POLYCYCLIC AROMATIC HYDROCARBONS IN DENSE CLOUD CHEMISTRY

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ABSTRACT

Virtually all detailed gas-phase models of the chemistry of dense interstellar clouds exclude polycyclic aromatic hydrocarbons (PAHs). This omission is unfortunate because from the few studies that have been done on the subject, it is known that the inclusion of PAHs can affect the gas-phase chemistry strongly. We have added PAHs to our network to determine the role they play in the chemistry of cold dense cores. Initially, only the chemistry of neutral and negatively charged PAH species was considered, since it was assumed that positively charged PAHs are of little importance. Subsequently, this assumption was checked and confirmed. In the models presented here, we include radiative attachment to form PAH\(^{+}\), mutual neutralization between PAH anions and small positively charged ions, and photodetachment. We also test the sensitivity of our results to changes in the size and abundance of the PAHs. Our results confirm that the inclusion of PAHs changes many of the calculated abundances of smaller species considerably. In TMC-1, the general agreement with observations is significantly improved, unlike in L134N. This may indicate a difference in PAH properties between the two regions. With the inclusion of PAHs in dense cloud chemistry, high-metal elemental abundances give a satisfactory agreement with observations. As a result, we do not need to decrease the observed elemental abundances of all metals, and we do not need to vary the elemental C/O ratio in order to produce large abundance differences of carbon species in TMC-1 (CP).

Subject headings: astrochemistry — ISM: abundances — ISM: individual (L134N, TMC-1 (CP)) — ISM: molecules — molecular processes

1. INTRODUCTION

Polycyclic aromatic hydrocarbons (PAHs) are hydrocarbon compounds with multiple six-membered aromatic rings. Such species are thought to be a common component of the interstellar medium in the Milky Way and in external galaxies (Draine 2003). PAHs are typically observed in infrared emission bands following excitation by either ultraviolet or visible radiation in assorted regions of the interstellar medium exposed to radiation (Léger & D'Hendecourt 1985; Léger et al. 1989; Allamandola et al. 1985; Salama et al. 1999). Despite all attempts, no specific types of PAHs responsible for these features have been identified (Mulas et al. 2006), and the form and intensity of the spectra vary strongly from one region to another (Draine & Li 2007), probably reflecting a variety of sizes and abundances. The origin of PAHs also remains an unanswered question (Williams 2003). Most believe that PAHs are formed in the atmospheres of carbon stars (Allain et al. 1997; Cau 2002) or in supernovae (Tappe et al. 2006).

Various observations show that the abundance of small free-flying PAHs has a tendency to decrease in dense clouds where UV photons do not penetrate (Boulanger et al. 1990; Bernard et al. 1993; Rapacioli et al. 2005). Rapacioli et al. (2005, 2006) proposed that PAHs form larger PAHs or PAH clusters in such dense regions. As a consequence, free-flying PAHs observed at the border of illuminated clouds may be the result of photodissociation of these larger species. Although they determined a minimal size of 400 carbons for these species using mid-infrared ISO C AM maps of photodissociation regions (PDRs), this number is very uncertain. If free-flying PAHs are not present in the dense interstellar medium, an efficient process is needed to aggregate them. Aggregation is poorly understood, and the time in which this process occurs is undetermined, as is the fraction of small PAHs remaining in dense clouds unexposed to intense radiation. Currently, constraints on the PAH size and abundance as a function of the physical conditions in the dense interstellar medium remain weak. In this article, we focus not on the size of PAHs, but on the chemical impact of their presence in dense cold interstellar cores. Nevertheless, the sensitivity of our results to size will be examined. We follow past authors by lumping all PAHs into three groups of species: neutral, positively charged, and negatively charged, designated PAH, PAH\(^{+}\), and PAH\(^{-}\), respectively. Of these three groupings, PAH\(^{+}\) species were found to be unimportant and were excluded from the detailed calculations.

The chemical effect of PAHs at steady state has been studied previously. In the diffuse interstellar medium, the inclusion of PAHs, among other effects, enhances the neutralization of atomic ions such as carbon, silicon, and sulfur, thus reducing their abundances (Lepp et al. 1988; Bakes & Tielens 1998). In their study of dense cloud chemistry, Lepp & Dalgarno (1988) concluded that for PAHs consisting of molecules with 30–50 carbon atoms, with fractional abundances greater than 10\(^{-3}\), PAH\(^{-}\) is the main source of negative charge, which considerably changes the abundances of some neutral species, enhancing, for example, the abundances of carbon-bearing molecules. Finally, Flower & Pineau des Forêts (2003) studied the importance of PAHs for C-shock models. In this article, we report a time-dependent study of the role of PAHs in the gas-phase chemistry of cold dense clouds and compare our results with observations. One focus of our study will be how the presence of PAHs affects the choice of elemental abundances.

The elemental abundances typically used for chemical models of dense clouds are referred to by the appellation “low metal”. Originally defined by Graedel et al. (1982), these abundances are based on the Copernicus satellite observations of the ζ Ophiuchi.
diffuse region (Morton 1974). However, in order to better reproduce the observation of molecules in dense clouds, the ζ Oph abundances of all the elements but He, C, N, and O are decreased by 2 orders of magnitude. This assumption implies that a strong depletion of the heavier elements occurs between the diffuse and dense cloud phases (Graedel et al. 1982). Although the details of this depletion are not understood, the depleted abundances are necessary to reproduce the abundances of molecular ions and, to a lesser extent, carbon-bearing species. Since the presence of PAHs in dense clouds should modify the ionization fraction, it may help to solve the problem of the choice of elemental abundances.

