Protonation enhancement by dichloromethane doping in low-pressure photoionization

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Doping has been used to enhance the ionization efficiency of analytes in atmospheric pressure photoionization, which is based on charge exchange. Compounds with excellent ionization efficiencies are usually chosen as dopants. In this paper, we report a new phenomenon observed in low-pressure photoionization: Protonation enhancement by dichloromethane (CH$_2$Cl$_2$) doping. CH$_2$Cl$_2$ is not a common dopant due to its high ionization energy (11.33 eV). The low-pressure photoionization source was built using a krypton VUV lamp that emits photons with energies of 10.0 and 10.6 eV and was operated at ~500–1000 Pa. Protonation of water, methanol, ethanol, and acetaldehyde was respectively enhanced by 481.7 ± 122.4, 197.8 ± 18.8, 87.3 ± 7.8, and 93.5 ± 35.5 times after doping 291 ppmv CH$_2$Cl$_2$, meanwhile CH$_2$Cl$_2$ almost does not generate noticeable ions itself. This phenomenon has not been documented in the literature. A new protonation process involving in ion-pair and H-bond formations was proposed to expound the phenomenon. The observed phenomenon opens a new prospect for the improvement of the detection efficiency of VUV photoionization.

Photoionization (PI), a widely used soft ionization technique, is usually coupled to various mass spectrometers for analyzing the chemical composition of samples1–3. Atmospheric pressure photoionization (APPI) is a new and highly attractive ionization technique4,5, which was developed ~10 years ago with the aim of improving the performance of liquid chromatography–mass spectrometry (LC-MS) for less polar compounds such as polycyclic aromatic hydrocarbons (PAHs). A krypton lamp, which emits VUV photons with energies of 10.0 and 10.6 eV, is usually chosen as the light source in APPI as it is cheap, compact, and robust. Different from classic vacuum photoionization, APPI shows characteristics more similar to those of chemical ionization (CI). The ionization mechanisms commonly observed in CI are also observed in APPI, such as the proton transfer reaction (PTR) and charge exchange. PTR typically takes place when the analyte in question has a higher proton affinity (PA), whereas charge exchange requires that the analyte possesses low ionization energy (IE).

Low-pressure photoionization (LPPI), defined as photoionization running under hundreds to thousands of Pa, has not been used as widely as APPI and conventional vacuum PI. LPPI has characteristics of both vacuum PI and APPI. Apart from molecular ions, protonated ions were found to be dominant for polar compounds. The proton transfer reactions in LPPI can be expressed as follows:

\[
A + R^+ \rightarrow AH^+ + [R-H] \tag{1}
\]

\[
A^+ + R \rightarrow AH^+ + [R-H] \tag{2}
\]

where A represents the analyte molecules and R is the reagent which offers a proton or hydrogen atom. The reagent could be the analyte or solvent molecules.

The use of dopants has been found to be very effective for enhancing the ionization efficiency of analytes6–11 in APPI and LPPI via charge exchange:

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where D and A represent dopant and analyte molecules, respectively. Benzene (IE = 9.24 eV)\(^ {13,14} \), acetone (IE = 9.70 eV)\(^ {10,15-17} \), toluene (IE = 8.83 eV)\(^ {6,7,10-12,18-20} \), and anisole (IE = 8.20 eV)\(^ {21} \) are often employed as dopants due to their excellent photoionization efficiencies under illumination of the krypton lamp. The resulting analyte ions may subsequently react with other molecules via proton transfer. The detection sensitivity could be enhanced by ~100 times via doping\(^ {22} \). However, these dopants cannot be applied to the detection of methanol (CH\(_3\)OH, IE = 10.84 eV), ethanol (C\(_2\)H\(_5\)OH, IE = 10.48 eV), and acetaldehyde (C\(_2\)H\(_4\)O, IE = 10.23 eV) due to their higher IEs. Dichloromethane has been chosen as a dopant for characterizing the molecular structures of analytes via secondary ion–molecule reactions, rather than for enhancing ionization efficiency\(^ {23} \).

Our previous studies revealed that LPPI with a specially designed photoionizer was super sensitive (~1000 counts/ppbv) to many organic compounds\(^ {24-26} \). However, the LPPI detection efficiency for CH\(_3\)OH, C\(_2\)H\(_5\)OH, and C\(_2\)H\(_4\)O is very low due to their low ionization efficiencies. In this paper, we report a new phenomenon: The detection efficiencies of the three small volatile organic compounds (VOCs) can be remarkably enhanced via CH\(_2\)Cl\(_2\) doping. The results and experimental method are described in the following sections.

