Ultra-Fast Polarity Switching, Non-Radioactive Drift Tube for the Miniaturization of Drift-Time Ion Mobility Spectrometer

Lingfeng Li, Hao Gu, Yanzhen Lv, Yunjing Zhang, Xingli He and Peng Li *

School of Electronic and Information Engineering, Soochow University, Suzhou 215006, China; lingfengli@suda.edu.cn (L.L.); hguguhao@stu.suda.edu.cn (H.G.); lvyanzhen110@gmail.com (Y.L.); yjzhang1223@suda.edu.cn (Y.Z.); hexingli@suda.edu.cn (X.H.)

Abstract: Drift-time ion mobility spectrometer (DT-IMS) is a promising technology for gas detection and analysis in the form of miniaturized instrument. Analytes may exist in the form of positively or negatively charged ions according to their chemical composition and ionization condition, and therefore require both polarity of electric field for the detection. In this work the polarity switching of a drift-time ion mobility spectrometer based on a direct current (DC) corona discharge ionization source was investigated, with novel solutions for both the control of ion shutter and the stabilization of aperture grid. The drift field is established by employing a switchable high voltage power supply and a serial of voltage regulator diode, with optocouplers to drive the ion shutter when the polarity is switched. The potential of aperture grid is stabilized during the polarity switching by the use of four diodes to avoid unnecessary charging cycle of the aperture grid capacitor. Based on the proposed techniques, the developed DT-IMS with 50 mm drift path is able to switch its polarity in 10 ms and acquire mobility spectrum after 10 ms of stabilization. Coupled with a thermal desorption sampler, limit of detection (LoD) of 0.1 ng was achieved for ketamine and TNT. Extra benefits include single calibration substance for both polarities and largely simplified pneumatic design, together with the reduction of second drift tube and its accessories. This work paved the way towards further miniaturization of DT-IMS without compromise of performance.

Keywords: ion mobility spectrometer (IMS); polarity switching; non-radioactive

1. Introduction

DT-IMS, normally working under ambient pressure, is a sensitive technique for the detection of gas phase charged molecules based on their different mobility in the electric field [1]. It can either work as a standalone device [2–4] or in conjunction with other analytical instruments like Gas Chromatograph (GC) as its detector [5–7]. Its popularity largely attributed to the successful application in the detection of trace explosives, narcotics, Chemical Warfare Agents (CWA) and Toxic Industrial Compounds (TIC) [8–11]. According to the chemical composition the ionization process could result positive or negative ions, such as majority of nitro-based explosives are detected in the negative mode, and peroxide explosives including HMTD and TATP in the positive mode [12–16]. Other molecules can produce both positive and negative ions simultaneously such as methyl salicylate, for which measurement in both modes can further improve the accuracy of identification [17].

To characterize ions of both polarities, common methods is to use two drift tubes of each to be setup for one polarity, which clearly brings along increased volume, power consumption and system complexity [18–23]. An alternative way is to switch the polarity on a single drift tube to analyze positive and negative ions alternately, which is so called polarity switching DT-IMS. With this method it is an important that the switching is fast enough so that no information will be lost, especially when the concentration and identity of analyte vary over time. An example is the detection of TATP, which forms positive adducts.
ion and evaporates very quickly under the temperature condition suitable to thermally
desorb explosives. If the switching of polarity takes a few seconds and the device happens
to work under the negative mode when the TATP is sampled, the analyte will be gone when
the polarity is switched back to positive mode, ending up a false negative result. In another
application, when used as a detector for GC the switching also has to be fast as possible,
since the elution peak only last for seconds and each measurement must take shorter time
than the standard deviation to restore the gaussian peak with high fidelity [24].

To achieve fast switching of polarity, there are four major considerations for the
design of the instrument. The first is to establish uniform electric filed in the drift tube
after the change of voltage, which requires minimal capacitance between the electrodes
used to define such electric field [24–26]. Another benefit for smaller capacitance is the
reduction of power consumption, allowing the use of smaller power supply module for
high fast switching instrument. The second is concerning the aperture grid in front of the
faraday plate, which plays a key role for the elimination of mirror current effect on the ion
detector [24,26]. The potential difference of aperture grid and faraday plate is defined by
circuit in between, comprising a resistor and a capacitor connecting in parallel. The use of
capacitor is to suppress the noise on the faraday plate. If its capacitance is too small then
the circuit is prone to noise, and if too large the switching speed will be limited. The third
is the design of ion shutter to meet the requirement of polarity switching [27]. And finally,
when DC corona ionization source is used, extra complexity for the arrangement of electric
field in the ionization/reaction region will be introduced to the system comparing to that
of radioactive ionization source.

