Stability of high-mass molecular libraries: the role of the oligoporphyrin core

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Molecular beam techniques are a key to many experiments in physical chemistry and quantum optics. In particular, advanced matter-wave experiments with high-mass molecules profit from the availability of slow, neutral and mass-selected molecular beams that are sufficiently stable to remain intact during laser heating and photoionization mass spectrometry. We present experiments on the photostability with molecular libraries of tailored oligoporphyrins with masses up to 25 000 Da. We compare two fluoroalkylsulfanyl-functionalized libraries based on two different molecular cores that offer the same number of anchor points for functionalization but differ in their geometry and electronic properties. A pentaporphyrin core stabilizes a library of chemically well-defined molecules with more than 1600 atoms. They can be neutrally desorbed with velocities as low as 20 m/s and efficiently analyzed in photoionization mass spectrometry.

Keywords: macromolecular beams; laser desorption; photoionization; TOF-MS; molecular libraries

Introduction

In recent years, advanced chemical synthesis and mass spectrometry have become enabling tools for matter-wave experiments with complex molecules, and quantum interferometry was able to provide new insights into molecular properties, structure or dynamics. The partnership between these disciplines has recently led to the development of new matter-wave interferometers. In particular, a Kapitza–Dirac–Talbot–Lau interferometer was successfully used for demonstrating the quantum wave nature of entire macromolecules with masses in excess of \( m = 10 000 \) Da. Complementary to that, a matter-wave interferometer with three pulsed laser light gratings (OTIMA) was established to visualize quantum delocalization even of clusters of molecules, for de Broglie wavelengths as small as \( \lambda_{dB} = 200 \) fm. Both devices were built to explore the limits of quantum physics in the regime of high masses and to enable new measurements on neutral nanoparticles. Here, we focus on new experiments with fluoroalkylsulfanyl-functionalized oligoporphyrins, two molecular libraries based on two cores of different aromaticity. We study their structural stability under laser desorption and photoionization up to masses around 25 000 Da.

Several methods have been developed to launch massive and fragile molecules: electrospray ionization (ESI) and matrix-assisted laser desorption ionization (MALDI) are soft-volatization techniques for particles up to the MDa mass range. However, ESI produces mainly highly charged ions and is initially accompanied by a dense carrier gas. In contrast to that, MALDI is typically operated under high vacuum, and the molecules are released carrying none or only a few charges. MALDI often uses an organic matrix, such as dihydroxybenzoic acid, to enhance absorption of the desorbing UV laser and to assist in charging the analytes. Moreover, without the acid matrix environment, proton exchange in the sample is reduced. We exploit this to generate neutral molecules. The analyte itself can act as its own matrix if it is sufficiently absorptive and photostable. The resulting beam velocities can be smaller by an order of magnitude than in common MALDI studies. The desorption of neutral molecules can be monitored by post-ionization mass spectrometry. In earlier studies, photoionization of organic molecules was typically limited to particles below 2000 Da. Large clusters of guanine, tryptophan and gramicidin could still be observed using single-photon ionization with masses up to 7000 Da. Molecules can also be tailored to facilitate volatilization and photoionization. The covalent tagging of biomolecules with ionizable chromophores has already been implemented before, and also the functionalization of a chromophore monomer with long fluoroalkylsulfanyl chains was successful. Here, we extend this concept by exploring different molecular oligomers in the attempt to enhance the molecular stability at even higher mass.

Functionalization of two oligoporphyrins

Applications in quantum interferometry will profit from a molecular library, i.e. a molecular ensemble with well-defined mass separation. At a fixed molecular beam velocity, mass variation...
implies a controlled tuning of de Broglie wavelengths ($\lambda_{DB} = h/mv$).

