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A selector for structural isomers of neutral molecules

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We have selected and spatially separated the two conformers of 3-aminophenol (C6H7NO) present in a molecular beam. Analogous to the separation of ions based on their mass-to-charge ratios in a quadrupole mass filter, the neutral conformers are separated based on their different mass-to-dipole-moment ratios in an ac electric quadrupole selector. For a given ac frequency, the individual conformers experience different focusing forces, resulting in different transmissions through the selector. These experiments demonstrate that conformer-selected samples of large molecules can be prepared, offering new possibilities for the study of gas-phase biomolecules.

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During the last decades, the properties of biomolecules in the gas phase have been studied in ever greater detail [1–3]. Although the study of biomolecules outside of their natural environment was met with skepticism in the beginning, spectroscopic studies on isolated species in a molecular beam have proven to be very powerful to understand their intrinsic properties. Moreover, their native environment can be mimicked by adding solvent molecules one by one. These studies on well-defined biomolecular systems are particularly relevant to benchmark theoretical calculations. Even in the cold environment of a molecular beam, biomolecules exist in various conformational structures. The existence of multiple conformers (structural isomers) has been observed in the study of jet-cooled glycine for the first time [4] and in numerous experiments since then. In many cases, the individual conformers are identified via their different electronic spectra [5, 6]. This has been exploited in multiple-resonance techniques to measure, for instance, conformer-specific infrared spectra from which the conformational structures can be deduced [7, 8]. Apart from this information on the local minima on the potential energy surface, information on the barriers separating the conformers has been obtained in sophisticated multiple-resonance experiments as well [9].

The preparation of conformer-selected samples of biomolecules would enable a new class of experiments to be performed on these systems, e.g., electron and X-ray diffraction [10, 11] and tomographic imaging experiments [12]. Also, ultrafast dynamics studies on the ground-state potential energy surface would benefit from the availability of these pure samples. For charged species the separation of structurally different molecules has been demonstrated using ion-mobility in drift tubes [13, 14]. For neutral molecules, no such separation method exists. It has been demonstrated that the abundance of the conformers in the beam can be partly influenced by selective over-the-barrier excitation in the early stage of the expansion [15] or by changing the carrier gas [16]. These methods, however, are not generally applicable nor able to specifically select each of the conformers.

Polar molecules experience a force in an inhomogeneous electric field given by the negative gradient of the Stark energy. If the molecule is in an eigenstate whose Stark energy increases with increasing electric field, a so-called low-field-seaking (lfs) state, it feels a force towards regions of low electric field. Molecules in lfs states can be focused using static inhomogeneous electric fields. This has been used, for example, to create the population inversion that was essential for the demonstration of the MASER [17] and for state-selection of small molecules for scattering experiments. Furthermore, small molecules in lfs states have been slowed down and trapped using time-varying electric fields [18, 19]. Large molecules have a high density of rotational states, and due to the interaction between these states all of them are high-field seeking (hfs), i.e., they feel a force towards regions of high electric field. Molecules in hfs states can be dynamically focused using the alternating gradient principle [20]. This has been demonstrated for ammonia in hfs states [21, 22]. It has also been applied in the deceleration of CO [23, 24], YbF [25], and benzonitrile [26] molecules in hfs states, as well as in the ac trapping of ND3 [27].

The conformers of a specific biomolecule all have the same mass m and the same connectivities between the atoms (constitu- tion) but differ by the orientations of their functional groups in the molecular frame, i.e., by their folding pattern. The vectorial sum of the local dipole moments of the functional groups largely determines the overall dipole moment of the molecule. The different dipole moments µ of the conformers can be exploited to select individual conformers using dynamic focusing with ac electric fields. This is most easily implemented in a setup using high voltage electrodes in a quadrupole arrangement around a molecular beam. The operation principle of such an m/µ-selector is equivalent to that of the m/q quadrupole mass filter for charged particles, where q is the charge [32]. In the m/µ-selector the most polar quantum states of a given conformer are focused most efficiently and have the highest transmission. These are also the quantum states that can be aligned or oriented best, using intense laser fields or strong static electric fields, respectively. Therefore, experiments that rely on highly oriented samples, such as tomographic or diffraction imaging experiments, would particularly benefit from the conformer-selected polar ensembles generated by the m/µ-selector.