Zhou et al. reported the change of metal electrode from thin rings to thick ones and the
switching recovery time was shortened from 20 s to 2 s, attributing to the reduced parasitic
capacitance and charge accumulation, although 2 s seems still too long for the simultaneous
detection of compound with different charge polarity [28]. Pershenkov et al. proposed to
use an independent power supply for the aperture grid capacitor to reduce the discharge
time, which requires extra cost and volume of the instrument [26]. Hitzemann et al. used a
much smaller capacitor (220 pF) for the aperture grid, in combination with field switching
ion shutter, to achieve a polarity switching time of 12 ms on a test system based on
radioactive ionization source [24]. Although fast switching was achieved in this design,
small aperture grid capacitor leads to the problem of noise control and faraday plate on
HV end requires more sophisticated design of amplification circuit.

In this work, we have developed methods and implementation for an ultra-fast polarity
switching DT-IMS based on DC corona discharge ionization. With a special design of
driving circuit, only a single HV power supply is needed for both the ionization source
and drift tube. For the ion shutter, a Tyndall-Powell (TP) ion gate was used for its simple
structure, with the voltage change controlled by an optocoupler to synchronize with the
polarity change of the system. A novel circuit was proposed to work with the aperture grid,
eliminating charging cycle and therefore the need of extra power supply, yet allowing the
use of large capacitor to filter the noise at same time. Meanwhile, with this implementation
only one calibration substance is needed for both polarities, further reduced the complexity
and overall volume. Coupled to a thermal desorption sampler and using narcotics and
explosives as example analytes, the developed instrument was characterized for both
switching speed and analytical performance.

2. Experiment
2.1. Instruments and Reagents
2.1.1. Instruments

The DT-IMS system investigated is mainly comprised of sampler, ionization source,
ion shutter, drift tube and ion detector (faraday plate), as shown in Figure 1. The DC corona
discharge ionization source employs a needle-hole structure, with the distance between
the needle and counter electrode set to be 2 mm, the corona discharge voltage is 2500 V.
The counter electrode and G1 of ion shutter encompassed a reaction region with length of
25 mm and field strength of 30 V/mm. A TP ion gate is used as ion shutter with 0.5 mm as shutter width between G1 and G2, 0.18 ms as opening time, 30 V/mm as ion gate opening field strength, −60 V/mm as ion gate closing field strength, 50 mm as length of drift tube and 30 V/mm as field strength. The distance between G1 and the end of the reaction region is 1.17 mm, and the distance between G1 and the front of the drift tube is also 1.17 mm. The ring electrodes used for both reaction region and drift tube are 16 mm for inner diameter, 1.37 mm for thickness. The insulator rings are 1.17 mm for thickness and 18 mm for inner diameter, with slightly larger inner diameter to reduce the parasitic capacitance and also the charge accumulation on the surface. The end of the drift tube is an aperture grid, which has a distance of 1 mm from the faraday plate and an aperture voltage of 60 V. Confirmed by the measurement results, with this structure design the parasitic capacitance of the drift tube is reduced to 1.15 pF. The working temperature of the drift tube was set to 120 °C to comply with the 200 °C working temperature of the thermal desorption sampler for the detection of explosives and narcotics with low volatility. The gas used in the system is clean air filtered by active charcoal and 13x molecular sieve for both sample stream at 150 mL/min and drift flow at 200 mL/min. The dew point after purification is below −60 °C. To maintain the stability of discharge and consistency of ionization products, another gas flow of 700 mL/min is introduced as protection for the corona discharge ionization source.

Figure 1. Structure illustration of the developed DT-IMS.

A commercially available high voltage power supply module (Shanghai Taizhu, SMO series) is used to switch the polarity of applied voltage, with output voltage of ±6000 V and polarity switch time of 10 ms. The ion current is converted to voltage signal by a homemade transimpedance amplifier with bandwidth of 8 kHz and gain of 5G V/A. The mobility spectrums are collected through a data acquisition card (National Instruments, USB6210) with sampling rate of 100 kHz. The signal for polarity changes and operation of ion shutter are also provided by the output of data acquisition card.

Illustration of electric circuit and time sequence of driving signal are shown in Figure 2a, b, respectively. For the polarity switching of DT-IMS there are 3 applied voltages need to be reversed, including the ion source (U_{source}), the drift tube (U_{drift}) and G2 of the TP ion gate (U_{gate}). Normally DT-IMS with corona discharge ionization source requires at least 2 HV modules to serve the ionization source and drift tube separately. In order to simplify the system for further miniaturization, in this work a single HV module was used for the entire system, with applied voltage and time sequence as shown in Figure 2b for U_{source}, U_{drift} and U_{gate}, respectively.
To implement the voltage settings with time sequence described in Figure 2b with a single HV supplier, a novel electronic circuit was designed as shown in Figure 2a. The end electrode of drift tube is connected to common ground so that the Faraday plate is also around the ground, which reduces the critical requirements for signal amplification circuit. Apart from the resistors in serial connection as HV dividing network, a certain number of Zener diodes are inserted between the counter electrode of the ionization source and the ground to define the drift field and also avoid voltage drift of HV module. In this implementation 25 Zener diodes (Nexperia, BZB100) are used for ±2500 V voltage drop across the drift region, with a protection resistor of 100 MΩ connecting to the HV output. The corona needle is connected to HV output through a large resistor of 1 GΩ for limiting the discharge current, to get the discharge voltage towards the counter electrode.

