal results. In actual spectrometers, even if the installation error is quite small, special optical design and aberration^{19,23} will affect the relationship between



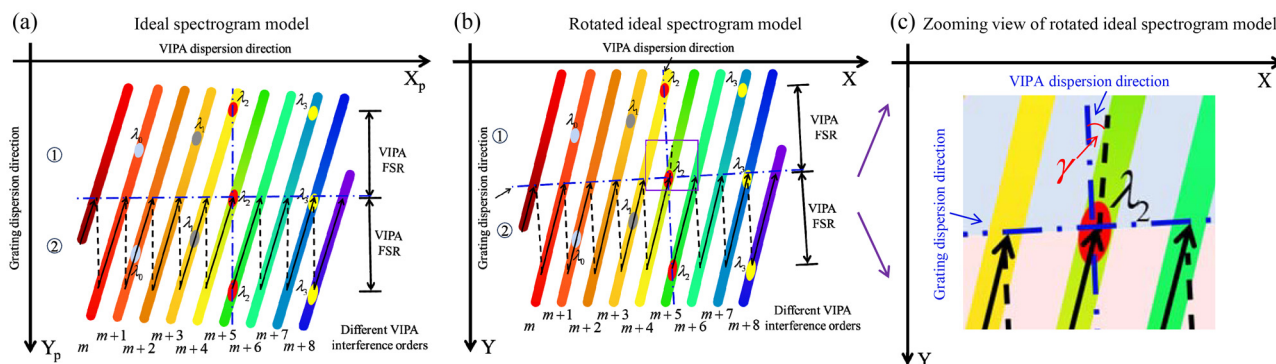


Fig. 1 The schematic spectrogram of a broadband light source and four beams of single longitudinal mode light (wavelengths are λ_0 to λ_3). (a) Ideal spectrogram model; (b) rotated spectrogram model. (c) The zoomed-in view of the rotated spectrogram model. The ellipses represent the spots of different single-mode light. In the vertical direction, different wavelengths (in different colours) are separated by the VIPA etalon. Different positions of the same wavelength represent different VIPA interference orders. In order to obtain a wider spectral coverage, a grating is used to separate the VIPA overlapping orders to form a fringe pattern. The region of one VIPA FSR can be roughly determined by a characteristic wavelength (such as λ_2). Then the fringes are connected end-to-end in the direction of the black arrows to obtain a one-dimensional spectrum. γ represents the angle between the VIPA dispersion direction and the y -axis of the camera.

the wavelength and pixel coordinates, resulting in a certain gap between the spectrum image collected by the spectrometer camera and the ideal image. For example, Diddams *et al.*³ and Zhou *et al.*²⁴ reported the camera rotation method, which rotated the camera that caused the coordinate axis to differ in the dispersion direction (as shown in Fig. 1(b)). To solve this problem, the model can be updated by simply adding the parameter γ (the angle between the VIPA dispersion direction and the camera y -axis).

Among the influence of aberrations, according to the third-order aberration theory, only distortion aberration²⁵ will affect the shape of the image, thereby affecting the interference position of a certain wavelength in the orthogonal dispersion system, causing bending of fringes, while other aberrations such as spherical aberration and coma aberration only affect the spectral resolution and fringe width of the spectrometer. In the cases of a large distortion effect,¹³ the spectral image can be corrected by using techniques such as the distortion correction algorithm.²⁷ Due to the small size of the VIPA etalon (which is comparable to the size of the F-P etalon) and the grating, the field of view of the imaging system of the VIPA spectrometer is small, and its distortion aberration can generally be ignored.²⁶ Therefore, in practical applications, the spectrogram model can be used for the wavelength calibration of the VIPA spectrometer.