In this Letter, we demonstrate the selective transmission of the cis and trans conformers of neutral 3-aminophenol (C6H7NO) through an ac quadrupole m/µ-selector. The cis
and trans conformers of 3-aminophenol are used here as prototypes for the different structural isomers of biomolecules. The rotational envelopes of the electronic origin transitions of the individual conformers provide information on the rotational state specific transmission of the device.

The experimental setup is shown in Fig. 1a. A sample of 3-aminophenol is heated to 110 °C and co-expanded in 2 bar of Kr through a pulsed nozzle operated at a repetition rate of 10 Hz. The mean velocity of the molecules in the beam is approximately 480 m/s with a velocity spread (full-width at half-maximum) of about 10 %. After passing two skimmers, placed 5 cm and 15 cm downstream from the nozzle, the molecules enter a second, differentially pumped vacuum chamber, in which the m/µ-selector is placed. The selector consists of four polished, 1 m long cylindrical stainless-steel electrodes of 4 mm diameter. High voltages of 12 kV against ground are applied as shown in Fig. 1b. The gaps are 0.9 mm between two adjacent electrodes and 3 mm between two opposing electrodes, resulting in a field strength of 45 kV/cm on the centerline and a maximum field strength of 135 kV/cm. Using three high-voltage switches, the field is switched rapidly (≪ 1 µs) between two electric field configurations shown in Fig. 1b. Switching from one configuration to the other interchanges the role of the x and y axes, i.e., it interchanges the directions of transverse focusing and defocusing as indicated by the white arrows in Fig. 1b. The resulting dynamic focusing of neutral molecules, which is very similar to the dynamic focusing of charged particles [28], has been described in detail elsewhere [20, 24].

The transmitted 3-aminophenol molecules are ionized 1.21 m downstream from the nozzle using two-color resonance-enhanced multi-photon ionization, (1+1′)-REMPI, as shown in Fig. 1c. Prior to entering the detection region, the molecules have to pass through a 2 mm diameter aperture positioned on the centerline of the selector. The axis of the m/µ-selector is tilted against the axis of the incoming molecular beam by 0.3 °, such that predominantly molecules that are transported through the selector enter the detection region. The laser beams used for excitation and ionization are unfocused, and have diameters of 4 mm and 2 mm, respectively. To minimize saturation effects, the energy of the frequency-doubled pulsed dye-laser for electronic excitation (290 nm) is reduced to 20 µJ/pulse, while the ionization laser (355 nm) is operated with 5 mJ/pulse. The ions are mass-selectively detected in a time-of-flight setup. The cis and trans conformers can be selectively detected due to their distinct frequencies, and have diameters of 4 mm and 2 mm, respectively.

From the precisely known rotational constants and dipole moments [29] the energies of the rotational states of cis-3-aminophenol and trans-3-aminophenol are calculated as a function of electric field strength. Fig. 2 shows the resulting Stark curves for the lowest rotational states of both species. The transmission characteristics of the selector depend on the effective dipole moment µ eff (the negative of the slope of the Stark curve), the electric field gradients, and the ac frequency. Similar to the frequency dependence in quadrupole mass-spectrometers, molecules with a given value of µ eff are only transmitted through the selector within a finite range of frequencies. At too low frequencies molecules are deflected and lost in one transverse dimension before they are refocused. For high frequencies the time-averaged potential becomes flat resulting in a strongly reduced transmission. The ac frequency for optimum transmission increases with increasing µ eff. When a constant ac frequency is applied, a µ eff selection is performed.

AC frequency scans for cis-3-aminophenol and trans-3-aminophenol are shown in Fig. 3. The transmission measurements are performed with the excitation laser frequency set close to the band origin of the respective conformer. The ac frequency is scanned from 0 kHz to 5 kHz in steps of 50 Hz.
For a given conformer, the number of transmitted molecules is measured with and without applied high voltages, and the ratio of these two measurements is plotted in Fig. 3. The start-phase of the switching cycle determines the overall transmission of the selector. In all measurements we start with a half-period of focusing along the horizontal axis (configuration 1). As a consequence, the phase of the switching cycle at the moment that the molecules exit the selector changes with the applied frequency. For cis-3-aminophenol a clear enhancement of the transmission is observed for ac frequencies in the range from 2–3.5 kHz, whereas for trans-3-aminophenol a weaker transmission maximum is observed around 1.5 kHz. Both, the higher frequency and the higher transmission for cis-3-aminophenol reflect its considerably larger dipole moment compared to that of the trans conformer. The central dip in the transmission curve at 2.7 kHz is due to effects of the exact phase of the ac switching cycle at the exit of the selector. This phase determines the shape of the molecular packet in the detection region, and thereby its overlap with the laser beams.