**Results**

**Protonation enhancement of water and LPPI mass spectrum of CH\(_2\)Cl\(_2\)**. Water (H\(_2\)O) is an important protonation agent for PTR mass spectrometry. The IE of water is 12.62 eV, which indicates that it cannot be photoionized directly by the photons emitted from the krypton lamp. However, H\(_3\)O\(^ + \) (m/z 19, 45 counts), (H\(_2\)O)\(_2\)H\(^ + \) (m/z 37, 214 counts), and (H\(_2\)O)\(_3\)H\(^ + \) (m/z 55, 24 counts) were observed in the LPPI mass spectrum of N\(_2\), as shown in Fig. 1(A). The concentration of water in the test chamber was <5 ppmv, as a result of impurities in high-purity N\(_2\) gas. Protonation of acetonitrile (IE = 12.20 eV) was observed in APPI with a krypton lamp as the VUV light source by Marotta et al. The authors speculate that photon irradiation leads first to the isomerization of acetonitrile molecules, affording species that exhibit IEs <10 eV and that consequently are able to generate photoionization products\(^ {27} \). The formation mechanism of protonated water and water clusters under illumination of 10.0 and 10.6 eV photons is not clear yet. In view of a tiny amount of N\(_2\)\(^ + \) (m/z 28, 34 counts) observed in Fig. 1(A), the photoelectrons in the photoionization region might lead to the formation of protonated water and water clusters. Figure 1(B) shows the mass spectrum obtained after injecting 291 ppmv CH\(_2\)Cl\(_2\) into the chamber. Surprisingly, the signal intensities of H\(_3\)O\(^ + \), (H\(_2\)O)\(_2\)H\(^ + \), and (H\(_2\)O)\(_3\)H\(^ + \) increased to 2.92 × 10\(^ 4 \), 1.24 × 10\(^ 5 \) and 2.29 × 10\(^ 4 \) counts, respectively. The signal intensity of protonated water was averagely enhanced by 481.7 ± 122.4 times, measured from three independent measurements. This phenomenon has never been reported.

CH\(_2\)Cl\(_2\) is a common solvent used in organic analysis. The IE of CH\(_2\)Cl\(_2\) is 11.33 eV. It cannot be directly ionized by the VUV photons emitted from the krypton lamp. As shown in Fig. 1(B), no noticeable ions were produced from direct photoionization of CH\(_2\)Cl\(_2\). A small mass peaks at m/z 47 is assigned to ethanol residual in the test chamber or minor impurity in the CH\(_2\)Cl\(_2\) reagent.

**Protonation enhancement of methanol, ethanol, and acetaldehyde**. Methanol (CH\(_3\)OH) is the simplest alcohol. Its IE is 10.84 eV, higher than the energy of the photons emitted from the krypton lamp. A weak signal of protonated methanol was observed when 4.6 ppmv methanol was sampled. Figure 2(A) shows the obtained LPPI mass spectrum of 4.6 ppmv methanol in nitrogen. The mass peaks at m/z 19, 37, and 55 correspond

\[
D + h\nu \rightarrow D^+ + e \quad (3)
\]

\[
D^+ + A \rightarrow A^+ + D \quad (4)
\]
to H₃O⁺, (H₂O)₂H⁺, and (H₂O)₃H⁺, respectively. The mass peaks at m/z 33, 51 and 65 are assigned to (CH₃OH)H⁺, (CH₃OH-H₂O)H⁺ and (CH₃OH)₂H⁺, respectively. The moderate mass peak at m/z 47 is assigned to ethanol, the impurity in the methanol reagent. The peak intensities of (CH₃OH)H⁺ and (CH₃OH)₂H⁺ are 559 and 171 counts, respectively. It is reported in the literature that dimers of methanol (CH₃OH)₂ with IE equal to 9.72 eV coexist with methanol monomers under ambient conditions and that protonated methanol is generated from the dissociation of (CH₃OH)₂⁺. Figure 2(B) shows the LPPI mass spectrum of 4.6 ppmv methanol doped with 291 ppmv CH₂Cl₂. The signal intensities of the mass peaks at m/z 33 and 65 reach 1.48 × 10⁵ and 6.06 × 10⁴ counts, respectively. The signal intensity of protonated methanol was averagely enhanced by 197.8 ± 18.8 times, measured from three independent measurements.