Spectral calibration algorithm

The process of the spectral calibration algorithm based on the spectrogram model is shown in Fig. 2. To achieve a high-accuracy calibration, multiple wavelengths and their corresponding pixel coordinates of different VIPA interference orders are required. Here, we use Nugent-Glandorf's method²⁴ to obtain the wavelengths of these absorption spots. Then the parameter γ is determined from these inputs. In the ideal spectrogram, the x -coordinate X_p is determined only by the grating dis-

persion, and the y -coordinate Y_p is determined only by the VIPA dispersion. Thus, γ can be determined by the following formula:

$$\gamma = \arg \min \sum_i |X_i - X_{pi}|^2 \quad (7)$$

where i is the number of absorption spots, (X_i, Y_i) is the pixel coordinate of a collected image and (X_{pi}, Y_{pi}) is the pixel coordinate of the corresponding ideal image. The conversion equation between the pixel coordinates (X_i, Y_i) and (X_{pi}, Y_{pi}) is as follows:²⁸

$$\begin{pmatrix} X_p \\ Y_p \\ 1 \end{pmatrix} = \begin{pmatrix} \cos \gamma & \sin \gamma & T_x \cos \gamma + T_y \sin \gamma - T_x \\ -\sin \gamma & \cos \gamma & -T_x \sin \gamma + T_y \cos \gamma - T_y \\ 0 & 0 & 1 \end{pmatrix} \begin{pmatrix} X \\ Y \\ 1 \end{pmatrix} \quad (8)$$

where (T_x, T_y) is the coordinate of the rotation center.

After determining the angle γ , the next step is to establish an ideal model input. As shown in Fig. 1, assuming that the VIPA interference order of one of the fringes is m , and the adjacent orders are $m \pm 1$. The matrix (X, Y, m, λ) of the spectral image and the corresponding matrix (X_p, Y_p, m, λ) of the spectrogram model can be determined. With the coordinates (Y_p, m, λ) as the ideal model input, the VIPA interference order m and the second-order polynomial parameters a_0, a_1 and a_2 can be derived using the least-squares fitting method, as shown in the following formula:

$$(m, a_0, a_1, a_2) = \arg \min_{m>0} |\lambda - \lambda(Y_p)|^2 \quad (9)$$

The (m, Y_p, λ) relationship of other pixels on the fringes can be calculated using eqn (5). The model in the horizontal direction is similar to that in the vertical direction. In particular, for broadband input light, the x -coordinate of each central pixel on the fringes (X_m, m) can be directly obtained by the



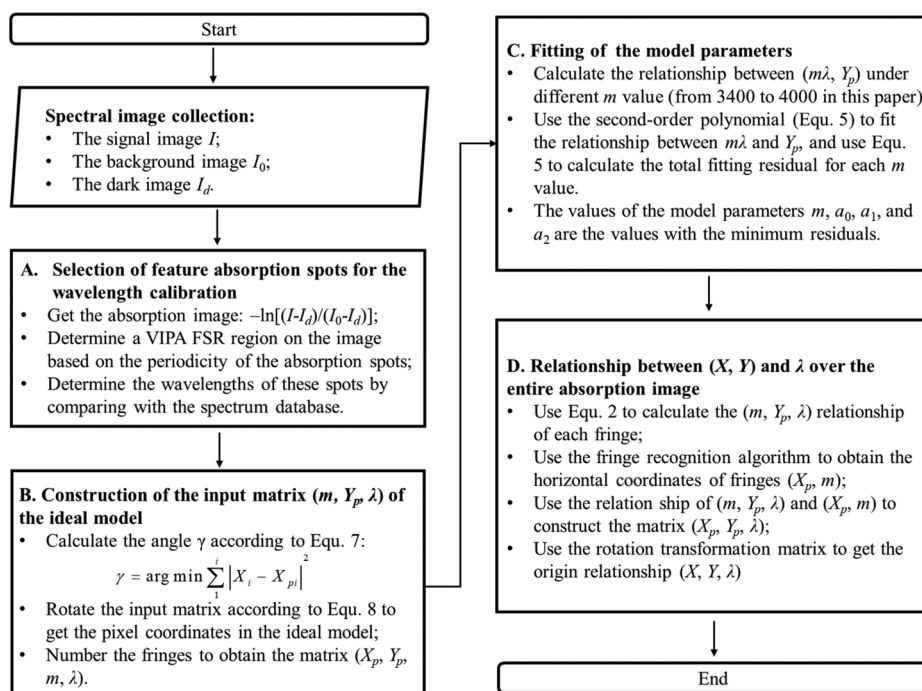


Fig. 2 Algorithm flow chart for the spectral calibration of the VIPA spectrometer.

fringe recognition algorithm.²² Finally, the (X, Y, λ) relationship is determined using (m, Y_p, λ) , (X_p, m) and angle γ .