As discussed above, the transmission of the selector depends on the effective dipole moment $\mu_{\text{eff}}$ of the individual quantum states. Although we cannot detect individual rotational states, there is a well-defined relation between the excitation laser frequency and the rotational states that are probed. In Fig 4 the (1+1')-REMPI spectrum of 3-aminophenol is shown. In the center of each of the vibronic bands, predominantly transitions from low-$J$ states are probed, whereas the wings of the rotational envelopes contain mostly transitions from high-$J$ states. The inset of Fig 4 shows the rotational contour of the origin transition of cis-3-aminophenol on an enlarged wavenumber scale measured with and without electric fields, for different ac frequencies. For a frequency of 1.6 kHz the wings of the rotational envelope are increased, whereas the intensity of the central part of the band is actually decreased. This directly reflects that for this ac frequency high-$J$ states are efficiently transported through the selector, whereas low-$J$ states, that generally have a larger $\mu_{\text{eff}}$, are over-focused and have a lower transmission. For a frequency of 2.95 kHz the whole rotational envelope is clearly increased. The largest enhancement is now observed for the central part of the rotational envelope, where mostly low-$J$ states are probed. For trans-3-aminophenol similar, albeit weaker, changes in the rotational envelope are observed.

Monte-Carlo trajectory calculations are performed to simulate the transmission curves for fixed excitation laser frequencies. For this, the rotationally resolved electronic excitation spectrum is calculated using the known rotational constants and transition moment orientations [29, 30]. A rotational temperature of 4 K yields a rotational envelope that agrees best with the observations. A rectangular spectral profile of the laser with a width of 0.15 cm$^{-1}$ is assumed. For all rotational states that are probed within this bandwidth of the laser, Monte-Carlo simulations are performed, and individual transmission curves are calculated. From the calculated line-strengths and populations, a weight for every single quantum state is determined. The weighted sum over the individual transmission curves is shown together with the experimental data in Fig 3 (dashed lines). These simulations nicely reproduce the peak position and the low-frequency cut-off of the experimental transmission curves. On the high-frequency side the experimentally observed transmission decreases faster than predicted, which we attribute to mechanical misalignment. Taking into account the phase-dependent shape of the molecular packet and its spatial overlap with the
detection laser beams, the observed modulation of the transmission peak for cis-3-aminophenol is correctly reproduced (dotted line).

In the experiments presented here, the selector is operated under conditions for optimum transmission, equivalent to the “rf-only” operation mode of quadrupole mass filters. The resolution $\mu_{\text{eff}}/\Delta \mu_{\text{eff}}$ of the selector is only about two in this case. In $m/q$ filters, the resolution $m/\Delta m$ is increased by adding a DC offset to the rf potentials, at the cost of a reduced transmission. In the $m/\mu$-selector a better resolution can be achieved by adding a static defocusing field to the two configurations of the electric field that we use here. This can be achieved by using different high voltages for the two electric field configurations or, more easily, by changing the duty cycle, i.e., by applying the presently used field configurations for different time-intervals. For many biomolecules, e.g., amino acids and peptides, the various conformers have large and widely different dipole moments. For phenylalanine, for instance, at least six conformers have been observed and their dipole moments are calculated to range from 1 D to 5.5 D. Therefore, the selection of its conformers would be feasible even at the present resolution.

In summary, we have selected and separated the cis and trans conformers of 3-aminophenol using switched electric fields in a quadrupole $m/\mu$-selector. The conformers are separated based on their distinct frequency dependent transmission characteristics. The dynamic focusing works best for the states with the lowest effective dipole moments, which are the lowest rotational states. Here, molecular packets with an excess of cis-3-aminophenol in low rotational states are created at an ac frequency of 2.95 kHz. Such conformer-selected molecular packets offer interesting perspectives for a variety of experiments. Since the selected states are the most polar ones, these samples are particularly useful for experiments in which aligned or oriented molecules are desired.

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[32] Alternatively, electric Stern-Gerlach-type beam deflection experiments could, in principle, provide partial spatial separation of conformers. However, it would only separate the most polar conformer from the others and would not provide an active confinement of the selected conformer. To the best of our knowledge, this application of beam-deflection has not been demonstrated yet.