This article is organized as follows. In § 2, we describe the parameters of the PAHs used for the analysis and the chemical processes involving them. In § 3, other details of the chemical model are given. The influence of PAHs on the abundances of smaller species is reported in the next section, while the sensitivity of the results to the PAH parameters is discussed in § 5. In § 6, we compare observations in molecular clouds with grids of models in which we vary several parameters, including PAH size and abundance. The paper ends with our conclusions.

2. PAH PARAMETERS AND CHEMISTRY

Draine & Li (2007) reproduced the extinction curves of various diffuse regions of the Milky Way from Spitzer observations and constrained the size distribution of carbonaceous grains including PAHs down to 3 Å in size (see Fig. 6 of Draine & Li 2007). Using the model $j_M = 1$ of Draine & Li (2007), which has the largest abundance of small PAHs, we integrated over a radius range of $3.8 - 6.0 \, \text{Å}$. The integration yields a mean radius of 4 Å ($N_C = 30$) and an integrated fractional abundance with respect to $n_1$ of $3.07 \times 10^{-7}$. Multiplying by 30 leads to a fractional abundance of C atoms in PAHs of $9.1 \times 10^{-6}$, or roughly $3\%$ of the elemental carbon abundance.

Because PAHs may not be free in dense clouds, we decided to also consider PAH clusters of radius 10 and 100 Å. With the formula of Draine & Li (2007), these limits correspond to PAHs with 468 and $4.68 \times 10^{10}$ carbon atoms, leading to fractional abundances of $2 \times 10^{-8}$ and $2 \times 10^{-11}$, respectively, if one assumes (1) the same atomic carbon abundance in PAHs as in the smaller species, and (2) that weakly bound clusters can also be approximated by the same relationship between number of carbon atoms and size.

Bakes & Tielens (1998) and Dartois & D’Hendecourt (1997) found that the abundance of PAH decreases strongly in clouds with an $A_V$ larger than 2, and Lepp & Dalgarno (1988) found that the abundance of PAH is smaller than the abundance of the other forms of PAHs by about 1 order of magnitude in their calculations. For this reason, in our study of dense cloud chemistry, we did not originally include cations, but only neutral and negatively charged PAHs. Prodded by the referee, however, we looked more carefully at the role of PAH+, but concluded again that the exclusion is justified. The reconsideration will be discussed in the next section. We start the chemistry with neutral PAHs only, based on the results of Bakes & Tielens (1998) and Dartois & D’Hendecourt (1997). Neutral PAHs become negatively charged very rapidly, and our results after a very small time interval do not depend on whether the chemistry commences with PAH or PAH-.

The main problem of including PAHs in chemical models is that few experimental measurements of reaction rates have been performed on these molecules, and these few tend to involve rather small cationic PAHs only. For example, Snow et al. (1998) studied the rates of the reactions between a variety of positively charged PAHs with 6, 10, and 16 carbon atoms and the neutrals H, O, N, and H₂, while Betts et al. (2006) studied reactions of the coronene cation (C₂₄H₁₂) with the same reactants. Although association reactions appear to be dominant, most of the experiments were done at sufficiently high densities such that the mechanism, three-body association, is not what occurs in the low-density interstellar medium. Moreover, cationic PAHs are not important in our study.

The processes in our model include electron attachment, recombination of negative PAH ions with smaller positive ions, and photodetachment of negative PAH ions by UV radiation. We consider these in turn.

Negative PAH ions are formed by radiative electron attachment:

$$\text{PAH} + e^- \rightarrow \text{PAH}^- + \text{hv}. \quad (1)$$

The rate coefficient for this reaction is thought to depend on the size of the PAH. Omont (1986) and Allamandola et al. (1989) proposed two different relations: $k_e = 10^{-7}(N_C)^{1/3} \text{cm}^3 \text{s}^{-1}$ and $k_e = 1.2 \times 10^{-7}(N_C)^{1/2} \text{cm}^3 \text{s}^{-1}$. These two formulae lead to differences of about a factor of 2 for PAHs with $N_C < 100$. We arbitrarily decided to follow Le Page et al. (2001) and use the first formula. The rate coefficient is $1.3 \times 10^{-6} \text{cm}^3 \text{s}^{-1}$ for the attachment of electrons to neutral PAHs with 30 carbon atoms. The results depend on this value only if $k_e$ is decreased by a factor of 100. In other words, an increase of this rate coefficient does not change the results. Calculations by E. Herbst (2008, unpublished data) based on the statistical approach of Terzieva & Herbst (2000) indicate that with typical electron affinities in the range of 0.5–1.0 eV, the large rate coefficients deduced by previous authors are valid for neutral PAHs with 30 carbon atoms, but not for those significantly smaller, in agreement with the experimental results of Moustefaoui et al. (1998).