Experimental setup

The schematic diagram of the experimental setup for CO₂ absorption spectroscopy is shown in Fig. 3. The experimental laser was a fiber output supercontinuum source (SC-5, YLS Photonics, with a wavelength range of 470 to 2400 nm). The

output light was collimated using a fiber collimator and then selected by two filters to have a wavelength range of 1425 to 1450 nm (6900 to 7018 cm⁻¹) for CO₂ absorption detection. The filtered light was coupled into the Chernin gas cell, which consisted of five plano-concave mirrors, each with a radius of 0.5 m and an optical path length of 4 m. The output beam of the gas cell was sent to the self-designed VIPA spectrometer.²⁹

The fiber-coupled input of the VIPA spectrometer was launched into free space and line-focused onto the entrance window of the VIPA etalon (free spectral range, FSR ~ 60 GHz) using a cylindrical lens ($f_c = 100$ mm). An orthogonally oriented reflective diffraction grating (GH25-24V, Thorlabs, 1200 lines per mm) was used to separate the overlapping orders. The two-dimensionally dispersed light was imaged onto the camera using a plano-convex imaging lens ($f = 200$ mm). The maximum distortion aberration of this type of imaging system is about 8 μ m (less than 1 pixel), which is estimated using Zemax. The camera (LD-SW640, Leading Optoelectronic) had a pixel array of 640 × 512 with a pixel pitch of 15 μ m and a maximum full frame rate of 23 Hz. The camera was slightly rotated to align the fringes with the camera's y-axis.

Results and discussion

Wavelength calibration of the VIPA spectrometer

The relationship between (X, Y) and λ of selected spots. In this paper, a CO₂ absorption spectrum was used as the model input to achieve high-precision spectral calibration. Pure N₂

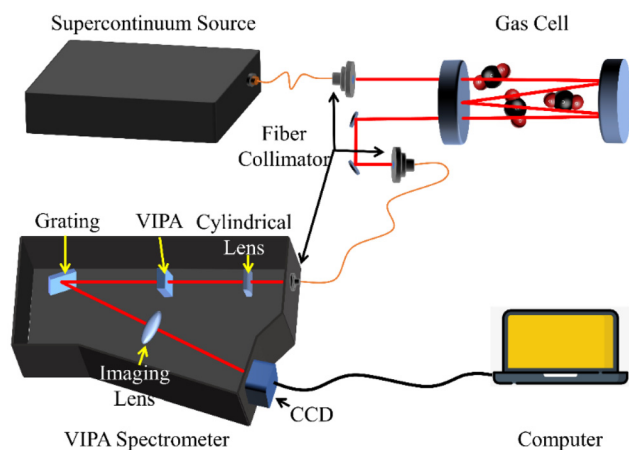


Fig. 3 The schematic diagram of the experimental setup. Filters are not shown in the figure.



(99.999%) and pure CO₂ (99.999%) were introduced into the gas cell to obtain the absorption spectrum. The cell pressure was stabilized at 200 mbar, and the temperature was 296 K. The collected images are shown in Fig. 4 ((a) is the N₂ background image (I_0) and (b) is the CO₂ signal image (I)). The dark image (I_d) was also collected.

The theoretical spectral coverage of the VIPA spectrometer was around 1425–1450 nm at an 85-degree grating incidence angle.²⁹ Since N₂ has no absorption in this region, the fringes in Fig. 4(a) are continuous and can be used as the absorption baseline. The black spots in Fig. 4(b) were caused by CO₂ absorption. The periodicity of these absorption spots can be used to identify different VIPA interference orders.^{1,16} By connecting these fringes end-to-end within one VIPA interference order, a one-dimensional spectrum without frequency axis calibration can be obtained. Using Klose's method,²² the wavelengths corresponding to the central pixels of these CO₂ absorption spots were obtained. In this paper, a portion of the data was used to fit the spectrogram model, and the rest was used as a control group to evaluate the accuracy of the wavelength calibration. The center pixel coordinates and their corresponding wavelengths used for the model inputs are shown in Table 1 and marked in Fig. 4(b).

