

Multivariate analysis and digital twin modelling: Alternative approaches to evaluate molecular relaxation in photoacoustic spectroscopy

A. Zifarelli^a, A.F.P. Cantatore^a, A. Sampaolo^{a,b,*}, M. Mueller^{c,d}, T. Rueck^c, C. Hoelzl^e, H. Rossmadl^e, P. Patimisco^{a,b}, V. Spagnolo^{a,b}

^a PolySense Lab, Dipartimento Interateneo di Fisica, University and Politecnico of Bari, Via Amendola 173, 70126 Bari, Italy

^b PolySense Innovations S.R.L. via Amendola 173, Bari, Italy

^c Sensorik-ApplikationsZentrum (SappZ), Regensburg University of Applied Sciences, 93053 Regensburg, Germany

^d Institute of Analytical Chemistry, Chemo, and Biosensors, University of Regensburg, 93053 Regensburg, Germany

^e Thorlabs GmbH, Münchner Weg 1, 85232 Bergkirchen, Germany

ARTICLE INFO

Keywords:

relaxation effects photoacoustic spectroscopy
multivariate analysis
digital twin model
QEPAS sensor
methane detection over a wide concentration range
water vapor influence on relaxation
Partial Least Squares Regression

ABSTRACT

A comparative analysis of two different approaches developed to deal with molecular relaxation in photoacoustic spectroscopy is here reported. The first method employs a statistical analysis based on partial least squares regression, while the second method relies on the development of a digital twin of the photoacoustic sensor based on the theoretical modelling of the occurring relaxations. Methane detection within a gas matrix of synthetic air with variable humidity level is selected as case study. An interband cascade laser emitting at 3.345 μm is used to target methane absorption features. Two methane concentration ranges are explored targeting different absorptions, one in the order of part-per-million and one in the order of percent, while water vapor absolute concentration was varied from 0.3 % up to 2 %. The results achieved employing the detection techniques demonstrated the possibility to efficiently retrieve the target gas concentrations with accuracy > 95 % even in the case of strong influence of relaxation effects.

1. Introduction

Gas sensing technologies based on optical spectroscopy have been widely investigated in the past decades with the aim of developing reliable sensors to be operated in real-world applications [1,2]. In the large panorama of available detection techniques, photoacoustic spectroscopy (PAS) represents a well-consolidated technology, characterized by versatility, robustness, and sensitivity. PAS demonstrated the detection of different gas species for several applications, as environmental monitoring [3,4], human healthcare [5,6], and industrial processes control [7,8]. In PAS, modulated optical radiation is absorbed by the target analytes and converted into acoustic waves are generated by photoacoustic effect. Then, the acoustic waves are detected by a transducer which in turn returns an electric signal proportional to the amplitude of the pressure wave [9]. Typically, in standard PAS setup resonant acoustic cells are used to amplify the generated photoacoustic waves and highly sensitive condenser microphones are used for detection. However, in the recent years different detection systems have been investigated, including: optical detection systems, e.g., interferometry

or fiber Bragg grating device; MEMS-based devices; and cantilever beams [10–13]. Quartz-enhanced photoacoustic spectroscopy (QEPAS) was proposed as a development of traditional PAS, exploiting a quartz tuning fork (QTF) as sharply resonant transducer [14]. Compared to the standard PAS setup the QTF acts as both acoustic resonator and electrical transducer, providing smaller footprint and increased ruggedness. Millimetric acoustic resonator tubes are usually coupled with the QTF to amplify the generated photoacoustic waves, acting like organ pipes. In recent years, QEPAS sensors have been widely employed for trace gas sensing, providing high sensitivity and versatility thus making them suitable to target multiple applications [15–19]. PAS and QEPAS fall within the category of indirect detection techniques, as the gas absorption is evaluated by means of the energy deposited in the sample by the light source and converted into pressure waves. Therefore, the transducer response is independent of the wavelength selected to excite the target analyte, and this characteristic makes this kind of sensors a suitable approach to multi-gas detection relying on broadband laser sources in particular [20]. However, the photoacoustic conversion of incident radiation into acoustic waves depends on gas sample

* Corresponding author at: PolySense Lab, Dipartimento Interateneo di Fisica, University and Politecnico of Bari, Via Amendola 173, 70126 Bari, Italy.

E-mail address: angelo.sampaolo@poliba.it (A. Sampaolo).

<https://doi.org/10.1016/j.pacs.2023.100564>

Received 16 July 2023; Received in revised form 28 September 2023; Accepted 6 October 2023

Available online 9 October 2023

2213-5979/© 2023 The Author(s).

Published by Elsevier GmbH. This is an open access article under the CC BY license

(<http://creativecommons.org/licenses/by/4.0/>).

composition both in terms of heat generation and pressure waves propagation [21]. Dealing with trace gas detection, the latter phenomenon has a minor effect on sensors response, while the heat conversion efficiency may be significant. The non-radiative relaxation process leading to heat generation in the gaseous sample is mainly determined by the transfer rate of the vibrational energy of excited target molecules into kinetic or vibrational energy of surrounding molecules, labelled as V-T and V-V relaxation, respectively [22–24]. Following the absorption of a photon flux with a harmonic modulation at frequency f , the influence of energy transfer processes on the photoacoustic waves generation can be expressed by the radiation-to-sound conversion efficiency parameter, labelled ϵ [25]. This parameter ranges from 0 to 1 depending on all the relaxation pathways of the targeted gas mixture, thus is strongly dependent on the energetic levels distribution of the molecules composing the sample [26–28]. For this reason, intentionally humidifying the gas sample has been widely employed in QEPAS sensing in the past years, since water (H_2O) is known to prevent incomplete radiation-to-sound conversion due to its promoting effect [29–31]. More recently, approaches based on multi-gas detection and signal compensation by means of an external detector have been developed to filter out the molecular relaxation dependencies [32,33]. These methods demonstrated a good efficiency, but the performed calibrations were limited to a narrow range of target gas concentrations as well as a narrow range of humidity levels, sufficiently high to maximize the conversion efficiency ϵ [34]. Aiming to develop a gas sensor for on field measurements capable to operate under different conditions, it is mandatory to employ more sophisticated data analysis techniques to model the sensor's response.

Recently, two opposite approaches aiming to deal with matrix effects in photoacoustic spectroscopy have emerged, one based on a multivariate analysis (MVA) as Partial Least Squares Regression (PLSR) [35] and one based on a digital twin (DT) of the developed sensor [36]. These methods address the same issue from two completely different perspectives. The PLSR-based approach relies on the statistical evaluation of the cross-correlation induced on QEPAS signal by the matrix effects while the DT-based approach relies on the theoretical computation of all the relaxation processes occurring in the gas sample.

In this work, we report on a compared investigation on the PLSR-based and DT-based data analysis technique with the aim of filtering out the molecular relaxation effects from QEPAS signal. The two approaches are tested on the same experimental datasets, collected targeting gas mixtures composed of methane (CH_4) and H_2O in synthetic air. An interband cascade laser (ICL) with central emission wavelength of $3.345 \mu\text{m}$ ($\sim 2989 \text{ cm}^{-1}$) is used as light source to target CH_4 absorption features. This spectral region is well-known in literature for the strong matrix effects occurring on the photoacoustic signal corresponding to CH_4 absorptions due to the influence of water vapor and oxygen [23,26,33,34]. Two CH_4 concentration ranges are explored separately, targeting different absorption features within the laser dynamic range: a “low-concentration range” in the order of part-per-million (ppm), and a “high concentration range” in the order of percent. The two data analysis techniques allowed the retrieval of CH_4 concentrations within different samples compositions and the comparative analysis provided an outline of the advantages and disadvantages of the presented methods.