Negatively charged PAHs can recombine with positive ions, designated X⁺. These recombination reactions differ from dissociative recombination of positive ions with electrons in two important ways (Bates & Herbst 1988). First, they are less exothermic because of the electron affinity of the PAH. Secondly, some of the reaction exothermicity can be deposited in internal energy of the PAH, so that less is available to dissociate the positive ion upon neutralization. We assume that two resulting channels are possible for these recombinations, simple charge exchange:

$$\text{PAH}^- + X^+ \rightarrow \text{PAH} + X, \quad (2)$$

and, if the positive ion contains hydrogen:

$$\text{PAH}^- + X^+ \rightarrow \text{PAH} + X' + \text{H}, \quad (3)$$

where $X'$ designates a neutral with 1 H atom fewer than the ion. The terms “neutralization” and “dissociative neutralization” are used for these two processes to distinguish them from recombination with electrons. We assume equal branching fractions for the two sets of products, if they exist. If the neutralization leads to a neutral species that does not exist or is not included in the model, only the second channel is considered. The products of the dissociation are very uncertain, and laboratory experiments are required.
The rate coefficients for these reactions have been computed by taking into account the Coulomb attraction between the charged species via the expression (Spitzer 1941; Weisheit & Upham 1978)

$$k = \frac{\pi a^2 \langle v_0 \rangle}{\mu} \left(1 + \frac{e^2}{\alpha_{kB} T} \right),$$

(4)

where $e$ is the electronic charge, $\alpha_{kB}$ is the Boltzmann constant, $a$ is the radius of the PAH (or PAH cluster), and $\langle v_0 \rangle$ is the mean relative velocity between PAH$^-$ and ions, which is given by the expression

$$\langle v_0 \rangle = \sqrt{\frac{8k_B T}{\pi \mu}},$$

(5)

where $T$ is the temperature and $\mu$ the reduced mass, which is mainly determined by the mass of $X^+$. In expression (4), we have assumed that the radius of the PAH is the critical radius at which reaction can occur. The radius at which the second term equals the first term on the right-hand side is approximately $1 \mu m$ at $10 K$. This radius is much larger than the PAH radii considered here, so that $e^2/\alpha_{kB} T \gg 1$. The rate coefficient at $10 K$ is approximately $2 \times 10^{-7} \text{cm}^3 \text{s}^{-1}$ for an ion of $30 \text{amu}$ and a PAH of $4 \text{Å}$.

Relation (4) is based on the fact that, due to the delocalization of the negative charge on the PAH, the trajectory of the positive ion as it approaches the PAH will always be affected by this charge. Taking into account the complex shape and large number of possible PAH clusters in dense clouds, several negative charges may stick, and the attraction between such a system and a positive ion may depend on the localization of the electrons in the cluster. As a consequence, expression (4) may be more likely to be correct for small PAHs of $4 \text{Å}$ radius than for larger ones. For this reason, we consider the alternative possibility that rate coefficients for the reactions between PAH$^-$ and $X^+$ are given by the simple ballistic relation

$$k = \frac{\pi a^2 \langle v_0 \rangle}{\mu}.$$  

(6)

For small PAHs, the rate coefficient varies as $a$, whereas for large PAHs, if the process is ballistic, it varies as $a^2$. For a $100 \text{Å}$ PAH and an ion of $30 \text{amu}$, the rate coefficient at $10 K$ is approximately $2 \times 10^{-6} \text{cm}^3 \text{s}^{-1}$, or $0.1$ of the value for the smaller PAHs.

In addition to these reactions, PAH$^-$ is neutralized by photodetachment of electrons due to ultraviolet, visible, and even near-infrared photons:

$$\text{PAH}^- + h\nu \rightarrow \text{PAH} + e^-.$$  

(7)

Following Lepp et al. (1988), we assume that immediately above the threshold, which is equal in size to the electron affinity of the neutral, the cross section rises from $0$ to a flat value of $10^{-16} \text{cm}^2$ (see also Malloci et al. 2007). For a standard interstellar radiation field, this leads to a conservative estimate for the rate coefficient in the absence of shielding of approximately $10^{-7} \text{s}^{-1}$, neglecting differences in electron affinities among PAHs. A crude estimate of the dependence on $A_F$ is given by the simple exponential $\exp(-A_F)$. Even with an $A_F$ of $10$, the rate is not negligible in dense clouds. Indeed, under most conditions, photodetachment is more rapid than neutralization, but it must also be remembered that the electrons produced in photodetachment then attach to a neutral PAH. We do not consider photodetachment by cosmic-ray-induced photons. If we assume a rate coefficient of $(2 \times 10^4) \zeta$ for this process (Flower & Pineau des Forêts 2003), the photodetachment by direct UV photons is then more efficient by $1$ order of magnitude, even at an $A_F$ of $10$.

