EXPERIMENTAL STUDIES OF REACTIONS OF NEUTRAL AND IONTZED PAHs

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Polycyclic Aromatic Hvdrocarbon (PAH) molecules have been postulated as the carriers of the unidentified infrared features [1] as well as of the visible absorption bands [2] that are observed in interstellar and circumstellar environments. A more recent study led to the suggestion that small PAH's are mostly responsible for the infra-red emission [3].

Very few experimental data are presently available on the reactions of neutral or ionized PAH's [4]. We performed measurements of dissociative recombination, electron attachment and ion-molecule reaction rates with naphthalene, anthracene and phenanthrene using the FALP-MS (Flowing Afterglow Langmuir Probe - Mass Spectrometer) apparatus. One of the major difficulties of the experiment is to inject the PAH as a gas into the low, and to control its flow rate precisely.

It appeared that anthracene on one hand and naphthalene and phenanthrene on the other have completely different behaviours concerning electron attachment: naphthalene and phenanthrene do not attach as anthracene does. A value of $9 \ 10^{-10} \text{ cm}^3 \text{s}^{-1}$ has been measured for anthracene, indicating that an activation barrier may exist.

When attachment is important, it becomes difficult to measure dissociative recombination rates because two different sinks for the electron compete. Anyway an order of magnitude for the dissociative recombination of anthracene can be evaluated at 2 10^{-6} cm³s⁻¹. For naphthalene and phenanthrene, the rates are 3 10^{-7} cm³s⁻¹ and 7 10^{-7} cm³s⁻¹ respectively. The reactions of He⁺ and C⁺ with PAH's are fast and close to the Langevin rate. Fragmentation of the PAH can be observed in both cases, but is more severe with He⁺ than with C⁺, and aggregates can form from the ionized products.

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