2. Partial least squares regression

The use of this statistical tool to model spectroscopic systems has already proven to be effective, benefitting from the large number of information acquired by the optical sensors and being able to deal with spectral and non-spectral interference occurring in PAS and QEPAS measurements [35]. Among the different possibilities, MVA represented a solid class of regression and classification algorithms which have been successfully applied to spectroscopic techniques [37]. Partial least squares regression (PLSR) is a MVA technique representing a

development of traditional multiple linear regression, developed to deal with noisy and highly correlated data [38,39]. When applied to optical spectroscopy, this tool has already demonstrated to be a reliable approach to complex gas mixtures [40,41]. PLSR extends the traditional linear regression model to include correlation effects, mathematically corresponding to collinearity in the matrices. The linear regression model can be expressed in matrix form as $\mathbf{Y} = \mathbf{X}\mathbf{B} + \mathbf{E}$, where \mathbf{X} is the matrix containing the experimental acquisitions, i.e., the spectra, \mathbf{Y} the matrix of the physical parameters to be estimated, i.e., the gas concentrations, \mathbf{E} is the residuals matrix, and \mathbf{B} is the matrix of the regression coefficients. To include correlation effects within the matrices, PLSR assumes that the system is described in terms of truly independent factors called latent variables (LVs) or components. LVs are extracted from the matrices \mathbf{X} and \mathbf{Y} by maximizing the covariance matrix $\text{cov}(\mathbf{X}, \mathbf{Y})$, thus projecting the matrices into a new vector space described by the LVs. This operation, called *projection on latent structures* allows the algorithm to perform a linear regression on truly orthogonal and independent vectors, thus returning solid regression coefficients and high predictive power [42]. The number of LVs represents an input parameter for the analysis, and thus it must be carefully evaluated to prevent the regression from being affected by under- or over-fitting of the data [43]. The high versatility of the algorithm and the possibility to describe different systems without a priori investigations make this approach a valuable tool for gas sensing spectroscopy.

3. Digital twin

The behavior of complex systems with regard to variations of different parameters is often difficult to predict. In 2003, Michael Grieves proposed the concept of a digital twin (DT) to address this challenge, promising many advantages, like better and more realistic predictions resulting in appropriate actions to be taken [44,45]. The most common application areas of the DT are the manufacturing sector, with keywords predictive maintenance, smart factory and industry 4.0 as well as prognostics and health management (PHM) [46]. Shafto et al. described a DT to be “an integrated multi-physics, multi-scale probabilistic simulation of a [...] system that uses the best available physical models, sensor updates [...] to mirror the life of its [...] twin” [47]. With the intention to improve the reliability of sensor systems, the concept of the DT was applied to a photoacoustic trace gas sensor for the first time in 2023 [36]. The quantities to be considered in view of DT compensation to finally associate an analyte concentration c_{CH_4} with a photoacoustic amplitude U are reported in Eq.(1):

$$U \sim (\gamma - 1) \frac{Q}{f} c_{\text{CH}_4} \sigma(\lambda) P_0 \epsilon \quad (1)$$

where, γ is the heat capacity ratio of the gas mixture, Q the quality factor of the resonator, $\sigma(\lambda)$ the absorption cross section at the selected wavelength, P_0 the incident laser power. For this purpose, the quality factor and the resonance frequency of the QTF, as well as the optical power used for photoacoustic signal generation are measured, while the absorption cross-section of the sample at the wavelength of excitation is simulated using HITRAN database. The core of the DT is the algorithm named CoNRad, which allows the calculation of the collision-based non-radiative relaxation efficiency ϵ , as well as the heat capacity ratio γ of the gas mixture, considering pressure, temperature, composition of the mixture, and the laser modulation frequency [48]. According to the definition used by Shafto et al., CoNRad represents the multi-physics, multi-scale probabilistic simulation based on physical models, while the sensor updates include the optical power, temperature, pressure, quality factor, the frequency of the QTF, and the humidity of the sample. Finally, the DT uses those sensor updates together with ϵ and γ , computed by CoNRad, to predict a photoacoustic signal U_{calc} for a random analyte concentration c_{CH_4} . The DT calculates a theoretical QEPAS signal U_{calc} for each measurement point, compares this to the

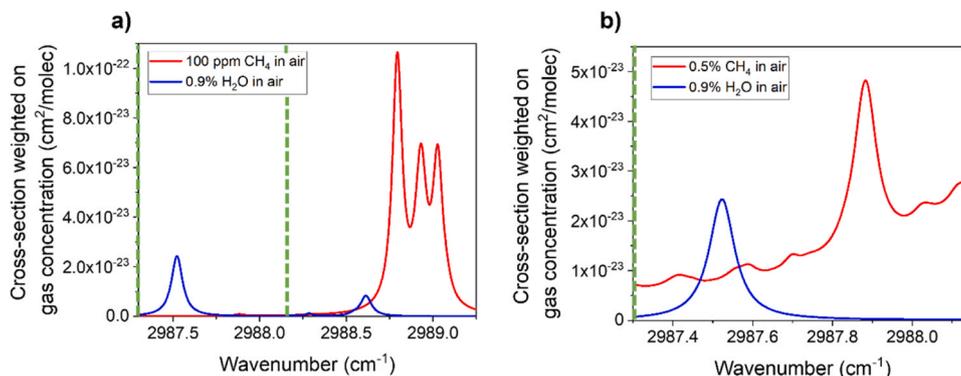


Fig. 2. HITRAN simulation at $P = 400$ Torr of the spectral regions selected to acquire QEPAS spectra of gas samples with (a) low CH_4 concentration (25–200 ppm), and (b) high CH_4 concentration (1100–11,000 ppm, 0.11–1.10 %). The absorption spectra of CH_4 (red lines) are simulated at concentrations representative for the concentration range, while the absorption spectra of H_2O (blue line) are simulated at a typical atmospheric concentration. Green dashed lines in panel (a) point out the spectral region shown in panel (b).

to select the methane dilution ratio in synthetic air and to set the flow rate in the gas line. The gas mixer is characterized by an accuracy of 1 % of the setpoint, for each channel. The output of the gas mixer was downstream connected to a humidifier (PermSelect PDMSXA-1000) to set the in-line H_2O concentration by varying the aperture of two needle valves, as depicted in Fig. 1, from 0.3 % up to 2 % of absolute humidity. A pressure controller (ALICAT EPC-15PSIAP01-BM0P), a flow meter (ALICAT BC-C1000), a needle valve, and a diaphragm pump (Thomas 1420VR 24 V) were embedded in the sensor box. These devices were employed to set the operating pressure and monitor the flow rate in the gas line: all the measurements were performed at 400 Torr and 50 sccm, respectively.

3.2. Target features selection

Due to the limitation on the voltage input of the employed digital lock-in amplifier, two different concentration ranges were selected to calibrate the QEPAS sensor and perform both the DT and PLSR analysis. The first range spans from 25 ppm to 200 ppm of CH_4 and is labelled hereafter as “low CH_4 concentration range”, while the second range spans from 1100 ppm to 11,000 ppm (0.11–1.10 %) and is labelled hereafter as “high CH_4 concentration range”. Thus, two spectral regions were selected within the DFB-ICL current dynamic range and simulated by using the HITRAN database [50], as reported in Fig. 2a-b together with the absorption spectrum of H_2O at typical atmospheric concentration of 0.9 %.

The methane triplet located between 2988.50 cm^{-1} and 2989.25 cm^{-1} (Fig. 2a) was targeted to detect CH_4 within the “low concentration range”. These features exhibit an absorption cross section weighted with CH_4 concentration in the order of $\sim 10^{-22}\text{ cm}^2/\text{molecule}$ (see Fig. 2a). Moving to the CH_4 detection at percentage level the use of these features would have led to two different issues, related to: i) nonlinearities in Lambert-Beer absorption; ii) lock-in signal saturation. Therefore, an absorption feature located at 2987.87 cm^{-1} was employed to target the “high CH_4 concentration range” (Fig. 2b), since at percentage level it provided an absorption cross section weighted with CH_4 concentration comparable to that estimated in the ppm range (Fig. 2a).

3.3. QEPAS sensor calibration

The gas sensor was calibrated for both the low CH_4 concentration range and for the high CH_4 concentration range, respectively, employing the same procedure for both data acquisition and data analysis.