The TF ion gate blocks the ions flux by setting up a reverse electric field between G1 and G2, with its driving circuit shown in Figure 2a. The trigger signal is applied on the MOSFET (PMV28ENEA) to control the open/close status of the ion shutter. An optocoupler (Infineon, PVA3055NPBF) is used to directly regulate the voltage of G2, so that the low voltage control signal can be separated from the HV. The voltages on both sides of the optocoupler are taken from the HV dividing network. When the optocoupler is closed, |Vc| is always smaller than |V1|, and therefore the electric field between G1 and G2 is in reverse direction to the drift region, the ion shutter is closed. On the other hand, when the optocoupler is open, exactly opposite situation is expected and the ion shutter is open. In this way the operation of the ion shutter worked with both polarities, with clear advantages over traditional half-bridge circuit, which cannot deal with the polarity switching due to the limitation that the gate voltage has to be higher than the drain voltage.

The purpose of aperture grid in IMS instrument is to prevent the generation of induced current as the ion swarm approaches the detector, so that the voltage stability of aperture grid is of key importance for the quality of signal measured on the detector [24]. Normally a large capacitor, which capacitance value is between 100 nF and 1 µF, is arranged between the aperture grid and ground, in parallel to the voltage dividing resistor, to stabilize the potential difference between the aperture grid and the ion detector and hence reduce the noise, as shown in Figure 3a. But when the polarity is switched, the charging/discharging of this large capacitor will largely slow down the establishment of desired voltage on the aperture grid. Here in this work, we use 4 switching diodes (BAV21W) to avoid the charging/discharging of the capacitor and yet to maintain a stable voltage on the aperture grid, as shown in Figure 3b. The circuit works as the following steps: when U_{drift} is at +HV, diode D2&D3 will be on and D1&D4 are off, so that with capacitor C node a is with ground and node b is with positive, the electric field within the capacitor is from b to a; when the polarity is switched and U_{drift} is at -HV, D2&D3 will be off and D1&D4 are on.

Figure 2. (a) Illustration of electric circuit; (b) time sequence of applied voltages.
on, node a is with negative voltage and node b is with ground, the electric field within the capacitor is still from b to a. Therefore, with this circuit the voltage of aperture grid can be established to the desired value without the charging/discharge of the capacitor, which is a key factor to realized ultra-fast polarity switching of the system. In the following sections the performance of this circuit was compared and analyzed against the traditional circuit.

![Figure 3. Aperture grid circuit: (a) traditional configuration; (b) this study.](image)

2.1.2. Reagents

HPLC grade acetonitrile was purchased from Tedia Co Ltd. (Fairfield, CA, USA). Standard explosive samples of TNT, PETN, HMX, RDX, HMTD and TATP with concentration of 1000 ng/μL were purchased from Beijing Shiji ‘aoke Biotechnology Co., Ltd. (Beijing, China). Standard narcotic samples of cocaine, ketamine, methamphetamine, heroin, morphine, and tetrahydrocannabinol with concentration of 1000 ng/μL were purchased from Shanghai Yuansi Standard Science and Technology Co., Ltd. (Shanghai, China). Molecular sieve 13x and activated charcoal were purchased from Sinopharm Chemical Reagent Co., Ltd. (Shanghai, China). Disposable Nomex swab was supplied by Suzhou Chuanche Specialty Materials Co., Ltd. (Suzhou, China).

2.2. Experimental Methods

2.2.1. Preparation of Standard Solution

Analyte solutions of explosives and narcotics were prepared by stepwise dilution of 1000 ng/μL standard solution to 100 ng/μL, 10 ng/μL, 5 ng/μL, 1 ng/μL and 0.1 ng/μL.

Mixture samples were prepared following these steps: the 1000 ng/μL standard solution was firstly diluted by 10 times to 100 ng/μL, the diluted solutions of chosen analytes were then mixed together with 100 μL of each, and finally the mixed solution was made up to 1 mL using acetonitrile to get 10 ng/μL concentration for each analyte.

2.2.2. Sampling Methods

For sample introduction 1 μL of sample solution was deposited on Nomex swab, which was then inserted into the sampling inlet of the thermal desorption unit after the solvent evaporated. The vapor of the analyte was then introduced into the DT-IMS system by the sample carrier gas flow. For each sample the test was repeated for at least 3 time to check the reproducibility.