The IE of ethanol (C₂H₅OH) is 10.48 eV, meaning it can be photoionized by the photons emitted from the krypton lamp (10.6 eV, 20%). Figure 3(A) shows the LPPI mass spectrum of 1.6 ppmv ethanol in nitrogen. As well as ions resulting from water and water clusters, mass peaks at m/z 45, 47, and 93 are assigned to ions produced from ethanol, i.e. C₂H₅O⁺ (551 counts), (C₂H₅OH)H⁺ (1923 counts), and (C₂H₅OH)₂H⁺ (222 counts). The mass peak of protonated ethanol was the strongest peak. After doping with 291 ppmv CH₂Cl₂, the intensities of the

Figure 2. LPPI mass spectra of 4.6 ppmv methanol before (A) and after (B) doping with 291 ppmv CH₂Cl₂.

Figure 3. LPPI mass spectra of 1.6 ppmv ethanol before (A) and after (B) doping with 291 ppmv CH₂Cl₂.
mass peaks at m/z 47 and 93 shown in Fig. 3(B) increased to $1.61 \times 10^5$ and $2.21 \times 10^4$ counts, respectively. The signal intensity of protonated ethanol was averagely enhanced by 87.3 ± 7.8 times, measured from three independent measurements. The mass peak at m/z 45 slightly increased to 2765 counts, while the mass peaks at m/z 29 (1.54 × 10^4 counts) and 65 (1.80 × 10^4 counts) assigned to C_2H_5^+ and (C_2H_5OH·H_2O)H^+ appeared.

Acetaldehyde (C_2H_4O) is one of the most important aldehydes; it occurs widely in nature and is produced industrially on a large scale. The IE of acetaldehyde is 10.23 eV. Figure 4(A) shows the LPPI mass spectrum of 0.66 ppmv acetaldehyde in pure nitrogen. The mass peaks at m/z 45 and 61 are assigned to (C_2H_4O)H^+ (1290 counts), and (C_2H_3O·H_2O)^+ (1307 counts), respectively. The molecular ion of acetaldehyde was not observed. Protonated acetaldehyde was the dominant ion. Figure 4(B) shows the LPPI mass spectrum of 0.66 ppmv acetaldehyde in nitrogen doped with 291 ppmv CH_2Cl_2. The signal intensity of protonated acetaldehyde (m/z 45) increased to $7.04 \times 10^4$ counts, while the signal at m/z 61 slightly increased to 2107 counts. The signal intensity of protonated acetaldehyde was averagely enhanced by 93.5 ± 35.5 times, measured from three independent measurements. Additionally, a mass peak at m/z 63 assigned to (C_2H_4O·H_2O)H^+ (1.71 × 10^4 counts) appeared.

Benzene (C_6H_6) is an important chemical and atmospheric pollutant. Its IE is 9.24 eV, lower than the energy of VUV photons emitted from the krypton lamp. Benzene and its derivatives have excellent photoionization efficiencies under illumination of a krypton VUV lamp. Figure 5(A) shows the LPPI mass spectrum of 0.42 ppmv benzene in nitrogen. The mass peaks at m/z 37 and 78 shown in Fig. 5(B) increased to 7.81 × 10^4 and 1.79 × 10^4 counts, respectively. The signal intensity of protonated benzene was averagely enhanced by 51.3 ± 17.2 times, measured from three independent measurements. Additionally, a mass peak at m/z 79 assigned to (C_6H_5OH·H_2O)H^+ (2.56 × 10^3 counts) appeared.

![Figure 4. LPPI mass spectra of 0.66 ppmv acetaldehyde before (A) and after (B) doping with 291 ppmv CH_2Cl_2.](image)

![Figure 5. LPPI mass spectra of 0.42 ppmv benzene before (A) and after (B) doping with 291 ppmv CH_2Cl_2.](image)
A stainless steel fan driven by a magnetic field was used to decrease by ~14% to 5.54 × 10^4 counts. The fluctuation of the signal intensities at m/z 78 was observed in separate experiments. No signal enhancement at m/z 79 (protonated benzene) was observed in all experiments.

### Discussion

Pure CH₂Cl₂ in LPPI almost does not generate noticeable ions as shown in Fig. 1, which implies that the observed protonation enhancement is not attributed to charge exchange. In order to reveal the mechanism of protonation enhancement, the doping effects of H₂, CH₄, CCl₄, and CH₂Cl₂ on the intensities of nitrogen doped with 291 ppmv CH₂Cl₂. The intensities of the mass peak at m/z 78 decreased by ~14% to 5.54 × 10^4 counts. The fluctuation of the signal intensities at m/z 78 was observed in separate experiments. No signal enhancement at m/z 79 (protonated benzene) was observed in all experiments.