The measurements were performed setting the H_2O level in the samples and varying the CH_4 concentration in the investigated range. The water vapor concentration was acquired by means of the hygrometer housed inside the ADM (see Fig. 1). The RH value was then converted into an absolute H_2O concentration, with known temperature and pressure inside the cell. Once set the H_2O concentration, two subsequent acquisition steps were performed for each CH_4 concentration. First, the QEPAS spectrum of the CH_4 - H_2O mixture in air was acquired operating the sensor in spectral scan mode. Then, the peak value of the selected

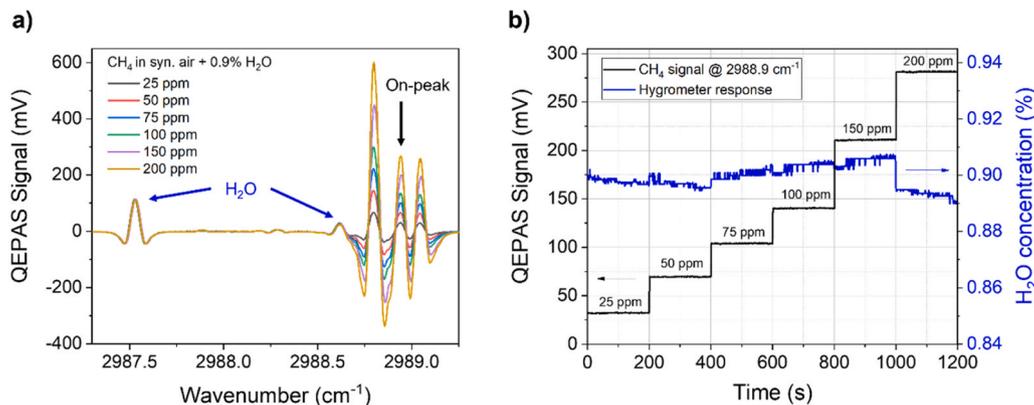


Fig. 3. (a) 2-f-QEPAS spectral scans acquired in the low CH_4 concentration range with a H_2O concentration of 0.9 %. Blue arrows point towards H_2O absorption features, black arrow points towards the CH_4 feature used for on-peak measurements. (b) Stepwise representation of CH_4 on-peak measurements (black line) and corresponding H_2O concentration (blue line) calculated using the capacitive hygrometer. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

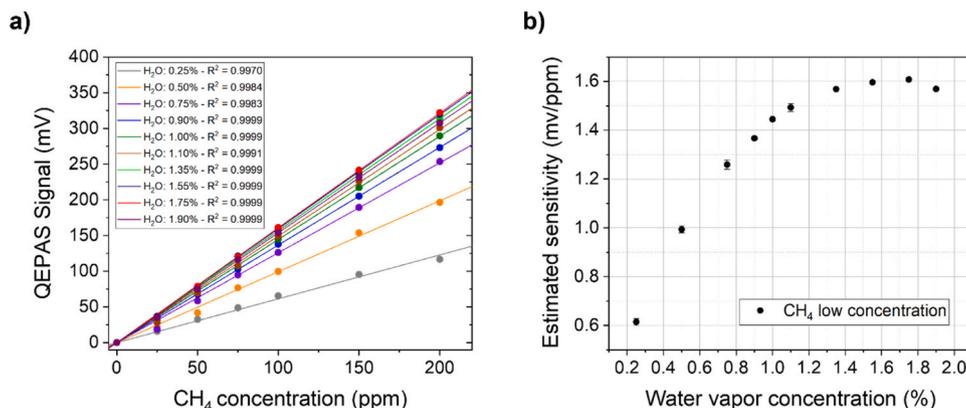


Fig. 4. (a) QEPAS peak signal of CH₄ in the low concentration range corresponding to the absorption feature at 2898.95 cm⁻¹, for each investigate humidity level (circular dots). A linear fit is superimposed to each measurement set (solid lines), and the corresponding value of R² is reported in the graph's legend. (b) QEPAS sensor sensitivities estimated by the best linear fit as a function of H₂O concentration.

CH₄ absorption feature was measured operating the sensor in on-peak mode. The QEPAS spectra have been employed as dataset for the PLSR analysis, while the on-peak acquisitions have been fed to the DT analysis.

3.4. Low CH₄ concentration range

The performances of the QEPAS sensor when targeting CH₄-H₂O mixtures with low CH₄ concentration were evaluated analyzing 60 gas samples with different composition. Six CH₄ concentrations, from 25 ppm up to 200 ppm, and ten H₂O concentrations, from 0.25 % up to 1.90 % were mixed in the samples starting from the gas cylinder with certified concentrations and diluted in synthetic air.

The QEPAS spectra acquired at different CH₄ concentrations with a H₂O level of 0.9 % are shown in Fig. 3a as representatives. Each spectrum was acquired by means of a single sweep and consists of 537 data points with a spectral resolution of ~0.0036 cm⁻¹ and total acquisition time was ~4 min. Such a wide scan allowed the detection of both the H₂O spectral features falling within the laser dynamic range as well as the CH₄ triplet.

The collected QEPAS spectral scans resemble the shape of the 2nd derivative of the HITRAN simulation shown in Fig. 2a. The methane triplet can be clearly observed, being characterized by three well separated peaks whose intensities scales proportionally to the CH₄ concentration. Two H₂O absorption features can be observed: one at ~2987.55 cm⁻¹ well separated from the triplet, and one at ~2988.60 cm⁻¹ partially merged to the first CH₄ peak. The detection phase for lock-in acquisition was set to the one maximizing the in-phase

signal generated by the central peak of CH₄ triplet at the lowest water concentration and was kept fixed for all the measurements.

After each spectral scan, the sensor was operated in on-peak mode, to acquire the data for DT analysis. To perform the analysis in the low CH₄ concentration range, the QEPAS peak located at 2988.93 cm⁻¹ was preferred to the adjacent and more intense one located at 2988.80 cm⁻¹ to avoid any influence from the nearby water peak at ~2988.60 cm⁻¹. Each QEPAS peak signal was acquired for 200 s before changing the sample composition in the gas line, to ensure that no signal drift occurs in the sensor. The average value was considered as reference value for DT analysis. The driving current of the DFB-ICL was locked to the selected CH₄ peak by means of the photodiode signal demodulated at the third harmonic, and employed as error signal [51]. The QEPAS peak signals acquired at different CH₄ concentrations with a nominal H₂O level of 0.9 % are shown in Fig. 3b as representatives. The water vapor monitoring acquired by the hygrometer is also reported in the same figure. During acquisitions at a fixed water vapor (Fig. 3b), small fluctuations around the mean value were observed. This effect occurs because the water vapor level is slightly influenced by the operating conditions: the valve opening dynamics of the mixer channels operates on a different time scale with respect to the passive humidifier, which reacts much slowly. Nevertheless, the measured relative variation around the mean value was within 3 %, for all the samples: this fluctuation leads to a negligible influence on CH₄ QEPAS signal. Indeed, the relative fluctuations of the CH₄ signal was below 2 %.

To point out the effects of H₂O on CH₄ photoacoustic response, the QEPAS peak signals were plotted in Fig. 4a as a function of the CH₄ concentration, for each humidity level.

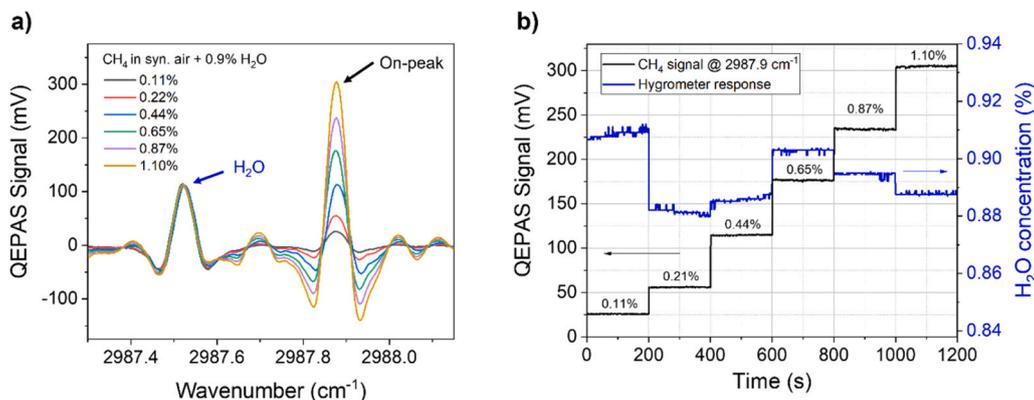


Fig. 5. (a) 2 f-QEPAS spectral scans acquired in the high CH₄ concentration range with a H₂O concentration of 0.9 %. Blue arrow points towards H₂O absorption feature. (b) Stepwise representation of CH₄ on-peak measurements (black line) and corresponding H₂O concentration (blue line).