3. Results and Discussion

3.1. Performance Evaluation of Polarity Switching DT-IMS

3.1.1. Comparison of Aperture Grid Circuits

Measured spectrums are shown in Figure 4, in comparison between the traditional aperture grid circuit and solution of this work. In both cases only Reactant Ion Peak (RIP)
was measured without sample introduction, with the ion shutter opened once every 20 ms and synchronized with the polarity switching. With traditional aperture grid circuit as shown in Figure 3a, the positive mode of IMS is maintained for 800 ms before switching to negative mode, with 1600 ms of data collected as shown in Figure 4a. From the results it can be observed that after the switching of HV which took 10 ms, the baseline needs around 600 ms to stabilize for valid measurement. With the proposed circuit as shown in Figure 3b, measured spectrum using 100 ms as switching interval are shown in Figure 4b. From the results, it only took 10 ms to get stable spectrum on top of another 10 ms for switching, demonstrating around 30 times improvement for the speed of polarity switching with the new approach.

![Figure 4. Aperture grid circuit: (a) traditional configuration; (b) this study.](image)

3.1.2. Mobility Spectrum with Fast Switching

A consecutive measurement of 40 ms of positive mode(red), 40 ms of negative mode(blue) and another 40 ms of positive mode(red) is shown in Figure 5. The experiment setting is similar, that only RIP was collected with 20 ms interval for ion shutter and synchronized polarity switch of 40 ms. It can be observed from the results that for the first opening of ion shutter after the change of polarity, baseline restoration and overload of the amplifier due to the HV switch exist, which made the collection of data invalid, as marked on the plot. However, for the second opening of ion shutter after switching the baseline is fully restored with expected RIP, as shown in Figure 5. Switching back to positive mode at 80 ms results identical spectrum as that of 0–40 ms, meaning the developed DT-IMS can work continuously with the ability to acquire full spectrum including both polarity within 80 ms.

![Figure 5. Effect of fast switching on the spectrum.](image)
To investigate the effect of polarity switching, measured spectrum with fast switching mode of 80 ms interval are put in comparison with results collected with single polarity under the same experimental conditions. For all three measurements, 5 ng of ketamine and TNT were used as analyte and the spectrum represented were all baseline corrected. From Figure 6 it can be observed that the drift time and intensity of both RIP and Product Ion Peak (PIP) are nearly identical, with maximum difference of peak intensity less than 3%. The results here demonstrates that fast polarity switching of the DT-IMS instrument develop in this work didn’t compromise the analytical performance.

Figure 6. Spectrum in comparison, measured under ultra-fast switching mode, positive mode and negative mode.

3.2. Polarity Switching DT-IMS for the Detection of Explosives and Narcotics

3.2.1. Effect of Switching Speed with Evaporated Samples

It is well known that detection of narcotics and explosives with IMS requires both polarities, since both positive and negative product ions are generated during the ionization process. Due to various volatility, typical sample introduction method is thermal desorption, with relatively high temperature to cover analytes with high boiling point such as HMX. The concentration of vapor phase analyte hence varied over time as shown in Figure 7 for 3 typical samples including ketamine, TNT and TATP. The measurements were carried out on the developed DT-IMS instrument in single polarity mode (positive mode for ketamine of 10 ng and TATP of 100 ng; negative mode for TNT of 10 ng) with 20 ms data collection interval and 200 °C for thermal desorption. All data was normalized for comparison, with the vertical axis representing relative concentration. It can be seen that for given conditions, the maximum concentration appears in large time window, from less than 0.5 s for TATP to around 5 s for TNT and ketamine.

Figure 7. Concentration profile over thermal desorption time of ketamine, TNT and TATP.

To examine the effect of polarity switching, an experiment was designed for the measurement of mixed sample comprising 100 ng of TATP and 10 ng of TNT, using the
developed DT-IMS instrument with different time and switching direction settings, as shown in Figure 8. Spectrums of three different switching period, 100 ms (a and b), 500 ms (c and d) and 1 s (e and f), were taken with both directions of switching including positive to negative (a, c and e) and negative to positive (b, d and f). From the results it can be observed that when the switching is fast enough, such as 100 ms or 10 Hz in this case, it doesn’t matter how the polarity is switched, analytes from both positive and negative modes can be detected with negligible difference. However, as the polarity switching got slower the results are substantially different until analyte from the second polarity mode cannot be detect with 1 s switching period. In combination of the concentration profile shown in Figure 7, it could be concluded that the variation of chemical information, including identity and concentration of analytes, can only be accurately captured by single drift tube DT-IMS with fast polarity switching capability.