In our original calculations, we did not include PAH$^+$. In dense clouds, positively charged PAHs can be produced by secondary UV photons induced by cosmic rays and by charge transfer with small positive ions $X^+$. The latter process can neutralize the $X^+$ abundance and so affect the small-molecule chemistry; PAH$^+$ can then be destroyed by recombination with free electrons. Following the suggestion of the referee, we tested the importance of positively charged PAH species using rate coefficients from Flower & Pineau des Forêts (2003) for PAH$^+$ production by photoionization and for recombination with electrons. For the charge-transfer reactions, we used rate coefficients computed with the Langevin theory and the polarizability from Omont (1986). As an example, the Coulomb-based rate coefficient for the recombination reaction between PAH$^-$ and $X^+$ is $2.0 \times 10^{-7} \text{cm}^3 \text{s}^{-1}$ for PAHs of $4 \text{Å}$ and an ion of $30 \text{amu}$, whereas the rate coefficient for the charge-transfer reaction between PAH and $X^+$ for the same ion is $3.2 \times 10^{-9} \text{cm}^3 \text{s}^{-1}$. In the various models that we ran, we found that the abundance of PAH$^+$ is at least $2$ orders of magnitude smaller than the PAH and PAH$^-$ abundances. Moreover, its inclusion does not affect any of the results discussed below. This conclusion holds even when the recombination rate coefficient is the ballistic one.

3. MODEL PARAMETERS

We utilize a pseudotime-dependent gas-phase model, which computes the chemical evolution at a fixed temperature and density. A version of this model (NAHOOON) is now available online. The chemical database is a recent version of the OSU network, osu_01_2007, which can also be found online. The reactions involving PAHs will also be available at the same address.

We use three different sets of elemental abundances (EA1, EA2, and EA3), which are listed in Table 1 with respect to the total nuclear hydrogen abundance, along with the initial choice of

| Species | EA1 | EA2 | EA3 | Reference for EA2 |
|---------|-----|-----|-----|--------------------|
| He...... | $1.40(-1)$ | $9.00(-2)$ | $9.00(-2)$ | see text |
| N........ | $2.14(-5)$ | $7.60(-5)$ | $7.60(-5)$ | Cardelli et al. (1993) |
| O........ | $1.76(-4)$ | $2.56(-4)$ | $2.56(-4)$ | Meyer et al. (1998) |
| C+....... | $3.70(-5)$ | $1.20(-4)$ | $1.20(-4)$ | Cardelli et al. (1993) |
| S+....... | $8.00(-8)$ | $1.50(-5)$ | $1.50(-5)$ | Federman et al. (1993) |
| Si+...... | $8.00(-9)$ | $1.70(-6)$ | $1.70(-6)$ | Cardelli et al. (1994) |
| Fe+...... | $3.00(-9)$ | $2.00(-7)$ | $2.00(-7)$ | Savage et al. (1992) |
| Na+...... | $2.00(-9)$ | $2.00(-7)$ | $2.00(-7)$ | Savage & Sembach (1996) |
| Mg+...... | $7.00(-9)$ | $2.40(-6)$ | $2.40(-6)$ | Savage et al. (1992) |
| Cl+...... | $1.00(-9)$ | $1.80(-7)$ | $1.80(-7)$ | Savage & Sembach (1996) |
| P+....... | $2.00(-10)$ | $1.17(-7)$ | $1.17(-7)$ | Savage & Sembach (1996) |
| F+....... | $6.68(-9)$ | $1.8(-8)$ | $6.68(-9)$ | Federman et al. (2005) |

Note.—Here “af(b)” means $a \times 10^{-b}$.

1. Low-metal abundances from Graedel et al. (1982) and Morton (1974).
2. We considered an additional depletion compared to the observed value to take into account the oxygen and carbon depleted on grain mantles (see text).
3. The sulfur abundance found by the authors is larger than what is usually assumed as the cosmic abundance. We use the cosmic abundance of $1.5 \times 10^{-5}$.
4. Depleted value from Neufeld et al. (2005).
ionization. EA1 represents the low-metal abundances discussed earlier. EA2 is based on the high-metal abundances observed in the ζ Oph diffuse cloud, but is modified based on recent observations. The abundance of helium has not been observed in cold cores, and we use a value of 0.09 for EA2 with respect to hydrogen, based on observations in the Orion Nebula (Baldwin et al. 1991; Osterbrock et al. 1992). To account for the oxygen and carbon that are depleted on grain mantles in the form of H2O and CO, we removed 2.4% of C and 13% of O (Shalabiea & Greenberg 1995). It is possible that the metals are more depleted in dense clouds than in diffuse regions. Thus, for EA3 we assumed that the silicon and magnesium are totally depleted on grains, and we assumed an abundance of 1.5 × 10⁻⁶ for iron, as suggested by Flower & Pineau des Forêts (2003). The complete depletion of Si is one explanation for the nondetection of SiO in dense clouds (see Ziurys et al. 1989b). We also considered an additional depletion of a factor of 10 for chlorine, as suggested by the observation of HCl in cold material around OMC-1 (Blake et al. 1985). Note that in all the elemental abundances that we considered, the elemental C/O ratio is 0.4.

The species are assumed to be initially in atomic form except for hydrogen, which is assumed to be entirely in its molecular form. We consider typical physical conditions for dense clouds: a temperature of 10 K, an H₂ density of 10⁴ cm⁻³, a cosmic-ray ionization rate ζ of 1.3 × 10⁻¹⁷ s⁻¹, and a visual extinction Aᵥ of 10.

4. GENERAL EFFECTS OF PAHS

In this section, we present results for the three sets of elemental abundances (EA1, EA2, and EA3), both with and without PAHs. In the models with PAHs, referred to as EA/PAH, we assume their fractional abundance to be 3.07 × 10⁻⁷, our standard value for small PAHs. Variations in this value will be considered in §§ 5 and 6.