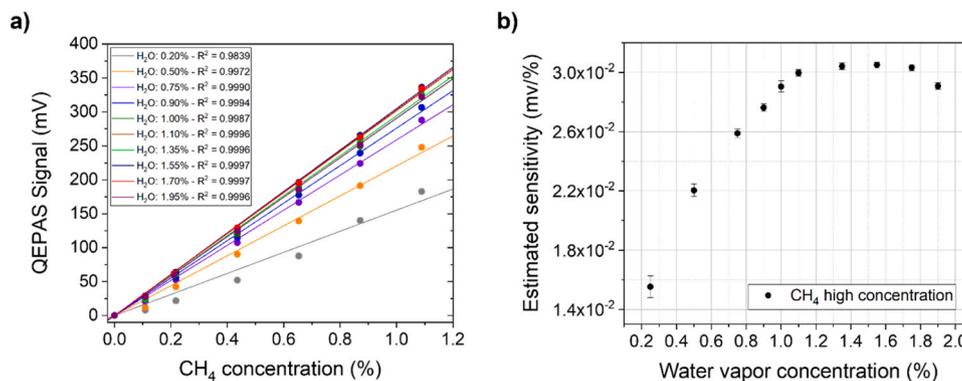


Fig. 6. QEPAS peak signal of CH₄ in the high concentration range corresponding to the absorption feature at 2897.87 cm⁻¹, for each investigate humidity level (circular dots). A linear fit is superimposed to each measurement set (solid lines), and the corresponding value of R² is reported in graph's legend. (b) QEPAS sensor sensitivities estimated by the best linear fit as a function of H₂O concentration.

The linear response of the sensor is verified in the investigated concentration range for each humidity level. As expected, the sensitivity the CH₄ QEPAS changes when the H₂O concentration is varied (Fig. 4b). For H₂O concentrations up to 1.0 % the sensitivity increases accordingly, while at higher H₂O concentrations the sensor's response reaches a plateau with a slight decrease at humidity level beyond 1.8 %.

3.5. High CH₄ concentration range

The performances of the QEPAS sensor when targeting CH₄-H₂O mixtures with high CH₄ concentration were evaluated analyzing 60 gas samples with different composition. Six CH₄ concentrations, from 0.11 % up to 1.10 %, and ten H₂O concentrations, from 0.20 % up to 1.95 % were mixed in the samples starting from the gas cylinder with certified concentrations to be diluted in synthetic air.

The QEPAS spectrum of each gas sample was acquired operating the sensor in spectral scan mode. The QEPAS spectra acquired at different CH₄ concentrations with a H₂O level of 0.9 % are shown in Fig. 5a as representatives. Each spectrum was acquired by means of a single sweep and consisted in 237 data points with a spectral resolution of ~0.0036 cm⁻¹ and total acquisition time was ~2 min.

The methane absorption feature located at 2987.87 cm⁻¹, as well as some other minor CH₄ features, can be observed in the full spectral scan. The H₂O absorption feature located at 2987.55 cm⁻¹ can be also observed in the collected scans. Differently from the previous case, a minor interference among CH₄ and H₂O absorptions is observed. The strongest CH₄ feature at 2987.87 cm⁻¹ was used for the on-peak measurements. The detection phase for lock-in acquisition was set to the one maximizing the in-phase signal generated by the selected CH₄ absorption feature and kept fixed for all the measurements.

Analogously to the measurements in low CH₄ concentration range, after each spectral scan the QEPAS peak signal was acquired for 200 s. The QEPAS peak signals acquired at different CH₄ concentrations with a nominal H₂O level of 0.9 % are shown in Fig. 5b, together with the H₂O concentration monitoring provided by the capacitive hygrometer. As for the measurements at low CH₄ concentrations, relative fluctuations < 3 % around the mean value were observed during the measurement sessions at fixed water concentrations, leading to a negligible effect on CH₄ peak signal amplitude which exhibited relative fluctuation < 2 % for all generated gas samples.

The QEPAS peak signals as a function of the CH₄ concentration for each humidity level were plotted in Fig. 6a.

The linear response of the sensor is verified also in the high CH₄ concentration range, for each humidity level. The sensitivity of the CH₄ QEPAS sensor depends on the humidity level (Fig. 6b), exhibiting a trend similar to the one observed for the low CH₄ concentrations (Fig. 4b). Up to H₂O concentrations of 1.0 %, the CH₄ response increases accordingly,

while at higher H₂O concentrations the sensor's response reaches a plateau with a slight decrease at humidity level beyond 1.80 %.

3.6. Data analysis and concentration retrieval

The information acquired from spectral scans and on-peak measurements were used to perform PLSR and DT analysis, respectively, aiming for filtering out the effects of variable water vapor on the CH₄ sensitivity and returning an accurate prediction of CH₄ concentration.

3.7. DT algorithm optimization

The core of the DT is the algorithm CoNRad presented in the previous section, which allows the user to calculate the expected relaxational efficiency, based on the measurement conditions. The on-peak QEPAS amplitude as well as its phase were included into the evaluation process since the phase also contains information about cross-influences. In cases where the measured amplitude is well reproduced by the theoretical one, but significant discrepancies occur between measured and theoretical phase shift, it must be assumed that the physical model of the DT lacks completeness, e.g. not considering relevant energy transitions, or assuming wrong transition rates. A complete relaxational diagram including the corresponding transition rates for mid-infrared methane detection in humidified air was already presented in Ref.[48]. The relaxations considered for the investigated samples as well as the reaction rate employed in the CoNRad algorithm are reported in Table S1 of supplementary file.

3.8. PLSR algorithm optimization

PLSR data analysis was performed independently for each concentration range, using the same algorithm. The predictors matrix \mathbf{X} was assembled starting from the acquired spectral scans, while the response matrix \mathbf{Y} contains the corresponding nominal concentrations. Both in-phase and quadrature spectra acquired from the lock-in amplifier were employed in the analysis to account for the phase shift on CH₄ QEPAS signals induced by different H₂O concentrations. The spectral acquisitions shown in Figs. 3a and 5a were employed for PLSR analysis, covering the spectral range from 2989.3 cm⁻¹ - 2987.3 cm⁻¹ and 2988.2 cm⁻¹ - 2987.3 cm⁻¹ for low and high CH₄ concentration, respectively. PLSR was performed in machine learning-like approach, splitting the dataset into a calibration set and a test set. To perform a consistent comparison with the DT analysis each sample was tested independently, thus with a single test sample and with 59 calibration samples.

A 10-fold cross-validation (CV) analysis was performed prior to the test step to determine the optimal number of LVs for each concentration

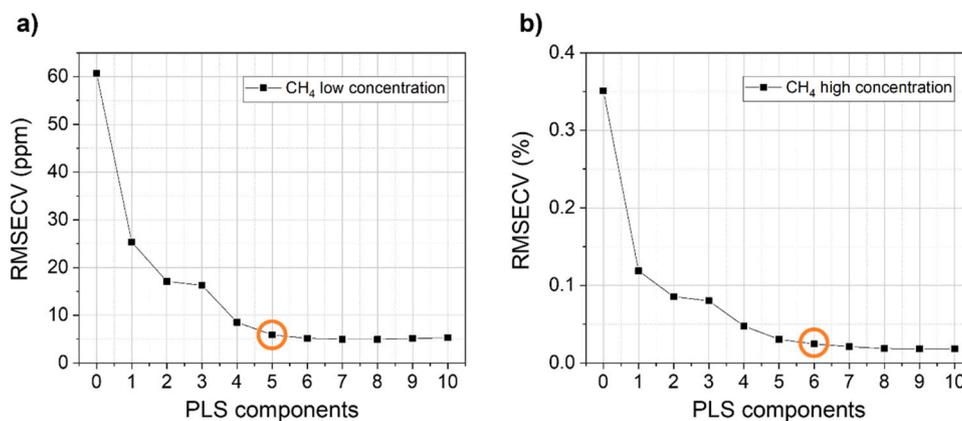


Fig. 7. RMSECV as a function of the number of PLS components for (a) low CH_4 concentrations and (b) high CH_4 concentrations. Orange circles point out the number of LVs selected for the analysis. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

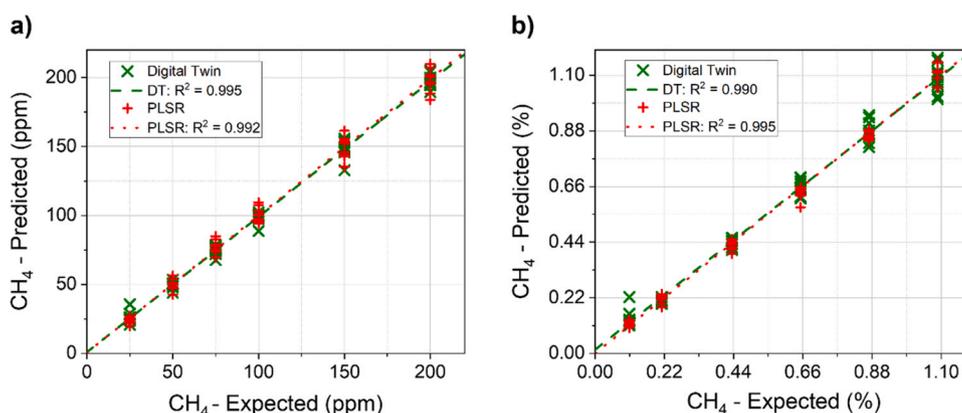


Fig. 8. Comparison between the DT (green \times) and PLSR (red $+$) results for (a) low CH_4 concentrations and (b) high CH_4 concentrations. Methane concentrations retrieved using the algorithms are plotted against the expected, nominal, concentrations in the samples. A linear fit is superimposed to the both the DT data (green dashed line) and PLSR data (red dotted line). The corresponding R^2 values are reported in graph's legend. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

range. The LVs evaluation was operated on the full measurement dataset. The CV errors, expressed as root-mean-square error (RMSECV), calculated for the two concentration ranges are shown in Fig. 7a-b.