The input matrix (m , Y_p , λ) of the ideal model. As shown in Fig. 4(b), the deviation of adjacent orders is about 9 pixels. Since the different VIPA interference orders of the same wavelength have the same x -coordinates in the ideal spectrogram model, the parameter γ can be determined using eqn (7). Taking the angle γ as the independent variable and 1×10^{-4} degrees as the iteration step size, the total differences of the rotated x -coordinates were calculated. The angle γ with the minimum total difference is -2.0293 degrees. The 21st fringe from the left to right in Fig. 4(b) is selected for analysis. Its VIPA order is set to m . The obtained rotated coordinates and their corresponding VIPA orders that can be used in the spectrogram model are shown in Table 2.

Fitting of the model parameters. The accuracy of the spectral calibration algorithm depends on the number of feature spots

Table 1 Coordinates of the absorption spot's central for different VIPA interference orders used to determine the γ angle. Columns (a) and (b) correspond to the yellow and pink squares in Fig. 4(b), respectively

Wavelength (nm)	(a) Coordinate of one interference order (X , Y)	(b) Coordinate of another interference order (X , Y)
1437.6679	(334, 103)	(343, 358)
1437.2197	(324, 129)	(333, 373)
1436.7871	(314, 141)	(324, 380)
1436.3699	(305, 141)	(314, 380)
1435.9681	(295, 129)	(304, 373)
1435.5817	(286, 105)	(295, 359)
1435.2107	(278, 62)	(285, 337)
1433.4446	(238, 151)	(247, 387)
1431.4172	(191, 116)	(199, 366)
1431.0323	(182, 89)	(189, 352)

Table 2 Coordinates and rotated coordinates of absorption spots used in the spectrogram model

Wavelength (nm)	VIPA order	Center coordinate (X , Y)	Corrected coordinate (X_p , Y_p)
1437.6679	$m - 16$	(343, 358)	(320.8421, 381.0921)
1437.2197	$m - 15$	(333, 372)	(310.3526, 394.7292)
1436.7871	$m - 14$	(324, 380)	(300.0756, 402.3701)
1436.3699	$m - 13$	(314, 380)	(291.0813, 402.0514)
1435.9681	$m - 12$	(304, 373)	(281.3354, 394.7017)
1435.5817	$m - 11$	(295, 359)	(272.8368, 380.3918)
1435.2107	$m - 10$	(285, 337)	(263.6221, 358.0515)
1433.4446	$m - 5$	(238, 151)	(223.2380, 170.5038)
1431.4172	$m - 1$	(199, 366)	(176.6491, 383.9880)
1431.0323	$m - 0$	(189, 352)	(167.1512, 369.6427)

and their location in the spectrogram image. Although the spectrogram model theoretically only requires three feature wavelengths to achieve the second-order polynomial fit, in practical applications, generally speaking, the more feature spots there are and the wider their distribution in the y -axis direction, the higher is the calibration accuracy. Here, we first used 37 different feature spots to achieve high accuracy fitting. Then, we used different numbers and combinations of feature