We compare two molecular libraries\cite{27} as shown in Fig. 1: a tetrahe- 
dral arrangement of four zinc-coordinated porphyrins arranged around one tetraphenylmethane core and a planar pentaporphyrin. The pentaporphyrin is expected to reduce its ionization energy, and the steric configuration is supposed to facilitate the escape of ejected electrons upon ionization. Both oligomers are functional-
ized by different numbers of fluoroalkylsulfanyl chains to boost the molecular mass while keeping the electronic polarizability low. The ionization energy of each porphyrin is smaller than 7 eV.\cite{28} The central oligomer therefore increases both the number of functional anchor points and the photoionization cross section for vacuum ultraviolet (VUV) light.\cite{29}

**Experimental design**

In order to assess the VUV photoionization properties of library 1 and 2, we devised an experiment as shown in Fig. 2. We dissolve 2.5 mg of library 1 in 1.5 ml of pure diethyl ether or 7.5 mg of library 2 in 1.5 ml pure tert-butyl methyl ether. Droplets of this solution without any matrix are deposited on a clean stainless steel plate and dried in air. The N$_2$ laser beam ($E = 90 \mu J$, $\tau = 3$ ns) is focused onto the sample plate under an angle of 30°. During desorption, the sample plate is laterally shifted to ensure that each laser shot hits a fresh sample spot.

The neutral molecular beam is ionized by the VUV light of an F$_2$ excimer laser beam ($\lambda = 157.6$ nm, $\tau = 8$ ns, $E < 1.4$ mJ), focussed into a rectangular shape of $8 \pm 1$ mm$^2$. The ions are detected by a linear time-of-flight mass spectrometer (Kaesdorf, $m/\Delta m \approx 100$) in Wiley McLaren\cite{30} configuration.

**Mass spectra**

Figure 3 shows the post-ionization mass spectra of the two libraries. Each scan was averaged over 30 laser shots. The desorption energy

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**Figure 1.** Molecular library 1 (a) consists of four porphyrin systems with a central metal atom (Zn). They are linked via a tetraphenylmethane unit, which breaks the conjugation between the porphyrins. The most abundant molecule has $n = 17$ chains and a mass of 11 676 Da. Library 2 (b) is based on a metal-free porphyrin pentamer that is fully conjugated in the core. The most abundant molecule has $n = 40$ chains and a mass of 22 152 Da.

**Figure 2.** Experimental scheme for short-pulse laser desorption and post-ionization. The focused 337-nm beam of an N$_2$ laser (waist $w_x = 1.5$ mm, $w_y = 0.8$ mm) releases a molecular plume that expands into the ion extraction region. A 157-nm laser beam (39 mm further above) intersects with the molecules and ionizes them. The cations are detected using time-of-flight mass spectrometry (TOF-MS).
was identical for both spectra. The spectrum of molecular library 1 consists of eight well-resolved peaks (Fig. 3(a)), each representing a different number \( n \) of fluoroalkylsulfanyl chains, where \( n \) varies by 3–4 around the most probable value \( n = 17 \). A molecule with 21 chains corresponds to a mass of 13 492 Da. The mass difference between neighbouring peaks is 450 ± 10 Da, resulting from the nominal mass of a single chain (479 Da) minus the fluorine atom that is substituted in order to attach the chain. The characteristic abundance of all peaks in this mass spectrum agrees well with MALDI spectra.

The post-ionization mass spectrum also shows a distribution of low masses, the most prominent one denoted by \( C_1 \) to \( C_{10} \). The mass difference between neighbouring peaks of 457 ± 18 Da corresponds to the molecular weight of a single chain. This signal is attributed to the clustering of fluoroalkylsulfanyl chains in the expanding plume. The \( C_1 \) chain is abundant during synthesis and was not eliminated.

The result of a similar experiment with library 2 is shown in Fig. 3(b). While low-mass fragments also exist here, we have only zoomed into the high-mass spectrum to better resolve its structure. The number of chains varies from 34 to 45, with small peaks up to around \( n = 50 \). Individual peaks can still be clearly resolved up to 25 000 Da. The characteristic distribution after laser desorption/photoionization is similar to MALDI measurements immediately after synthesis.

Slow neutral beams of high-mass molecules

Slow molecules are essential for many experiments, such as beam deflection\([31,32]\) or advanced matter-wave studies. Because state-of-the-art interferometers are capable of operating with de Broglie wavelengths down to \( \lambda_{\text{dB}} = 200 \text{ fm} \), we are aiming for neutral particle beams with a momentum of less than 25 000 Da × 40 m/s.