4.1. Ionization Fraction

Figure 1 shows the ionization fraction (sum of the negatively or positively charged species) as a function of time for the three sets of elemental abundances, each with and without PAHs. The initial ionization is typical of a diffuse cloud, since we start with ionized carbon in all models. With low-metal elemental abundances (EA1) and no PAHs, the ionization fraction stays high for only 10⁴ yr and then decreases sharply, reaching a steady state value of 3 × 10⁻⁸ by 10⁵ yr. With both EA2 and EA3, and no PAHs, the ionization fraction decreases slowly, remaining larger than 10⁻⁶ for about 10⁴ yr and eventually reaching a steady state value of a few × 10⁻⁷. When PAHs are added, the ionization fraction does not depend significantly on the elemental abundances. The main effect is to sharply decrease the earlier ionization fraction of the gas; within the first 300 yr, the ionization fraction goes from ~10⁻⁴ to ~5 × 10⁻⁶, its steady state value, which is only slightly larger than that achieved with the EA1 model. Thus, the inclusion of PAHs reduces the ionization fraction for EA2 and EA3 to a value close to that achieved with low-metal abundances and no PAHs. The steady state ionization fraction for model EA1/PAH is larger by a factor of 2 than that of model EA1 without PAHs. The ionization fraction of model EA2/PAH at steady state can be compared with the very similar value of 3.8 × 10⁻⁸ reported by Lepp & Dalgarno (1988) for a ratio of total PAH to hydrogen of 10⁻⁷ and high-metal abundances.

Figure 2 depicts the fractional abundances of neutral and negatively charged PAHs, as well as the electron abundance as functions of time for model EA2/PAH. The other PAH models are not shown because these abundances do not depend significantly on the chosen elemental abundances. By 300 yr, the fractional abundances of electrons and PAH⁻ drop to 10⁻⁹ and ~7 × 10⁻⁸, respectively, so that PAH⁻ becomes the dominant negatively charged species. The conversion of neutral to negatively charged PAHs occurs via electron attachment, which is very efficient at early times because of the high electron abundance. However, once formed, PAH⁻ is efficiently destroyed by reactions with...
atomic ions, which are also very abundant at the beginning of the chemical evolution. Note that the recombination of atomic ions is much more efficient with PAHs than with electrons, since the latter must occur radiatively. For instance, the neutralization of C$^+$ by PAH$^-$ (with $a = 4$ Å) is 4 orders of magnitude larger than by electrons at 10 K. Because of this efficient neutralization, the ionization fraction of the gas decreases much more quickly when PAHs are present (see Fig. 1). We also find that the abundance of neutral PAHs exceeds that of the anionic PAHs by a factor of 5–10 after 300 yr, which is also in agreement with the steady state results of Lepp & Dalgarno (1988).

4.2. Results for Selected Species

In the absence of PAHs, the higher metal elemental abundances EA2 and EA3 lead to a very different chemistry from EA1, and tend to worsen the agreement between the modeling results and the observations for selected classes of molecules (Graedel et al. 1982). With models EA2/PAH and EA3/PAH, however, the chemistry is much closer to that computed with the low-metal elemental abundances in the absence of PAHs. The major change in the chemistry is caused by the high abundance of PAH$^-$, which neutralizes atomic positive ions efficiently and, upon reaction with positive molecular ions, tends to neutralize them without significant dissociation, unlike dissociative recombination reactions with electrons. In addition, the decrease of the ionization fraction at initial stages of the chemistry increases the abundance of ionic species such as H$_3^+$ and H$_2$O$^+$ because they are depleted more slowly. At later times, some of the C-, S-, and N-bearing species are enhanced in the models EA2/PAH and EA3/PAH because of (1) larger elemental abundances and (2) a smaller electron abundance. We now present the results for the most important species and for selected molecules that show significant differences among models.

4.2.1. C, C$^+$, and CO

In Figure 3, we show our results for C and C$^+$ as functions of time for the three elemental abundances with and without PAHs. As expected, the fractional abundance of ionized carbon is decreased most rapidly when the PAHs are included. The agreement at steady state for the model EA2/PAH is in reasonable agreement with the result of Lepp & Dalgarno (1988). A consequence of the C$^+$ neutralization by PAH$^-$ is that atomic carbon becomes the main reservoir of carbon earlier than with the models without PAHs. At the age of dense clouds, between 10$^5$ and 10$^6$ yr, atomic carbon is more abundant in EA2 and EA3 than in EA1 because of a higher elemental abundance of carbon. Our steady state fractional abundance of neutral carbon ($\sim 3 \times 10^{-10}$) in model EA2/PAH is 3 orders of magnitude below that of the analogous model in Lepp & Dalgarno (1988). After early time, CO becomes the main reservoir of carbon in all models; the CO abundance (not shown in the figure) is insensitive to the model used.

4.2.2. H$_3^+$, HCO$^+$, H$_2$O$^+$, OH, H$_2$O, and O$_2$

The rapid decrease of the electron abundance in the models with PAHs leads to strong increases in the abundances of molecular ions such as H$_3^+$, H$_2$O$^+$, and HCO$^+$, as well as neutral species formed from them at the initial stage of the chemistry, as can be seen in Figure 4. At longer times, the ionic abundances show much less time dependence, with two general but not universal patterns: (1) they are more abundant in the models with PAHs than in the analogous models without them; and (2) they are most abundant in models with EA1 abundances and least abundant in models with EA2 abundances, both with and without PAHs. General reasons for the trends include the higher electron abundances in models EA2 and EA3, the slower depletion rates by negatively charged species (mainly PAH$^-$) in the PAH-containing models, and differences in abundances of important neutral species involved in depletion processes among the PAH-containing models.