Calibration dataset for low CH_4 concentrations shows the decrease of RMSECV up to 5 PLS components, namely LVs, while the error values corresponding to a larger number of components are characterized by negligible variations. Conversely, the calibration dataset for high CH_4 concentrations shows a significant RMSECV decrease up to 6 PLS components. PLSR is not meant to be interpreted as an explainable AI [52], but it is possible to correlate the LVs with independent, physically relevant parameters contributing to the spectra [42]. In this case, it would be possible to assign a LV to: i) CH_4 concentration; ii) H_2O concentration; iii) the photoacoustic relaxation rate of CH_4 through H_2O ; iv) phase of the acquired signal for each water concentration; and v) resonance properties of the QEPAS spectrophone affected by variations of fluid dynamics properties in the gas samples. The additional components observed at high CH_4 concentration can be ascribed to the self-relaxation of CH_4 , which can be assumed negligible at low concentrations [26].

3.9. Results comparison

All the acquired spectra were tested employing the algorithms configurations presented in the previous paragraphs. The CH_4 concentrations retrieved employing DT and PLSR analysis targeting the low and high CH_4 concentration range are shown in Fig. 8a-b, respectively.

Table 1

Results of best linear fit superimposed to the predicted versus expected concentration graphs.

	Low CH_4 concentration range			High CH_4 concentration range		
	R^2	Slope	Intercept (ppm)	R^2	Slope	Intercept (%)
DT	0.995	0.981 ± 0.009	0.928 ± 1.038	0.990	0.983 ± 0.013	0.015 ± 0.009
PLSR	0.992	0.992 ± 0.012	0.725 ± 1.372	0.996	1.001 ± 0.009	0.001 ± 0.006

The achieved results reveal a linear trend of the predicted values versus the expected values for both the concentration ranges and both the analysis methods, as demonstrated by the superimposed best linear fit, whose calculated parameters are reported in Table 1.

Table 2

Results comparison between DT and PLSR analysis for both low and high CH_4 concentration range in terms of calculated AREP and absolute deviation.

	DT – AREP (rel.%)	PLSR – AREP (rel.%)	DT - Abs. dev. (ppm)	PLSR - Abs. dev. (ppm)
Low CH_4 range	3.8 %	4.6 %	2.9	3.9
High CH_4 range	7.6 %	4.9 %	248	179

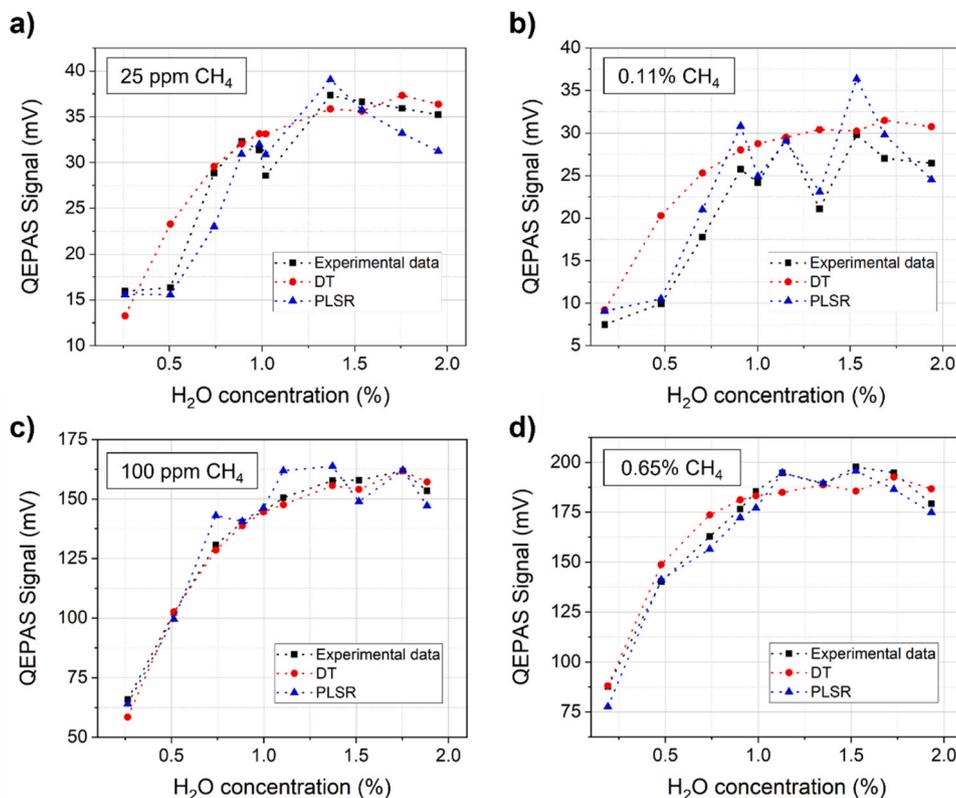


Fig. 9. Comparison among the QEPAS signal acquired experimentally (black squares), calculated theoretically using DT (red dots) and regressed from PLSR predicted concentrations (blue triangles) for the set characterized by CH₄ concentration of (a) 25 ppm; (b) 0.11 %; (c) 100 ppm; (d) 0.65 %. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

The R^2 values are ≥ 0.990 and the calculated slopes approach the ideal value of 1. Both the analysis techniques point out a negligible intercept within the error limits, $< 1\sigma$ for all the datasets except for DT at high concentrations ($< 2\sigma$).

The average relative error of prediction (AREP) as well as the mean absolute deviation from the expected values were used to evaluate the accuracy of the predicted concentrations. These values, corresponding to each data analysis technique and each concentration range, are shown in Table 2.

In the low CH₄ concentration range, DT results are slightly better than PLSR predictions providing lower discrepancies and thus higher accuracies. Conversely, in the high CH₄ concentration range, PLSR shows a higher accuracy compared to DT. Comparing the two concentration ranges, the high concentration one shows lower accuracy for both analysis methods, with a slight degradation (from 4.6 % to 4.9 %) for PLSR and a more significant one (from 3.8 % to 7.6 %) for DT. The one-to-one comparison of the retrieved results is reported in Table S2 and Table S3 of supplementary files. The achieved results show variable discrepancies between the nominal and predicted concentrations. In particular, the test samples characterized by lowest CH₄ concentrations in both concentration ranges (25 ppm and 0.11 %) as well as the test samples characterized by the lowest H₂O level (0.20 % and 0.25 %) pointed out the lowest relative accuracy in the dataset.

The interpretation of the obtained results can be effectively pursued in the perspective of the intrinsic characteristics and differences between the two analysis approaches. If the components of the gas matrix are known and their physical properties in terms of relaxation pathways fully and accurately determined, DT analysis is expected to be more accurate than PLSR, being supported by a systematic computation of all the contributions to the QEPAS signal generation rather than the statistical interpretation of the spectra. In addition, DT analysis can be easily implemented for real-time conversion of QEPAS signal into target

gas concentration. Moreover, relying on a full characterization of energy dynamics, the algorithm can be easily adapted to target another gas species or to address the effect of a matrix variation, once provided collisional partners and related energy levels. The main drawback of DT approach is that it requires a complete knowledge of all the involved physical phenomena as well as a full control of all the parameters characterizing the complex gas mixtures. Instrumental parameters, high inaccuracy of instruments when operating close to their limit (as for the generation of the lowest concentrations, i.e., the 25 ppm and 0.11 % CH₄-mixtures), variations of relaxation dynamics when a large number of target molecules are involved (potentially responsible for the low accuracy of DT in the high CH₄ concentrations range) are hard to be modelled. Conversely, PLSR points out a higher versatility characterized by stable performances which can be easily generalized to different datasets. Due to the multivariate nature of the analysis, PLSR can deal with overlapping features and be less prone to inaccuracy due to external instruments, i.e., the hygrometer, when the information is included in the spectra. The main drawback is that the results obtained for a sensor cannot be generalized for another sensor and time-consuming acquisitions to build a training dataset are always mandatory. To visualize these effects, the collected QEPAS peak signals are compared with the signals theoretically calculated by the DT. In addition, we included the QEPAS signal regressed from the predicted PLSR concentrations exploiting the linear response of the sensor, and the results are shown in Fig. 9. These signals are close but not coincident to the peak signals acquired in the spectral scans like the ones showed in Fig. 5, as the PLSR analysis account for all the data to retrieve the concentrations.