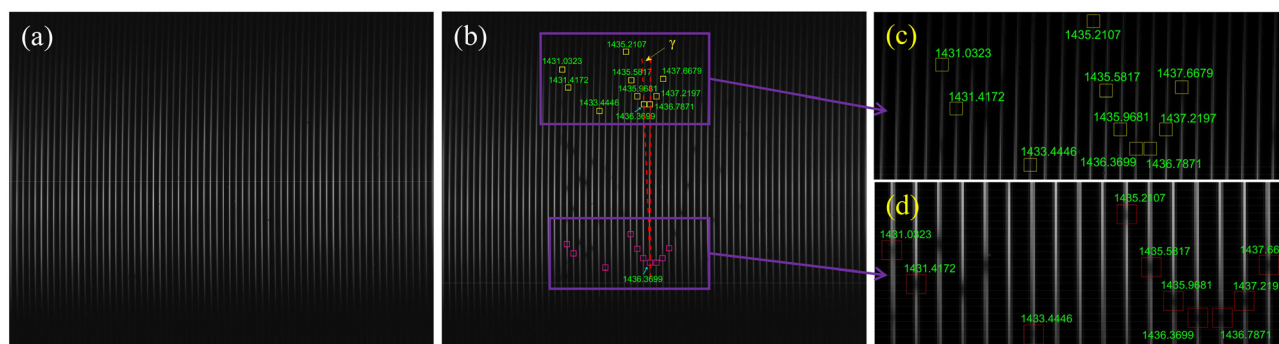


Fig. 4 Spectral images collected by the VIPA spectrometer. (a) The background image I_0 of pure N₂ and (b) the signal image I of pure CO₂. (c) The zoomed-in view of one VIPA FSR in figure b and (d) the zoomed-in view of another VIPA FSR. The yellow squares represent the absorption spots given in Table 1. Yellow and pink represent different VIPA regions, in a one-to-one correspondence. According to the Beer–Lambert law, the absorption image can be obtained by $-\ln[(I - I_d)/(I_0 - I_d)]$.



spots as model inputs and compared the fitting results. It was found that for an accurate fitting of model parameters, it is ideal to have 10 or more feature spots spread broadly (such as Table 2) as the model inputs.

According to eqn (5), $m\lambda$ (OPD) and the pixel vertical axis Y_p satisfy a quadratic polynomial ($m\lambda = \text{OPD} = a_2 Y_p^2 + a_1 Y_p + a_0$).¹⁶ Therefore, the closer the assumed value of m is to the real VIPA interference order, the smaller the quadratic polynomial fitting residual between $m\lambda$ and Y_p . Using the parameter m as an independent variable, the least-squares method is used to fit $m\lambda$ and Y_p . The fitting residual can be expressed as follows:

$$\text{residual} = \sum_{i=1}^n [m\lambda_i - a_2 Y_{pi}^2 - a_1 Y_{pi} - a_0]^2 \quad (10)$$

where n is the number of input relationships (λ and pixel coordinate).

Fig. 5(a), (b), and (c) show the second-order polynomial fitting results of the corrected y-coordinates and $m\lambda$ values of 37 absorption spots for three different m values (3413, 3454, and 3493), respectively. Fig. 5(d) shows the total residuals under different m values (from 3400 to 3500). The total

residuals were down and then up, which illustrates that the m value approaches the real VIPA interference order first and then away. When $m = 3454$, the fitting residual is the smallest, which represents the real order of the 21st fringe being 3454. Under this condition, the parameters a_2 , a_1 and a_0 of the second-order polynomial are -0.00647 , -2.3535 and $4\,944\,539.5519$, respectively.

Relationship between (X, Y) and λ over the entire absorption image. The same fringe recognition algorithm as Nugent-Glandorf *et al.*'s²² was used to obtain the central pixel coordinates of the fringes in images I and I_0 . Their corresponding coordinates were calculated using eqn (8) to obtain the matrix (X, Y, m) and the corresponding rotated matrix (X_p, Y_p, m) . Fig. 6 shows the theoretical spectrum calculated with the fitting parameters and different m values. The matrix (Y_p, m, λ) can be obtained from the curve. The one-to-one relationship between these three matrices allows us to determine the wavelength (λ) of each central pixel (X, Y) on the fringes in images I and I_0 . The wavelengths of the absorption spots within one VIPA FSR were labeled as shown in Fig. 7.

The accuracy of the spectrogram model

In general, the centroid of light spots in the measured spectrogram is regarded as the position corresponding to the absorption line.²³ To verify the accuracy of the spectrogram model, the centroid coordinates of some absorption peaks (including the absorption spots that are not involved in model parameter fitting) were calculated and compared with the peak coordinates calculated using the spectrogram model. The comparison results of 36 CO₂ absorption lines are shown in Fig. 8, where the red and green points represent the error of the x-coordinate and the y-coordinate, respectively. The results