3.2.2. Detection of Narcotics and Explosives

To further evaluate the instrument samples of six narcotics and seven explosives have been tested with switching time of 100 ms, example spectrums of 10 ng ketamine and 10 ng TNT are shown in Figure 9. Spectrums of other analytes can be found in Figures S1 and S2 of the supporting information.
Table 1. Reduced mobility of 6 common narcotics.

| No. | Sample   | Molecular              | Generated Ion     | Drift Time/ms | \(K_0/\text{cm}^2 \cdot \text{V}^{-1} \cdot \text{s}^{-1}\) (Measured) | \(K_0/\text{cm}^2 \cdot \text{V}^{-1} \cdot \text{s}^{-1}\) (Literature [31–33]) |
|-----|----------|------------------------|-------------------|---------------|---------------------------------------------------------------|--------------------------------------------------------------------------------|
| 1   | Cocaine  | C_{17}H_{21}NO_{4}     | [M+H]^+          | 11.90         | 1.170                                                         | 0.003                                                                           |
| 2   | Ketamine | C_{19}H_{20}NOCl       | [M+H]^+          | 10.34         | 1.348                                                         | 0.002                                                                           |
| 3   | Methamphetamine | C_{19}H_{20}N   | [M+H]^+          | 9.05          | 1.536                                                         | 0.007                                                                           |
| 4   | Heroin   | C_{21}H_{23}NO_{5}     | [M+H–CH_{3}COOH]^+ | 11.91        | 1.169                                                         | 0.002                                                                           |
| 5   | Morphine | C_{17}H_{20}NO_{3}     | [M+H–H_{2}O]^+   | 10.82         | 1.253                                                         | 0.027                                                                           |
| 6   | Marijuana| C_{21}H_{30}O_{2}      | [M+H]^+          | 11.88         | 1.172                                                         | 0.002                                                                           |

Figure 9. Example IMS spectrums: (a) 10 ng ketamine; (b) 10 ng TNT.

3.2.3. Validation of Calibration Method

For traditional DT-IMS with separate drift tubes, normally two calibration substances are needed to compensate the variation and inconsistency of the drift field, since the variation of voltage is independent for each polarity. However, with the polarity switching DT-IMS developed in this work, because the voltage across the drift tube is defined by the Zener diode for both polarities, in principle it is possible to use a single calibration substance.

Validation of the hypnosis was carried out with experiments using niacinamide, a common positive mode calibrator for IMS instrument with well recognized mobility, to calculate the reduced mobility of narcotic and explosive samples in comparison with the values reported in the literature [29,30]. The reduced mobility of niacinamide was reported to be 1.860 \(\text{cm}^2 \cdot \text{V}^{-1} \cdot \text{s}^{-1}\), so in the system with niacinamide as calibrator, reduced mobility of other sample in the same mode can be calculated as:

\[
K'_0 = \frac{K_0 \cdot t}{t'}
\]  

(1)

where \(K'_0\) is the reduced mobility of analyte of concern, \(K_0\) is the reduced mobility of the calibrator, \(t'\) and \(t\) are the drift time for analyte and calibrator, respectively. Spectrum data of narcotic and explosive samples from Section 3.2.2 was used to calculate the reduced mobility, which was summarized in Tables 1 and 2.
Table 2. Reduced mobility of 7 common explosives.

| No. | Sample     | Molecular Formula | Generated Ion | Drift Time/ms | \( K_0/\text{cm}^2\cdot\text{V}^{-1}\cdot\text{s}^{-1} \) (Measured) | \( K_0/\text{cm}^2\cdot\text{V}^{-1}\cdot\text{s}^{-1} \) (Literature [31,34,35]) |
|-----|------------|-------------------|---------------|--------------|-----------------------------------------------------------------|-----------------------------------------------------------------|
| 1   | TNT        | \( C_7H_5N_3O_6 \) | \([M-H]^–\)   | 9.18         | 0.006                                                          | 1.519                                                          | 0.001                                                          | 1.53                                                          |
| 2   | Black Powder | S                | \([S]^-\)     | 6.47         | 0.010                                                          | 2.147                                                          | 0.006                                                          | 2.26                                                          |
| 3   | PETN       | \( C_6H_4N_2O \)  | \([M+NO_2]^–\) | 11.90        | 0.012                                                          | 1.171                                                          | 0.002                                                          | 1.19                                                          |
| 4   | HMX        | \( C_4H_8N_4O_8 \) | \([M+NO_2]^–\) | 10.86        | 0.010                                                          | 1.282                                                          | 0.002                                                          | 1.28                                                          |
| 5   | RDX        | \( C_3H_6N_6O_6 \) | \([M+NO_2]^–\) | 9.84         | 0.020                                                          | 1.415                                                          | 0.003                                                          | 1.42                                                          |
| 6   | HMTD       | \((\text{CH}_2\text{N}_2\text{O}_2)_3\) | \([M+H]^+\)   | 11.38        | 0.150                                                          | 1.225                                                          | 0.016                                                          | 1.55                                                          |
| 7   | TATP       | \( C_9H_{18}O_6 \) | \( [(\text{CH}_3)_2\text{C(O)OO}]^+\) | 6.60         | 0.012                                                          | 2.120                                                          | 0.023                                                          | 2.15                                                          |

As shown in Tables 1 and 2, majority of the analytes tested demonstrated good consistency of measured \( K_0 \) to that reported in the literature, with HMTD as the only exception for reasons to be further investigated [31–35]. This result, the successful mobility correction of both polarities using a single calibrator with only positively charged product ion, proves the feasibility of not only to reduce a drift tube from the system, but also to eliminate the need for the second calibrator with associated container and supporting accessories, which is a substantial step for further miniaturization of the DT-IMS instrument. The underlying reason can be ascribed to the stable and consistent drift field of both working mode. In the developed system the voltage across the drift tube is defined by the bidirectional voltage regulator diodes for both modes, and since a single drift tube was used there will be no size inconsistency either from the manufacturing or elongated working, therefore the electric field for ion drifting will be strictly the same for either polarity, enabling mobility correction with a single calibrator.