The data reported in Figs. 9a and 9b were acquired setting the same experimental parameters in the gas line (pressure, flow rate, temperature, and gas mixer apertures), changing the starting certified concentration. The same applies to the data reported in Figs. 9c and 9d. The first

pair of graphs exhibits the higher signal fluctuations ascribed to instrumental accuracy, while the second pair points out the excellent theoretical modelling at low concentrations compared to a less accurate one in the high concentrations range.

The possibility to model sensor's behaviors that are repeatable and related to minor but physically relevant phenomena can be an advantage of statistically based approach like PLSR, however the occurrence of overfitting as well as systematic errors should be carefully evaluated. Conversely, the advantages of theoretically based approaches like DT relies on the possibility to simulate the sensor behavior upstream filtering the possible experimental fluctuations. Aiming to develop a gas sensor for on field operation this may represent a disadvantage or an advantage as moving outside the controlled laboratory environment it is easy to run into unpredictable and uncontrolled fluctuations affecting sensor's calibration.

The effects of the measurements set characterized by the higher discrepancies on the overall accuracy can be evaluated in terms of relative and absolute accuracy. Removing the measurements with 25 ppm of CH₄ the DT-AREP reduces from 3.8 % to 2.9 % while its absolute deviation increases from 2.9 ppm to 3.0 ppm. In turn the PLSR-AREP reduces from 4.6 % to 4.2 % while its absolute deviation increases from 3.9 ppm to 4.4 ppm. Removing the measurements with 0.11 % of CH₄ the DT-AREP reduces from 7.6 % to 3.5 % while its absolute deviation reduces from 248 ppm to 237 ppm. In turn the PLSR-AREP reduces from 4.9 % to 3.5 % while its absolute deviation increases from 179 ppm to 189 ppm.

4. Conclusions

In this work, a comparative investigation upon two different data analysis techniques used to compensate the effects of molecular relaxation in photoacoustic spectroscopy is presented. The first one was based on PLSR analysis while the second one was based on the DT representation of the experimental system. To address this task, a dedicated QEPAS sensor was developed and embedded in a shoe-size box containing the sensor head, the electronic instrumentation, and the gas line management devices. For this case study, CH₄ was selected as target molecule and the effects of variable H₂O concentration (from 0.3 % up to 2.0 %) within a synthetic air matrix were investigated in the spectral range around 3.345 μm. Two CH₄ concentration ranges were explored, one in the ppm range and one in the percent range, to account for different applications. PLSR-based approach relies on the statistical evaluation of the collected spectral scans to model the cross-correlation among the analytes in the gas sample and retrieve the target concentration. DT-based approach relies on an algorithmic approach to compute the whole relaxation dynamics occurring in the gas sample by exploiting the collected on-peak target signal. Despite of the strong relaxation effects induced by the different humidity levels in the samples, both analysis tools were able to return target concentrations with average accuracy ~95 %. In the low concentration range, DT performed better compared to PLSR benefitting of its strong theoretical background. In the high concentration range, an opposite situation was observed that can be ascribed to a partial spectral interference among H₂O and CH₄. Further developments of both analysis algorithms will be focused on improving the achieved accuracy as well as on testing more complex mixtures, in terms of number of analytes as well as spectral and non-spectral cross-interference effects.

Declaration of Competing Interest

The authors declare the following financial interests/personal relationships which may be considered as potential competing interests: Angelo Sampaolo reports was provided by Polytechnic University of Bari.

Data availability

Data will be made available on request.

Acknowledgements

The authors from Dipartimento Interateneo di Fisica di Bari acknowledge financial support from THORLABS GmbH within the Pol-ySenSe joint research laboratory.

Appendix A. Supporting information

Supplementary data associated with this article can be found in the online version at doi:10.1016/j.pacs.2023.100564.

References

- [1] A. Fathy, Y.M. Sabry, I.W. Hunter, D. Khalil, T. Bourouina, Direct absorption and photoacoustic spectroscopy for gas sensing and analysis: a critical review, *Laser Photon. Rev.* 16 (2022) 2100556.
- [2] J. Hodgkinson, R.P. Tatam, Optical gas sensing: a review, *Meas. Sci. Technol.* 24 (2013), 012004.
- [3] C. Feng, M. Giglio, B. Li, A. Sampaolo, P. Patimisco, V. Spagnolo, L. Dong, H. Wu, Detection of hydrogen sulfide in sewer using an erbium-doped fiber amplified diode laser and a gold-plated photoacoustic Cell, *Molecules* 27 (2022) 6505.
- [4] H. Xiao, J. Zhao, C. Sima, P. Lu, Y. Long, Y. Ai, W. Zhang, Y. Pan, J. Zhang, D. Liu, Ultra-sensitive PPB-level methane detection based on NIR all-optical photoacoustic spectroscopy by using differential fiber-optic microphones with gold-chromium composite nanomembrane, *Photoacoustics* 26 (2022), 100353.
- [5] C. Popa, A.M. Bratu, M. Petrus, A comparative photoacoustic study of multi gases from human respiration: mouth breathing vs. nasal breathing, *Microchem. J.* 139 (2018) 196–202.
- [6] Y. Zhang, Y. Xie, J. Lu, J. Zhao, Y. Wu, J. Tong, J. Shao, Continuous real-time monitoring of carbon dioxide emitted from human skin by quartz-enhanced photoacoustic spectroscopy, *Photoacoustics* 30 (2023), 100488.
- [7] X. Yin, Y. Su, T. Xi, B. Chen, L. Zhang, X. Zhang, L. Liu, X. Shao, Research progress on photoacoustic SF₆ decomposition gas sensor in gas-insulated switchgear, *J. Appl. Phys.* 131 (2022), 130701.
- [8] X. Yin, H. Wu, L. Dong, W. Ma, L. Zhang, W. Yin, L. Xiao, S. Jia, F.K. Tittel, Ppb-level photoacoustic sensor system for saturation-free CO detection of SF₆ decomposition by use of a 10 W fiber-amplified near-infrared diode laser, *Sens. Actuators B Chem.* 282 (2019) 567–573.
- [9] S. Palzer, Photoacoustic-based gas sensing: a review, *Sensors* 20 (2020) 2745.
- [10] X. Zhang, L. Liu, Y. Liu, L. Zhang, X. Yin, H. Huan, T. Xi, X. Shao, Detectors for gas-phase photoacoustic spectroscopy: a review, *Microw. Opt. Technol. Lett.* (2022).
- [11] T. Yang, W. Chen, P. Wang, A review of all-optical photoacoustic spectroscopy as a gas sensing method, *Appl. Spectrosc. Rev.* 56 (2021) 143–170.
- [12] T. Strahl, J. Steinebrunner, C. Weber, J. Wöllenstein, K. Schmitt, Photoacoustic methane detection inside a MEMS microphone, *Photoacoustics* 29 (2023), 100428.
- [13] T. Tomberg, M. Vainio, T. Hieta, L. Halonen, Sub-parts-per-trillion level sensitivity in trace gas detection by cantilever-enhanced photo-acoustic spectroscopy, *Sci. Rep.* (2018) 1–7.
- [14] A.A. Kosterev, Y.A. Bakhirkin, R.F. Curl, F.K. Tittel, Quartz-enhanced photoacoustic spectroscopy, *Opt. Lett.* 27 (2002) 1902.
- [15] R. De Palo, A. Elefante, G. Biagi, F. Paciolla, R. Weih, V. Villada, A. Zifarelli, M. Giglio, A. Sampaolo, V. Spagnolo, P. Patimisco, Quartz-enhanced photoacoustic sensors for detection of eight air pollutants, *Adv. Photon. Res.* (2023).
- [16] B. Li, G. Menduni, M. Giglio, P. Patimisco, A. Sampaolo, A. Zifarelli, H. Wu, T. Wei, V. Spagnolo, L. Dong, Quartz-enhanced photoacoustic spectroscopy (QEPAS) and Beat Frequency-QEPAS techniques for air pollutants detection: a comparison in terms of sensitivity and acquisition time, *Photoacoustics* (2023), 100479.
- [17] Y. Liu, H. Lin, B.A.Z. Montano, W. Zhu, Y. Zhong, R. Kan, B. Yuan, J. Yu, M. Shao, H. Zheng, Integrated near-infrared QEPAS sensor based on a 28 kHz quartz tuning fork for online monitoring of CO₂ in the greenhouse, *Photoacoustics* 25 (2022), 100332.
- [18] B. Li, C. Feng, H. Wu, S. Jia, L. Dong, Calibration-free mid-infrared exhaled breath sensor based on BF-QEPAS for real-time ammonia measurements at ppb level, *Sens. Actuators B Chem.* 358 (2022), 131510.
- [19] G. Menduni, A. Zifarelli, E. Kniazeva, S. Dello Russo, A.C. Ranieri, E. Ranieri, P. Patimisco, A. Sampaolo, M. Giglio, F. Manassero, E. Dinuccio, G. Provolò, H. Wu, D. Lei, V. Spagnolo, Measurement of methane, nitrous oxide and ammonia in atmosphere with a compact quartz-enhanced photoacoustic sensor, *Sens. Actuators B Chem.* 375 (2023), 132953.
- [20] A. Sampaolo, P. Patimisco, M. Giglio, A. Zifarelli, H. Wu, L. Dong, V. Spagnolo, Quartz-enhanced photoacoustic spectroscopy for multi-gas detection: a review, *Anal. Chim. Acta* 1202 (2022), 338894.
- [21] B. Berne, R. Pecora. *Dynamic Light Scattering: With Applications to Chemistry, Biology, and Physics*, Springer, Boston, MA, 2000.
- [22] G. Wysocki, A.A. Kosterev, F.K. Tittel, Influence of molecular relaxation dynamics on quartz-enhanced photoacoustic detection of CO₂ at λ = 2 μm, *Appl. Phys. B Lasers Opt.* 85 (2006) 301–306.