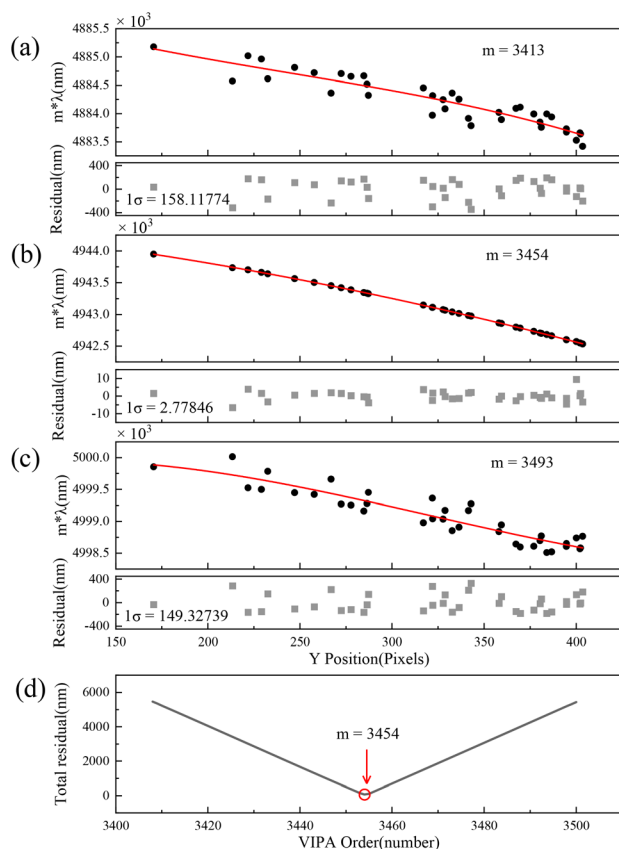


Fig. 5 Schematic diagram of the inversion of optimized model parameters. The second-order polynomial fitting results under the conditions of (a) $m = 3413$, (b) $m = 3454$, and (c) $m = 3493$. (d) The sum of the fitting residuals at different m values.

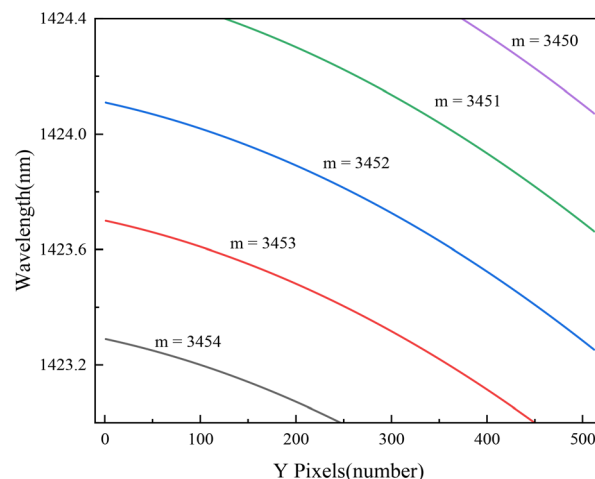


Fig. 6 Theoretical spectrum calculated by fitting parameters with different m values. Different lines (different m values) represent different fringes on the detector plane. The X coordinate of the pixel can be obtained by the fringe recognition algorithm. The Y coordinate of the pixel and its wavelength can be obtained from the curve, and thus the wavelength corresponding to each central pixel on the fringe can be obtained.



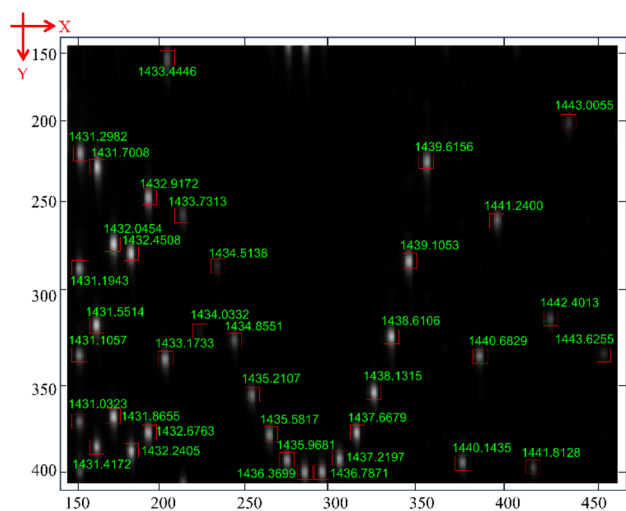


Fig. 7 Zoomed-in CO₂ absorption image. Absorption spots within a VIPA FSR are labelled with their wavelength.