3.2.4. Detection of Mixed Samples

With the hardware performance established such as switching speed and mobility accuracy, the polarity switching DT-IMS is tested with mixed samples of narcotics and explosives for the examination of detection capability. The mixed samples are prepared as described in the experiment section and tested with 100 ms switching time, as shown in Figure 10. Each prepared mixed sample intentionally include analytes of both polarities for demonstration. From the results it can be seen that with the current instrument all of the narcotic and explosive we had can be successfully detected, regardless of polarity and volatility.

![Figure 10](image-url). Measured spectrums of mixed samples: (a) 10 ng TNT and 10 ng ketamine; (b) 10 ng TNT and 100 ng TATP; (c) 100 ng HMX and 100 ng TATP; (d) 10 ng RDX and 100 ng HMTD.
3.2.5. Limit of Detection

LoD of the developed instrument is also investigated with the standard of signal-to-noise ratio greater than 3. Commercial instruments normally claim LoD of 1 ng or even lower [29]. In this work two representative analytes, ketamine and TNT, are chosen to characterize the LoD by stepwise dilution. As shown in Figure 11 for both analytes 0.1 ng can be reliably detected.

![Figure 11](image_url)

Figure 11. Example IMS spectrums of (a) 0.1 ng ketamine; (b) 0.1 ng TNT.

4. Conclusions

In this work an ultra-fast polarity switching IMS drift tube with non-radioactive ionization source was designed and implemented, employing novel methods on the electronic design to switch the aperture grid and ion shutter. Stable spectrum can be measured within 20 ms after the change of polarity. Testing results demonstrated that the developed instrument can effectively detect common narcotics and explosives regardless of ion polarity and volatility. The LoD was lower than 0.1 ng and only a single calibrator was needed for mobility correction for both positive and negative modes. The proposed electronics and system design paved the path for the future miniaturization of DT-IMS, which can either be developed into a miniaturized hand-held instrument, or a sensor to be used for GC and other separation technologies.

Some limitations of this work still exist:
1. The delay of optocoupler which was used to control the TP ion gate could possibly introduce systematic errors for the measurement of mobility.
2. The power consumption of the DC corona discharge ionization source is relatively large.
3. The integration of the system needs to be improved.

The future work we have planned includes optimization of the ion gate, replacing DC corona ion source with AC, and integration of not only the drift tube but also other peripherals such as flow control components etc.

Supplementary Materials: The following supporting information can be downloaded at: https://www.mdpi.com/article/10.3390/s22134866/s1, Figure S1: Photos of the drift tube device; Figure S2: Example IMS spectrums of 5 narcotics: (a) 10 ng cocaine; (b) 50 ng methamphetamine; (c) 50 ng heroin; (d) 50 ng morphine; (f) 50 ng marijuana; Figure S3: Example IMS spectrums of 6 explosives: (a) 50 ng black powder; (b) 50 ng PETN; (c) 100 ng HMX; (d) 50 ng RDX; (f) 100 ng HMTD; (g) 100 ng TATP.

Author Contributions: Conceptualization, L.L. and P.L.; methodology, L.L.; validation, H.G. and Y.L.; formal analysis, X.H. and Y.Z.; investigation, L.L., H.G. and Y.L.; resources, P.L.; data curation, L.L.; writing—original draft preparation, L.L., H.G. and Y.L.; writing—review and editing, X.H., Y.Z. and P.L.; visualization, H.G. and Y.L.; supervision, P.L.; project administration, P.L.; funding acquisition, P.L. All authors have read and agreed to the published version of the manuscript.
**Funding:** This work was supported by the National Natural Science Foundation of China (No. 21904091 and 62001318), Natural Science Foundation of Jiangsu Province (No. BK20200876), and Chinese Postdoctoral Science Foundation (No. 2020M671578).

**Institutional Review Board Statement:** Not applicable.

**Informed Consent Statement:** Not applicable.

**Data Availability Statement:** The data presented in this work are available in the article and Supplementary Materials.