Here, we measure the molecular velocity distribution by varying the delay time between the desorbing and the ionizing laser pulse. Figure 4 shows data for the molecular libraries 1 and 2. For a velocity analysis, we select three of the most prominent mass peaks of Fig. 3. Our data show a bimodal velocity distribution: the first component, best fitted by a Maxwell–Boltzmann distribution (dashed lines), is attributed to ions that are created in the source. The slow component (solid lines) represents neutral molecules that can only be detected if the VUV laser is on.

For library 1, the neutral part of the distribution is best fitted with a Gaussian distribution centred at around \( v = 90 \text{ m/s} \). There is still a substantial signal at \( v = 40 \text{ m/s} \) (Fig. 4(a)). Library elements of higher mass arrive with slightly lower speed. The slow fraction of library 2 (Fig. 4(b)) is centred at around \( v = 45 \text{ m/s} \) extending to velocities around 20 m/s for the most abundant mass peak. The functionalized pentaporphyrins are thus considerably more massive and substantially slower than similar macromolecular beams in earlier experiments.\([33,34]\)
Thermal robustness and photostability

The mass spectra and velocity curves of Figs 3 and 4 prove already the feasibility of neutral laser desorption and VUV photoionization of even the most massive elements of both libraries. Here, we are additionally interested in the stability of these compounds under thermal load and VUV exposure.

Even though the measured velocities are low on an absolute scale in molecular beam physics, the most probable kinetic energy of the 22,000 Da particle at 45 m/s corresponds to 0.24 eV. If the molecular beam were thermal, the corresponding temperature of about 2700 K would certainly disintegrate all particles. However, Fig. 4 indicates already that the desorption is not purely thermal.

In Fig. 5, we search for indications of thermal fragmentation between desorption and detection, by analyzing the evolution of the molecular mass ratios in library 1 and library 2 as a function of the post-desorption time. All mass peaks are normalized to the signal of the most abundant mass (n = 17 in library 1, n = 40 in library 2).

Figure 5. Evolution of the mass peaks in library 1 (a) and library 2 (b) as a function of the waiting time between laser desorption and post-ionization (all parameters as in Fig. 2): n = 16, n = 36 (black squares), n = 19, n = 38 (red circles), n = 20, n = 42 (blue triangles), n = 21 and n = 44 (pink inverted triangles). All mass peaks are normalized to the signal of the most abundant mass (n = 17 in library 1, n = 40 in library 2).

Figure 6 shows the signals of selected library members as a function of the ionizing laser energy. In both cases, the ion yield increases with growing pulse energy up to a saturation point. This corroborates the design assumption that the VUV ionization cross section of these molecules can be made sufficiently large by starting from an oligoporphyrin. The signal decrease with higher energy (library 1) is attributed to photoinduced fragmentation. Interestingly, this is not observed for library 2, because it has a larger number of fluoroalkylsulfanyl chains and more relaxation channels. Thus, it is expected that the excess energy is dissipated to more constituents.

Figure 6. Photoionization signal versus laser energy of library 1 (a) and library 2 (b). The distinct symbols describe different numbers of fluoroalkylsulfanyl chains attached to the porphyrin core. The lines are drawn to guide the eye. The signals were taken for a delay of 400 μs (library 1) and 800 μs (library 2) between desorption and ionization laser pulse.

Conclusions

We have studied the volatilization and post-ionization properties of two new molecular libraries with different core conjugation but similar chains. Both can be efficiently launched at low velocities and photoionized at a laser wavelength of 157 nm even at masses up to 25,000 Da. For the most abundant library element, we still find a substantial signal of neutral particles with velocities v < 45 m/s. This matches the initial de Broglie wave requirement λ_{dB} ≥ 200 fm. Fluoroalkylsulfanyl-functionalized porphyrins therefore fulfill important expectations for future matter-wave studies. The synthesis of such compounds is demanding, but the concept of fluoroalkylsulfanyl-functionalized particles has certainly not yet reached its limits. It will be intriguing to explore ways to launch and detect neutral mass-specified particles even more complex than those presented here.
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