The ion H$_3^+$ is a partial exception to the trends in that its abundance is not enhanced by PAHs at later times, mainly because it is destroyed principally by heavy neutral species such as CO and H$_2$O rather than by charged species. The abundance in EA2 and EA3 is smaller than in EA1 because of dissociative recombination by more abundant electrons, whereas it is smaller in EA2/PAH and EA3/PAH than in EA1/PAH because of reaction with H$_2$O, which is more abundant in the high-metal models. The enhancement in the ion HCO$^+$ for models with PAHs in the initial stages of the chemistry is even more dramatic than that of H$_3^+$ because of its formation from abundant carbon atoms by reactions that lead first to CH$_3^+$. At later times, HCO$^+$ is formed by the reaction between H$_3^+$ and CO in all models. Without PAHs, HCO$^+$ is destroyed by electrons, whereas it is destroyed by H$_2$O in models with PAHs.

The abundance of H$_2$O$^+$ is also dramatically enhanced in the initial stages of the chemistry in the presence of PAHs. Without PAHs, H$_2$O$^+$ is destroyed by electrons, whereas it is destroyed by PAH$^-$ in EA1/PAH and EA3/PAH. In EA2/PAH, due to a large abundance of elemental silicon, H$_2$O$^+$ reacts with gas-phase...
SiO, which is as abundant as $1.5 \times 10^{-6}$ at $10^6$ yr. Such a large abundance of SiO is in disagreement with observations in dense clouds (Ziurys et al. 1989b).

One consequence of the large H$_3$O$^+$ abundance at initial stages of the chemistry is that the abundances of OH, H$_2$O, and O$_2$, which stem from the recombination and neutralization of H$_3$O$^+$, are also enhanced in PAH-containing models. The situation is somewhat more complex at later times, with only water showing a much higher abundance in PAH-containing models. The predicted enhancement of H$_2$O in PAH-containing models is a problem, because the calculated abundances far exceed the observational upper limits of $\approx 3 \times 10^{-7}$ in L134N and $7 \times 10^{-8}$ in TMC-1 (Snell et al. 2000). The result for molecular oxygen is, on the other hand, not affected by PAHs at times after $10^5$ yr. Our theoretical O$_2$ abundance is about $10^{-9}$ at $10^5$ yr using EA2/PAH or EA3/PAH, which compares favorably with the observational upper limits of $1.7 \times 10^{-7}$ in L134N and $7.7 \times 10^{-8}$ in TMC-1 (Pagani et al. 2003).

4.2.3. Assorted Hydrocarbons and Oxygen-containing Organic Molecules

The computed abundances of the hydrocarbons c-C$_3$H$_2$ (cyclopropenylidene), C$_3$H$_4$ (methyl acetylene), C$_6$H$_2$ (triacetylene), and CH$_3$C$_4$H (methyl diacetylene) are shown in Figure 5 as functions of time. Without PAHs, the abundances peak at so-called early times ($10^5$ yr) for low-metal abundances (EA1), a well-known effect. For EA2 and EA3, on the other hand, the early-time peaks are weak to nonexistent and the peak abundances very low, an effect caused by the high electron abundances, which disrupt the growth of molecular ions. When PAHs are added, the abundances of the four hydrocarbons depicted are enhanced for all three models. Although the enhancement is significantly greater for EA2 and EA3, the peak EA1 values are still the largest. The enhancements are caused mainly by the lesser destructive effect of PAH compared with electrons when it reacts with intermediate ions in the chain of synthetic reactions, although in individual cases secondary effects can be isolated. The major effect, which comprises both lower rates and larger product neutral channels, is stronger than the enhancement in destruction rates of hydrocarbons due to often larger abundances of molecular ions. That differences remain among the PAH models despite the fact that PAH has the same abundance shows, once again, that other processes are also important. The computed abundances of hydrocarbons at steady state can be very different from the results of Lepp & Dalgarno (1988). For example, the C$_3$H$_2$ abundance is $\approx 10^{-15}$ in model EA2/PAH, whereas it is $10^{-9}$ in Lepp & Dalgarno (1988). This difference may be due to discrepancies in the network for carbon chemistry.
As can be seen in Figure 6, the abundances of the oxygen-containing organic neutral molecules HCOOH, CH2CO, CH3OH, and CH3CHO in models containing PAHs are enhanced at almost all times by at least 1 order of magnitude over the abundances in models without PAHs. Once again, the reason is generally the same as for the hydrocarbons: neutralization reactions between precursor ions with PAH− are not as destructive as dissociative recombination reactions with electrons. Even if a species, such as ketene, is formed by a neutral-neutral reaction, its precursor is likely formed at least partially through ion-molecule channels. As an example, consider the case of methanol. Methanol (CH3OH) is produced through the ion CH3OH+2, which itself is formed via the radiative association between CH+3 and water. In models without PAHs, the protonated ion can recombine dissociatively.