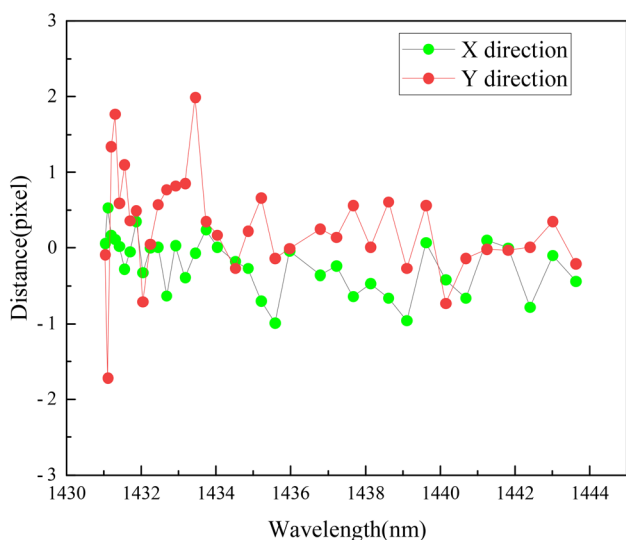


Fig. 8 The errors of the spectrogram model. The red point represents the error in the X direction and the green point represents the error in the Y direction.

show that the maximum calibration error is less than 2 pixels and the average error is about 1 pixel. Using the linear dispersion equation of the VIPA etalon,¹⁰ the maximum frequency error can be calculated to be 2.6 pm and the average frequency error is 0.88 pm. Compared to the deviation of the proportional model (~ 20 pm)¹⁶ and the linear model (~ 6 pm),¹⁶ the accuracy of the spectrogram model is better. Compared to the orthogonal dispersive model, although the accuracy is similar, the spectrogram model is more concise and easier to solve.

Since the fringe recognition algorithm can offset the effect of distortion to a certain extent,²¹ the deviation in the y-direction is larger than that in the x-direction. It is expected that

the calibration accuracy can be improved to the sub-pixel level by combining distortion correction algorithms. In addition, it is expected that self-calibration can be further achieved by adding compensation functions (such as the method of Shen *et al.*¹⁹) or automatically calculating the deviation (such as the method of Sadler *et al.*³⁰). Therefore, the spectrogram model is a promising technique that will play an important role in the applications of VIPA spectrometers.

Conclusion

In this work, a new simple, easy-to-solve and accurate spectrogram model calibration algorithm was developed for the wavelength calibration of VIPA spectrometers. The method simplified the dispersion model into 5 parameters and further considered the angle direction. The average accuracy was better than 1 pixel, which corresponds to a frequency accuracy of 0.88 pm. By combining with the distortion correction algorithm, the accuracy can be further improved. The reported method provides a useful tool for precision measurements.

Author contributions

HZ: investigation, methodology, and writing – original draft. WXZ: conceptualization, resources, supervision, and writing – review & editing. WHC and BXL: validation. FB, NNY and GFX: data curation. WJZ, LHD and WDC: resources and review. All authors contributed to discussions and the final version of the manuscript.

Data availability

All data are presented in the manuscript, and the files are available from the authors upon request.

Conflicts of interest

The authors declare no conflicts of interest.