- [23] N. Barreiro, A. Vallespi, G. Santiago, V. Slezak, A. Peuriot, Influence of oxygen on the resonant photoacoustic signal from methane excited at the ν_3 mode, *Appl. Phys. B* 104 (2011) 983–987.
- [24] B. Lang, P. Breittegger, G. Brunnhofer, P. Valero Jordi, A. Klug Simon Schweighart, W. Hassler, Alexander Bergmann, Molecular relaxation effects on vibrational water vapor photoacoustic spectroscopy in air, *Appl. Phys. B Aims Scope* 126 (2020) 64.
- [25] Y.-H. Pao (Ed.), *Optoacoustic Spectroscopy and Detection*, Academic Press, 1977.
- [26] N. Barreiro, A. Peuriot, G. Santiago, V. Slezak, Water-based enhancement of the resonant photoacoustic signal from methane-air samples excited at 3.3 μm , *Appl. Phys. B Lasers Opt.* 108 (2012) 369–375.
- [27] J. Hayden, B. Baumgartner, B. Lendl, Anomalous humidity dependence in photoacoustic spectroscopy of CO explained by kinetic cooling, *Appl. Sci.* 10 (2020) 843.
- [28] F. Sgobba, A. Sampaolo, P. Patimisco, M. Giglio, G. Menduni, A.C. Ranieri, C. Hoelzl, H. Rossmadl, C. Brehm, V. Mackowiak, D. Assante, E. Ranieri, V. Spagnolo, Compact and portable quartz-enhanced photoacoustic spectroscopy sensor for carbon monoxide environmental monitoring in urban areas, *Photoacoustics* 25 (2022), 100318.
- [29] A.A. Kosterev, Y.A. Bakhrin, F.K. Tittel, S. McWhorter, B. Ashcraft, QEPAS methane sensor performance for humidified gases, *Appl. Phys. B Lasers Opt.* 92 (2008) 103–109.
- [30] L. Dong, R. Lewicki, K. Liu, P.R. Buerki, M.J. Weida, F.K. Tittel, Ultra-sensitive carbon monoxide detection by using EC-QCL based quartz-enhanced photoacoustic spectroscopy, *Appl. Phys. B Lasers Opt.* 107 (2012) 275–283.
- [31] Y. Ma, R. Lewicki, M. Razeghi, F.K. Tittel, QEPAS based ppb-level detection of CO and N₂O using a high power CW DFB-QCL, *Opt. Express* 21 (2013) 1008.
- [32] H. Wu, L. Dong, X. Yin, A. Sampaolo, P. Patimisco, W. Ma, L. Zhang, W. Yin, L. Xiao, V. Spagnolo, S. Jia, Atmospheric CH₄ measurement near a landfill using an ICL-based QEPAS sensor with V-T relaxation self-calibration, *Sens. Actuators B Chem.* 297 (2019), 126753.
- [33] A. Elefante, G. Menduni, H. Rossmadl, V. Mackowiak, M. Giglio, A. Sampaolo, P. Patimisco, V.M.N. Passaro, V. Spagnolo, Environmental monitoring of methane with quartz-enhanced photoacoustic spectroscopy exploiting an electronic hygrometer to compensate the h₂o influence on the sensor signal, *Sensors* 20 (2020) 2935.
- [34] J. Pangerl, M. Müller, T. Rück, S. Weigl, R. Bierl, Characterizing a sensitive compact mid-infrared photoacoustic sensor for methane, ethane and acetylene detection considering changing ambient parameters and bulk composition (N₂, O₂ and H₂O), *Sens. Actuators B Chem.* 352 (2022), 130962.
- [35] G. Menduni, A. Zifarelli, A. Sampaolo, P. Patimisco, M. Giglio, N. Amoroso, H. Wu, L. Dong, R. Bellotti, V. Spagnolo, High-concentration methane and ethane QEPAS detection employing partial least squares regression to filter out energy relaxation dependence on gas matrix composition, *Photoacoustics* 26 (2022), 100349.
- [36] T. Rück, M. Müller, S. Jobst, S. Weigl, J. Pangerl, R. Bierl, F.M. Matysik, Digital Twin of a photoacoustic trace gas sensor for monitoring methane in complex gas compositions, *Sens. Actuators B Chem.* 378 (2023), 133119.
- [37] A.C. Olivieri, *Introduction to Multivariate Calibration: a Practical Approach*, Springer, Netherlands, 2018.
- [38] S. Wold, A. Ruhe, H. Wold, W.J. Dunn III, The collinearity problem in linear regression. the partial least squares (PLS) approach to generalized inverses, *SIAM J. Sci. Stat. Comput.* 5 (1984) 735–743.
- [39] S. Wold, M. Sjöström, L. Eriksson, PLS-regression: a basic tool of chemometrics, *Chemom. Intell. Lab. Syst.* 58 (2001) 109–130.
- [40] A. Loh, M. Wolff, Multivariate analysis of photoacoustic spectra for the detection of short-chained hydrocarbon isotopologues, *Molecules* 25 (2020).
- [41] A. Zifarelli, M. Giglio, G. Menduni, A. Sampaolo, P. Patimisco, V.M.N. Passaro, H. Wu, L. Dong, V. Spagnolo, Partial least-squares regression as a tool to retrieve gas concentrations in mixtures detected using quartz-enhanced photoacoustic spectroscopy, *Anal. Chem.* 92 (2020).
- [42] H. Abdi, Partial least squares regression and projection on latent structure regression (PLS Regression), *WIREs Comput. Stat.* 2 (2010) 97–106.
- [43] M.P. Gómez-Carracedo, J.M. Andrade, D.N. Rutledge, N.M. Faber, Selecting the optimum number of partial least squares components for the calibration of attenuated total reflectance-mid-infrared spectra of undesigned kerosene samples, *Anal. Chim. Acta* 585 (2007) 253–265.
- [44] M. Grieves, Digital twin: manufacturing excellence through virtual factory replication, *White Pap.* (2014) 7.
- [45] M. Singh, E. Fuenmayor, E. Hinchy, Y. Qiao, N. Murray, D. Devine, Digital twin: origin to future, *Appl. Syst. Innov.* 4 (2021) 36.
- [46] F. Tao, H. Zhang, A. Liu, A.Y.C. Nee, Digital twin in industry: state-of-the-art, *IEEE Trans. Ind. Inform.* 15 (2019) 2405–2415.
- [47] M. Shafto, M.C. Rich, D.E. Glaessen, C. Kemp, J. Lemoigne, L. Wang, *Draft Modeling, Simulation, Information Technology & Processing Roadmap Technology Area 11*, (2010).
- [48] M. Müller, T. Rück, S. Jobst, J. Pangerl, S. Weigl, R. Bierl, F.M. Matysik, An algorithmic approach to compute the effect of non-radiative relaxation processes in photoacoustic spectroscopy, *Photoacoustics* 26 (2022), 100371.
- [49] P. Patimisco, A. Sampaolo, M. Giglio, S. Dello Russo, V. Mackowiak, H. Rossmadl, A. Cable, F.K. Tittel, V. Spagnolo, Tuning forks with optimized geometries for quartz-enhanced photoacoustic spectroscopy, *Opt. Express* 27 (2019) 1401.
- [50] I.E. Gordon, L.S. Rothman, R.J. Hargreaves, R. Hashemi, E.V. Karlovets, F. M. Skinner, E.K. Conway, C. Hill, R.V. Kochanov, Y. Tan, P. Wcislo, A.A. Finenko, K. Nelson, P.F. Bernath, M. Birk, V. Boudon, A. Campargue, K.V. Chance, A. Coustenis, B.J. Drouin, J.M. Flaud, R.R. Gamache, J.T. Hodges, D. Jacquemart, E.J. Mlawer, A.V. Nikitin, V.I. Perevalov, M. Rotger, J. Tennyson, G.C. Toon, H. Tran, V.G. Tyuterev, E.M. Adkins, A. Baker, A. Barbe, E. Cané, A.G. Császár,