**Conflicts of Interest:** The authors declare no conflict of interest.

**References**

1. Eiceman, G.A.; Karpas, Z. *Ion Mobility Spectrometry*, 3rd ed.; CRC Press: Boca Raton, FL, USA, 2013.

2. Baumbach, J.; Berger, D.; Leonhardt, J.; Klockow, D. Ion mobility sensor in environmental analytical chemistry—Concept and first results. *Int. J. Environ. Anal. Chem.* 1993, 52, 189–193. [CrossRef]

3. Bocos-Bintintan, V.; Thomas, C.P.; Ratiu, I.A. Sensors’ array of aspiration ion mobility spectrometer as a tool for bacteria discrimination. *Talanta* 2020, 206, 120233. [CrossRef] [PubMed]

4. Kirk, A.T.; Zimmermann, S. Ionenmobilitätspektrometrie. *Chem. Unserer Zeit* 2016, 50, 5. [CrossRef]

5. Aguilera-Herrador, E.; Cárdenas, S.; Ruizsanyi, V.; Sielemann, S.; Valcárcel, M. Evaluation of a new miniaturized ion mobility spectrometer and its coupling to fast gas chromatography multicapillary columns. *J. Chromatogr. A* 2008, 1214, 143–150. [CrossRef]

6. Ahrens, A.; Zimmermann, S. Towards a hand-held, fast, and sensitive gas chromatograph-ion mobility spectrometer for detecting volatile compounds. *Anal. Bioanal. Chem.* 2021, 413, 1009–1016. [CrossRef] [PubMed]

7. Langejuergen, J.; Wagner, C.; Beutel, S.; Schepner, T.; Zimmermann, S. Non-invasive monitoring of bacterial growth and autono-induced protein production in a bioreactor with a closed-loop GC-IMS. *Int. J. Ion Mobil. Spectrom.* 2015, 18, 9–15. [CrossRef]

8. Borsdorf, H.; Mayer, T. Response of halogenated compounds in ion mobility spectrometry depending on their structural features. *Talanta* 2011, 83, 815–822. [CrossRef]

9. Ewing, R.G.; Atkinson, D.A.; Eiceman, G.; Ewing, G. A critical review of ion mobility spectrometry for the detection of explosives and explosive related compounds. *Talanta* 2001, 54, 515–529. [CrossRef]

10. Utriainen, M.; Kärpänoja, E.; Paakkanen, H. Combining miniaturized ion mobility spectrometer and metal oxide gas sensor for the fast detection of toxic chemical vapors. *Sens. Actuators B Chem.* 2003, 93, 17–24. [CrossRef]

11. Yamaguchi, S.; Asada, R.; Kishi, S.; Sekioka, R.; Kitagawa, N.; Tokita, K.; Yamamoto, S.; Seto, Y. Detection performance of a portable ion mobility spectrometer with 63Ni radioactive ionization for chemical warfare agents. *Forensic Toxicol.* 2010, 28, 84–95. [CrossRef]

12. Drees, C.; Vautz, W.; Liedtke, S.; Rosin, C.; Althoff, K.; Lippmann, M.; Zimmermann, S.; Legler, T.J.; Yildiz, D.; Perl, T. GC-IMS headspace analyses allow early recognition of bacterial growth and rapid pathogen differentiation in standard blood cultures. *Appl. Microbiol. Biotechnol.* 2019, 103, 9091–9101. [CrossRef] [PubMed]

13. Kanu, A.B.; Hill, H.H.; Jr. Ion mobility spectrometry detection for gas chromatography. *J. Chromatogr.* 2008, 1177, 12–27. [CrossRef] [PubMed]

14. Oxley, J.C.; Smith, J.L.; Luo, W.; Brady, J. Determining the vapor pressures of diacetone diperoxide (DADP) and hexamethylene triperoxide diamine (HMTD). *Propellants Explos. 2009*, 34, 539–543. [CrossRef]

15. Puton, J.; Nousiainen, M.; Sillanpää, M. Ion mobility spectrometers with doped gases. *Talanta* 2008, 76, 978–987. [CrossRef] [PubMed]

16. Schulte-Ladbeck, R.; Edelmann, A.; Quintas, G.; Lendl, B.; Karst, U. Determination of peroxide-based explosives using liquid chromatography with on-line infrared detection. *Anal. Chem.* 2006, 78, 8150–8155. [CrossRef] [PubMed]

17. Lippmann, M.; Kirk, A.T.; Hitzemann, M.; Zimmermann, S. Compact and Sensitive Dual Drift Tube Ion Mobility Spectrometer with a New Dual Field Switching Ion Shutter for Simultaneous Detection of Both Ion Polarities. *Anal. Chem.* 2020, 92, 11834–11841. [CrossRef]

18. Atkinson, J.R.; Clark, A.; Taylor, S.J. Ion Mobility Spectrometer Comprising Two Drift Chambers. U.S. Patent 7,994,475, 9 August 2011.

19. Machlinski, K.J.; Pompeii, M.A.; Johnson, G.P.; Fitzgerald, R.A.; Byrne, J.A. (Chemical Agent) Point Detection System (IPDS) employing Dual Ion Mobility Spectrometers. U.S. Patent 6,627,878, 30 September 2003.

20. Scott, J.R.; Dahl, D.A.; Miller, C.J.; Tremblay, P.L.; McJunkin, T.R. Dual Mode Ion Mobility Spectrometer and Method for Ion Mobility Spectrometry. U.S. Patent 7,259,369, 21 August 2007.