Table 1 lists IEs, PAs, molecular dipole moments, H-bond formation possibilities, and protonation enhancements of the compounds investigated.

Table 1. Ionization energies (IEs), proton affinities (PAs), molecular dipole moments, H-bond formation possibilities, and protonation enhancements of the compounds investigated. ahttp://webbook.nist.gov/. bhttp://www.kayelaby.npl.co.uk/. cObtained from three independent measurements.

| Compounds | Ionization Energy (eV) | Proton Affinity (kJ/mol) | Molecular Dipole Moment (C·m) | H-bond Formation Possibilities as H Acceptor | Protonation Enhancement by CH₂Cl₂ (times) |
|-----------|-----------------------|--------------------------|-------------------------------|---------------------------------------------|------------------------------------------|
| H₂O       | 12.62                 | 6917/22                  | 6.2                           | Yes                                        | 481.7 ± 122.4                            |
| CH₃OH     | 10.84                 | 7543/7.82                | 5.5                           | Yes                                        | 197.8 ± 18.8                             |
| C₂H₅OH    | 10.48                 | 7764/8.05                | 5.7                           | Yes                                        | 87.3 ± 7.8                               |
| C₃H₆O     | 10.23                 | 7685/7.97                | 8.3                           | Yes                                        | 93.5 ± 35.5                              |
| C₂H₅H₂O   | 9.24                  | 7504/7.78                | 0                             | No                                         | 0                                        |
| CH₂Cl₂    | 11.33                 | 628/6.51                 | 6.0                           | No                                         | 0                                        |

benzene. The mass peak at m/z 78 is assigned to [12C₆H₆]+ (6.42 × 10^4). Figure 5(B) shows the LPPI mass spectrum of 0.42 ppmv benzene in nitrogen doped with 291 ppmv CH₂Cl₂. The intensities of the mass peak at m/z 78 decreased by ~14% to 5.54 × 10^4 counts. The fluctuation of the signal intensities at m/z 78 was observed in separate experiments. No signal enhancement at m/z 79 (protonated benzene) was observed in all experiments.

#### Method

The experimental setup has been described in detail elsewhere. Briefly, it mainly consisted of a 120 L test chamber and a LPPI mass spectrometer.

The 120 L test chamber was mainly built with an open-head stainless steel drum and covered with a thin Tedlar bag to ensure one atmospheric pressure during sampling. A stainless steel fan driven by a magnetic field was...
μ1.6, 0.66, and 0.42 ppmv, respectively. The injection of 100 and emitted VUV photons with energies of 10.0 eV (~80%) and 10.6 eV (~20%). The sample gas was introduced into the photoionization source and controlled by a needle valve. The sample flow was maintained at ~1 cm³ s⁻¹. The pressure in the photoionization source was 500–1000 Pa. The mass spectrometer was a simple V-shaped time-of-flight mass spectrometer with a free-field flight distance of 460 mm. The cycle time of detection was 10 s.

In the experiments, a small amount of bottle-contained chemical was first injected into the test chamber. Then, 100 µL CH₂Cl₂ was injected into the test chamber and the mass spectra were subsequently acquired after each injection. The amount of methanol, ethanol, acetaldehyde, and benzene injected into the nitrogen-filled test chamber was 1.0, 0.5, 0.5, and 0.2 µL, respectively. The resulting mixing ratios for the prepared gases were 4.6, 1.6, 0.66, and 0.42 ppmv, respectively. The injection of 100 µL CH₂Cl₂ resulted in 291 ppmv in the mixing ratio.

In this study, methanol (A. R., Sinopharm), ethanol (A. R., Sinopharm), acetaldehyde (40% in water, Sinopharm), benzene (A. R., Beijing Shiji), CH₂Cl₂ (HPLC grade, Cleman Chemical), CHCl₃ (A. R., Beijing Shiji), and CCl₄ (A. R., Sinopharm) were used. High-purity nitrogen (>99.99%), hydrogen (>99.999%), and methane (>99.9%) were purchased from Beijing Huayuan Gas Co., Ltd.

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**Author Contributions**
J.S. conceived the idea, conducted the experiment, and wrote the paper. Y.Z., C.X., and Z.L. conducted the experiment; W.S. did the preliminary study which indicated the possibility of protonation enhancement. B.Y. and X.Z. were involved in preparing the manuscript and supplying chemicals. P.Z. and P.M. managed the experimental setup.

**Additional Information**

**Competing financial interests:** The authors declare no competing financial interests.

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