A. Dudaryonok, O. Egorov, A.J. Fleisher, H. Fleurbaey, A. Foltynowicz, T. Furtenbacher, J.J. Harrison, J.M. Hartmann, V.M. Horneman, X. Huang, T. Karman, J. Karns, S. Kassi, I. Kleiner, V. Kofman, F. Kwabia-Tchana, N. N. Lavrentieva, T.J. Lee, D.A. Long, A.A. Lukashovskaya, O.M. Lyulin, V. Y. Makhnev, W. Matt, S.T. Massie, M. Melosso, S.N. Mikhailenko, D. Mondelain, H. S.P. Müller, O.V. Naumenko, A. Perrin, O.L. Polyansky, E. Raddaoui, P.L. Raston, Z. D. Reed, M. Rey, C. Richard, R. Tóbiás, I. Sadiék, D.W. Schwenke, E. Starikova, K. Sung, F. Tamassia, S.A. Tashkun, J. Vander Auwera, I.A. Vasilenko, A.A. Viganin, G.L. Villanueva, B. Vispoel, G. Wagner, A. Yachmenev, S.N. Yurchenko, The HITRAN2020 molecular spectroscopic database, *J. Quant. Spectrosc. Radiat. Transf.* 277 (2022).

- [51] Q. Wang, Z. Wang, W. Ren, Wavelength-stabilization-based photoacoustic spectroscopy for methane detection, *Meas. Sci. Technol.* 28 (2017), 065102.
- [52] F. Akulich, H. Anahideh, M. Sheyyab, D. Ambre, Explainable predictive modeling for limited spectral data, *Chemom. Intell. Lab. Syst.* 225 (2022), 104572.



Andrea Zifarelli received the M.S. degree (cum laude) in Physics in 2018 from the University of Bari and his Ph.D. in Physics from the University of Bari in 2022. His research activities were mainly focused on the development of spectroscopic techniques based on laser absorption for the analysis of complex gas mixtures by employing quartz tuning forks as sensitive elements. This investigation was performed by using innovative laser sources as well as developing new algorithms for multivariate analysis approaches. Currently, his research activities are carried out at the PolySense Lab, joint-research laboratory between Technical University of Bari and THOR-LABS GmbH.



Aldo Francesco Pio Cantatore received his M.S. degree in Physics (cum laude) in 2022 from the University of Bari. From the same year, he is a Ph.D. student at the Physics Department of the University of Bari, developing his research work at PolySense Lab, joint research laboratory between the Polytechnic of Bari and THORLABS GmbH. His current research activities are mainly focused on the Quartz-Enhanced Photoacoustic Spectroscopy-based analysis of complex gas mixtures, as well as on the development of compact and portable gas sensors based on Light-Induced Thermoelastic Spectroscopy.



Angelo Sampaolo obtained his Master degree in Physics in 2013 and the Ph.D. Degree in Physics in 2017 from University of Bari. He was a visiting researcher in the Laser Science Group at Rice University from 2014 to 2016. Since March 2021, he is an assistant professor at the Technical University of Bari. His research activity has included the study of the thermal properties of heterostructured devices via Raman spectroscopy. Most recently, his research interest has focused on the development of innovative techniques in trace gas sensing, based on Quartz-Enhanced Photoacoustic Spectroscopy and covering the full spectral range from near-IR to THz. His achieved results have been acknowledged by a cover paper in *Applied Physics Letters* of the July 2013 issue.



Max Mueller received his master's degree in electrical and microsystems engineering (M. Eng.) from the Ostbayerische Technische Hochschule (OTH) of Regensburg in Germany in 2020. Currently he is pursuing his Doctorate of Natural Sciences (Dr. rer. nat.) in cooperation with the Institute for Analytical Chemistry, Chemo- and Biosensors at the University of Regensburg and the Sensorik-ApplikationsZentrum (SappZ) in Germany. Since 2018, he has been conducting research in the field of photoacoustic trace gas sensing and is focusing on vibrational energy transfer and classical acoustic phenomena.



Thomas Rueck studied Chemistry (Dipl. Chem.) at the University of Regensburg in Germany. In the course of his diploma thesis in cooperation with Continental Automotive GmbH, he started his research on the photoacoustic gas sensing technique in 2009. By now he has 13 years of experience in this field and received his Doctor of Science (Dr. rer. nat.) in 2017. Currently, Thomas Rück is head of the gas sensing team of the Sappz, which is affiliated to the Ostbayerische Technische Hochschule (OTH) of Regensburg.



Pietro Patimisco obtained the Master degree in Physics (cum laude) in 2009 and the Ph.D. Degree in Physics in 2013 from the University of Bari. Since 2020, he is Assistant professor at the University of Bari. He was a visiting scientist in the Laser Science Group at Rice University in 2013 and 2014. Dr. Patimisco's scientific activity addressed both micro-probe optical characterization of semiconductor optoelectronic devices and photoacoustic gas sensors. Recently, his research activities included the study and applications of trace-gas sensors, such as quartz enhanced photoacoustic spectroscopy and cavity enhanced absorption spectroscopy in the mid infrared and terahertz spectral region, leading to several publications, including a cover paper in Applied Physics Letter of the July

2013 issue.



Christine Hoelzl received the Dipl.-Phys. and Dr. rer. nat. degrees from Technical University Munich in 2006 and 2012, respectively. During her Ph.D. thesis, her research interest was in the field of ultrafast optical orientation and time-resolved spectroscopy of carrier spin dynamics in semiconductor materials. Currently, she is contributing to the development of spectroscopy solutions at THORLABS GmbH.



Vincenzo Spagnolo obtained the Ph.D. in physics in 1994 from University of Bari. From 1997–1999, he was researcher of the National Institute of the Physics of Matter. Since 2004, he works at the Technical University of Bari, formerly as assistant and associate professor and now as full Professor of Physics. Starting from 2019, he became Vice-Rector of the technical university of Bari - Deputy to Technology Transfer. He is the director of the joint-research lab PolySense between Technical University of Bari and THORLABS GmbH, fellow member of SPIE and senior member of OSA. His research interests include photoacoustic gas sensing and spectroscopic techniques for real-time monitoring. His research activity is documented by more than 220 publications and 3 filed patents. He has given more than 50 invited presentations at international conferences and workshops.

than 50 invited presentations at international conferences and workshops.



Hubert Rossmadl obtained his Diploma degree in Physics in 2006 while working at the ASDEX-Upgrade Experiment from the Max-Planck Institute for Plasma Physics and the Ph.D. degree in Physics in 2013 from the Technical University of Munich. His research activity has included the study of solvated electrons with ultrafast time-resolved absorption and fluorescence-spectroscopy reached via non-linear optics. After two years of industrial production of frequency-converted single-mode lasers at Toptica, he is since 2016 at Thorlabs developing innovative compact sensors for trace gas detection based on Quartz Enhanced Photoacoustic Spectroscopy.