21. Spangler, G.E.; Wroten, J.F., Jr. Apparatus for Simultaneous Detection of Positive and Negative Ions in Ion Mobility Spectrometry. U.S. Patent 4,445,038, 24 April 1984.

22. Zimmermann, S.; Kirk, A.; Lippmann, M.; Bohnhorst, A. Ion Mobility Spectrometer and Method for Analyzing Samples by Ion Mobility Spectrometry. U.S. Patent 11,293,899, 5 April 2022.
23. Shaltaeva, Y.; Golovin, A.; Vasilyev, V.; Gromov, E.; Matusko, M.; Malkin, E.; Ivanov, I.; Belyakov, V.; Pershenkov, V. The Review of Bipolar Ion Mobility Spectrometers. In Proceedings of the International Conference on Nanotechnologies and Biomedical Engineering, Chisinau, Moldova, 18 September 2019.

24. Hitzemann, M.; Kirk, A.T.; Lippmann, M.; Bohnhorst, A.; Zimmermann, S. Miniaturized Drift Tube Ion Mobility Spectrometer with Ultra-Fast Polarity Switching. *Anal. Chem.* **2022**, *94*, 777–786. [CrossRef]

25. Gromov, E.; Matusko, M.; Shaltaeva, Y.; Pershenkov, V.; Belyakov, V.; Golovin, A.; Malkin, E.; Ivanov, I.; Vasilyev, V. Dual Mode Ion Mobility Spectrometer High Voltage Formation Circuit. In Proceedings of the IEEE 31st International Conference on Microelectronics (MIEL), Nis, Serbia, 16–18 September 2019.

26. Pershenkov, V.; Belyakov, V.; Shaltaeva, Y.; Malkin, E.; Golovin, A.; Ivanov, I.; Vasilyev, V.; Matusko, M.; Gromov, E. Fast Switching of the Polarity of Dual Mode Ion Mobility Spectrometer. In Proceedings of the IEEE 31st International Conference on Microelectronics (MIEL), Nis, Serbia, 16–18 September 2019.

27. Chen, C.; Tabrizchi, M.; Li, H. Ion gating in ion mobility spectrometry: Principles and advances. *Trends Anal. Chem.* **2020**, *133*, 116100. [CrossRef]

28. Zhou, Q.; Peng, L.; Jiang, D.; Wang, X.; Wang, H.; Li, H. Detection of nitro-based and peroxide-based explosives by fast polarity-switchable ion mobility spectrometer with ion focusing in vicinity of Faraday detector. *Sci. Rep.* **2015**, *5*, 10659. [CrossRef]

29. Armenta, S.; Esteve-Turrillas, F.; Alcala, M. Analysis of hazardous chemicals by “stand alone” drift tube ion mobility spectrometry: A review. *Anal. Methods* **2020**, *12*, 1163–1181. [CrossRef]

30. Sorribes-Soriano, A.; Esteve-Turrillas, F.A.; Armenta, S.; Amorós, P.; Herrero-Martínez, J.M. Amphetamine-type stimulants analysis in oral fluid based on molecularly imprinting extraction. *Anal. Chim. Acta* **2019**, *1052*, 73–83. [CrossRef] [PubMed]

31. Du, Z.; Sun, T.; Zhao, J.; Wang, D.; Zhang, Z.; Yu, W. Development of a plug-type IMS-MS instrument and its applications in resolving problems existing in in-situ detection of illicit drugs and explosives by IMS. *Talanta* **2018**, *184*, 65–72. [CrossRef] [PubMed]

32. Liu, H.; Xia, L.; Shen, C.; Huang, C.; Chu, Y. Dopant for detection of methamphetamine in the presence of nicotine with ion mobility spectrometry. *Anal. Bioanal. Chem.* **2021**, *413*, 4237–4246. [CrossRef] [PubMed]

33. Reese, E.S.; de Harrington, P. The analysis of methamphetamine hydrochloride by thermal desorption ion mobility spectrometry and SIMPLISMA. *J. Forensic Sci.* **1999**, *44*, 68–76. [CrossRef]

34. Crawford, C.L.; Boudries, H.; Reda, R.J.; Roscioli, K.M.; Kaplan, K.A.; Siems, W.F.; Hill, H.H., Jr. Analysis of Black Powder by Ion Mobility—Time-of-Flight Mass Spectrometry. *Anal. Chem.* **2010**, *82*, 387–393. [CrossRef]

35. Jiang, D.; Peng, L.; Wen, M.; Zhou, Q.; Chen, C.; Wang, X.; Chen, W.; Li, H. Dopant-assisted positive photoionization ion mobility spectrometry coupled with time-resolved thermal desorption for on-site detection of triacetone triperoxide and hexamethylene trioxide diamine in complex matrices. *Anal. Chem.* **2016**, *88*, 4391–4399. [CrossRef]