Fig. 5.—c-C3H2 (cyclopropenylidene), C6H6 (methyl acetylene), C6H4 (triacetylene), and CH3C4H (methyl diacetylene) abundances as a function of time for EA1 (solid lines), EA2 (dashed lines), and EA3 (dash-dotted lines) without PAHs (gray lines) and with PAHs (black lines). For the last three molecules, the abundance produced by model EA2 and EA3 without PAHs is smaller than 10−15.

Fig. 6.—Abundances of formic acid (HCOOH), ketene (CH2CO), methanol (CH3OH), and acetaldehyde (CH3CHO) plotted as functions of time for the three elemental abundances (solid lines: EA1; dashed lines: EA2; dash-dotted lines: EA3) without PAHs (gray lines) and with PAHs (black lines). The acetaldehyde abundance in model EA2 and EA3 is smaller than 10−15.
with electrons to form CH$_3$OH + H, although this channel is now known to be a minor one (Geppert et al. 2006). With PAH$^-$, on the other hand, we assume that this channel is the only one, since the neutral form CH$_3$OH$_2$ does not exist, and we only allow one hydrogen to be ejected from the species (see §2.1). Another reason for the enhanced abundance of methanol lies in the enhanced abundance of water. We note that with PAHs, the peak-predicted fractional abundance of methanol is still 1–2 orders of magnitude below the typical value in cold dense clouds of 10$^{-9}$ (Garrod et al. 2007).

Unlike the situation with the hydrocarbons, the abundances in models EA2/PAH and EA3/PAH are not significantly less, and sometimes even greater, than those in model EA1/PAH. The small differences suggest that PAH$^-$ plays an even more dominant role than in the case of the hydrocarbons. In the non-PAH models, although the EA1 abundances tend to be greater than those in EA2 and EA3, the effect is smaller than for hydrocarbons, except for acetaldehyde.

### 4.2.4. Cyanopolyynes

Cyanopolyynes are very abundant toward the famous cyanopolyyne peak (CP) of the cold core TMC1, yet their abundances are underproduced by gas-phase models unless a high-C/O elemental abundance ratio is assumed (Terzieva & Herbst 1998). As shown in Figure 7, the PAHs have a strong enhancing effect on the formation of these molecules, especially on the more complex cyanopolyynes. Once again, there is little difference among the results for the three PAH models. In general, the dominant formation mechanism for each cyanopolyyne more complex than HC$_5$N involves the next smaller member of the series. At 10$^5$ yr, the HC$_5$N, HC$_7$N, and HC$_9$N abundances are enhanced by more than 3 orders of magnitude over their values without PAHs. The peak fractional abundances are indeed larger than the values observed in TMC1 (CP) of 2 × 10$^{-9}$, 5 × 10$^{-10}$, and 2.5 × 10$^{-10}$ for HC$_5$N, HC$_7$N, and HC$_9$N, respectively (see Markwick et al. 2000 and references therein). Similar enhancements are seen for other nitrogen-containing molecules such as CH$_2$CHCN, CH$_3$CN, CH$_3$C$_3$N, and C$_3$H$_2$N$^+$.

### 4.2.5. Sulfur-bearing Species

The elemental abundance of sulfur is smaller by more than 2 orders of magnitude in EA1 than the two other sets of elemental abundances. As a consequence, most of the sulfur-bearing species have larger abundances with high-metal abundances at all times, with and without PAHs. In Figure 8, we follow the temporal evolution of four species: OCS, H$_2$S, CS, and C$_3$S. Three of these molecules tend to have enhanced peak abundances in the presence of PAHs, although the situation is more complex toward steady state. The exceptional cases are CS for all elemental abundances and H$_2$S for the low-metal case. The CS molecule is produced by the dissociative recombination of HCS$^+$ and HC$_2$H$^+$ by electrons or dissociative neutralization of HCS$^+$ by PAH$^-$. The larger ion HC$_2$H$^+$ does not react with PAH$^-$ to produce CS with our assumptions. As a consequence, the abundance of CS is smaller in the models including the PAHs. The late-time situation regarding H$_2$S is more complex. The primary formation mechanism for H$_2$S at most times is through the protonated precursor H$_3$S$^+$, which is formed by a chain of reactions starting with S + H$_3$ and proceeds through HS$^+$ and H$_2$S$^+$. Without PAHs, H$_2$S is also produced by the charge exchange between H$_2$S$^+$ and S. The ion H$_3$S$^+$ is more abundant in the low-metal case without PAHs, because it is produced by the reaction S$^+$ + H$_2$CO → H$_2$S$^+$ + CO, where S$^+$ is more abundant than the neutral form in the absence of PAHs.

### 5. Sensitivity to Several PAH Parameters

There are some observational and theoretical indications that PAHs in dense clouds may be aggregated into clusters with radii up to 100 Å. We have performed additional calculations using the three elemental abundances and PAH clusters of radii 10 and 100 Å with the reduced abundances discussed in §2. In our discussion of the results, we use the term PAH regardless of the size.
Fig. 8.—Abundances of OCS, H$_2$S, CS, and C$_3$S plotted as a function of time for the three elemental abundances (solid lines: EA1; dashed lines: EA2; dash-dotted lines: EA3) without PAHs (gray lines) and with PAHs (black lines).