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References

- 1 M. J. Thorpe, D. Balslev-Clausen, M. S. Kirchner, J. Ye, K. D. Moll, R. J. Jones, B. Safdi and J. Ye, *Opt. Express*, 2008, **16**, 2387–2397.
- 2 K. C. Cossel, E. M. Waxman, I. A. Finneran, G. A. Blake, J. Ye and N. R. Newbury, *J. Opt. Soc. Am. B*, 2017, **34**, 104–128.
- 3 S. A. Diddams, L. Hollberg and V. Mbele, *Nature*, 2007, **445**, 627–630.
- 4 X. Zhu and J. He, *J. Opt.*, 2019, **21**, 025703.
- 5 W. Bao, Z. Ding, P. Li, Z. Chen, Y. Shen and C. Wang, *Opt. Express*, 2014, **22**, 10081–10090.
- 6 C. Wang, Z. Ding, S. Mei, H. Yu, W. Hong, Y. Yan and W. Shen, *Opt. Lett.*, 2012, **37**, 4555–4557.
- 7 X. Zhu, D. Lin, Z. Zhang, X. Xie and J. He, *Astron. J.*, 2023, **165**, 228.
- 8 A. J. Fleisher, B. J. Bjork, T. Q. Bui, K. C. Cossel, M. Okumura and J. Ye, *J. Phys. Chem. Lett.*, 2014, **5**, 2241–2246.
- 9 B. J. Bjork, T. Q. Bui, O. H. Heckl, P. B. Changala, B. Spaun, P. Heu, D. Follman, C. Deutsch, G. D. Cole, M. Aspelmeyer, M. Okumura and J. Ye, *Science*, 2016, **354**, 444–448.
- 10 S. Xiao, A. M. Weiner and C. Lin, *IEEE J. Quantum Electron.*, 2004, **40**, 420–426.
- 11 S. Xiao and A. M. Weiner, *IEEE Photonics Technol. Lett.*, 2005, **17**, 372–374.
- 12 S. K. Scholten, J. D. Anstie, N. B. Hébert, R. T. White, J. Genest and A. N. Luiten, *Opt. Lett.*, 2016, **41**, 1277.
- 13 L. Nugent-Glandorf, T. Neely, F. Adler, A. J. Fleisher, K. C. Cossel, B. Bjork, T. Dinneen, J. Ye and S. A. Diddams, *Opt. Lett.*, 2012, **37**, 3285–3287.
- 14 S. Xiao and A. M. Weiner, *Opt. Express*, 2004, **12**, 2895–2902.
- 15 X. Zhu, D. Lin, Z. Hao, L. Wang and J. He, *Astron. J.*, 2020, **160**, 135.
- 16 G. Kowzan, K. F. Lee, M. Paradowska, M. Borkowski, P. Ablewski, S. Wójtewicz, K. Stec, D. Lisak, M. E. Fermann, R. S. Trawiński and P. Masłowski, *Opt. Lett.*, 2016, **41**, 974–977.
- 17 A. Klose, G. Ycas, F. C. Cruz, D. L. Maser and S. A. Diddams, *Appl. Phys. B*, 2016, **122**, 78.
- 18 G. Kowzan, D. Charczun, A. Cygan, R. S. Trawiński, D. Lisak and P. Masłowski, *Sci. Rep.*, 2019, **9**, 8206.
- 19 M. Shen, Z. Hao, X. Li, C. Li, L. Guo, Y. Tang, P. Yang, X. Zeng and Y. Lu, *Opt. Express*, 2018, **26**, 34131–34141.
- 20 J. Zhu, C. Sun, J. Yang, T. Ma, X. Guo and J. Zhang, *Opt. Precis. Eng.*, 2020, **28**, 1627–1633.
- 21 R. M. Bernstein, S. M. Burles and J. X. Prochaska, *Publ. Astron. Soc. Pac.*, 2015, **127**, 911–930.
- 22 L. Nugent-Glandorf, F. R. Giorgetta and S. A. Diddams, *Appl. Phys. B*, 2015, **119**, 327–338.
- 23 L. Yin, Bayanheshig, J. Yang, Y. Lu, R. Zhang, C. Sun and J. Cui, *Appl. Opt.*, 2016, **55**, 3574–3581.
- 24 H. Zhou, W. Zhao, B. Lv, W. Cui, B. Fang, N. Yang and W. Zhang, *Acta Opt. Sin.*, 2023, **43**, 1899914.
- 25 J. E. Greivenkamp, *Field Guide to Geometrical Optics (SPIE)*, 2004.
- 26 D. M. Bailey, G. Zhao and A. J. Fleisher, *Anal. Chem.*, 2020, **92**, 13759–13766.
- 27 A. Wang, T. Qiu and L. Shao, *J. Math. Imaging Vis.*, 2009, **35**, 165–172.
- 28 Y. Zhang, *Image Engineering*, Tsinghua University Press, 4th edn, 2018.
- 29 H. Zhou, W. Zhao, B. Fang, B. Lv, W. Cui, W. Zhang and W. D. Chen, *Analyst*, 2023, **148**, 4421–4428.
- 30 D. A. Sadler, D. Littlejohn and C. V. Perkins, *J. Anal. At. Spectrom.*, 1995, **10**, 253–257.