Fig. 9.—Ionization fraction as a function of time for small and large PAHs using the three elemental abundances. For the two larger PAHs, we use expressions for the rate coefficients of PAH-"X" neutralization reactions that include (left) and exclude (right) charge attraction.
For the larger PAHs (10 and 100 Å), we consider two expressions for the rate coefficients of the neutralization reactions between PAH and positive ions: one involving charge attraction, as expressed in equation (4), and the other a simple ballistic one, as expressed in equation (6), for which the ballistic rate is considerably lower. For radii of 10 and 100 Å and a temperature of 10 K, the ballistic rate is, respectively, about 1000 times and 10 times smaller than the rate involving charge attraction. Another process with a rate coefficient that depends on the PAH size is the radiative attachment of electrons to neutral PAHs, which increases with increasing radius. In Figure 9, we show the ionization fraction as a function of time with each set of elemental abundances and PAHs of 4, 10, and 100 Å. As can be seen, the ionization fraction depends highly on the size of the PAH, and on whether or not charge attraction is considered. We do not show the dependence on the total abundance of C atoms incorporated into PAHs, but a larger abundance implies a more rapid decrease of the ionization fraction to a smaller value at late times.

In the left panel of Figure 9, neutralization proceeds with charge attraction for all three sizes of PAHs. For the two smaller sizes of PAHs, the main negatively charged species in the model remains PAH⁻. For the PAHs of 100 Å radius, the total abundance of PAHs is 2 \times 10^{-11}, much smaller than the abundance of electrons. The evolution of the ionization fraction is then closer to the case without PAHs, but the steady state ionization fraction can still be somewhat lower. For the case with model EA1 and the largest PAHs, the ionization fraction is the same with or without charge attraction and is not different from the model in which there are no PAHs at all. As a consequence, the 10 Å and (for EA2 and EA3) the 100 Å cases represent intermediate stages between the models with 4 Å PAHs and those without PAHs. The right-hand panel of Figure 9 shows the results when positive ion–PAH⁻ neutralization occurs ballistically for the larger PAHs. In this case, the 10 and 100 Å cases give results very similar to the model without PAHs, so that the effect of large PAHs at the chosen abundances is negligible in dense cloud chemistry. This effect occurs because the major depletion mechanism for PAH⁻ is photodetachment rather than recombination with positive ions, so that lowered neutralization rates have the sole effect of decoupling PAH⁻ from the ion-molecule chemistry.

The modification of the ionization fraction has consequences for the abundances of species. In general, PAHs of 100 Å (regardless of charge attraction) and of 10 Å without charge attraction lead to abundances similar to the models without PAHs. For most of the species, PAHs of 10 Å with charge attraction give intermediate abundances between the case of very small 4 Å PAHs and the case without PAHs. Many species such as C, C⁺, H⁺, HCO⁺, and OH show a difference smaller than a factor of 2 while using larger PAHs of 10 Å instead of smaller PAHs of 4 Å. Some species are largely underproduced using the elemental abundances EA2 and EA3 without PAHs. This is the case for C₂H₂, C₃H₄, C₅H₂, CH₃C₂H, CH₃CHO, and cyanopolyynes. In the
The presence of 10 Å PAHs, these species are almost as abundant as with smaller PAHs.

6. COMPARISON WITH OBSERVATIONS

The objective of this section is to study the impact of PAHs on the comparison with observations in the two cold cores L134N (north peak) and TMC-1 (cyanopolyyne peak). These two clouds are known to show differences in the abundances of C-rich molecules that are sometimes explained by differences in age or C/O ratio (Wakelam et al. 2006). Here we explore the possibility that they are also explained by the amount and size of PAHs present.

6.1. Method

The observed abundances in L134N (north peak) and TMC-1 (cyanopolyyne peak) are listed in Tables 3 and 4 of Garrod et al. (2007). In addition to these species, which comprise 42 molecules for L134N and 53 for TMC-1, we considered SiO, for which only upper limits of $3 \times 10^{-3}$ and $2 \times 10^{-4}$ for TMC1 have been determined (Ziurys et al. 1989a).

In this comparison, we utilize the three different sets of elemental abundances listed in Table 1. Of all these elements, sulfur is the least constrained by abundances of gas-phase molecules, such that the depleted fraction of sulfur is uncertain (Ruffle et al. 1999). PAH size and abundance are also poorly determined parameters. Instead of fixing these parameters to previously used values, we try to give some trends by comparing observations with grids of models in which these three parameters are varied in addition to time.

We computed 441 models by varying the abundance of elemental sulfur between $10^{-8}$ and $10^{-3}$, the abundance of PAHs between $10^{-10}$ and $10^{-7}$, and the size of PAHs between 4 and 400 Å. Note that the largest values of PAH sizes and abundances used here are not in agreement with the relation $N_C = 460(a/10 \text{ Å})^3$, which was used in the first part of the article. Considering how little information is available on the relation of size to number of carbons for PAHs, in particular for PAH clusters, we consider the PAH size and abundance as independent. The agreement between present models and observations quantified using statistical comparisons, such as minimization, is not good enough to give reasonable constraints on model parameters. For this reason, we did not use such methods, but instead for each model and each time we counted the number of species with fractional abundance $X$ reproduced by the model within 1 order of magnitude ($X_{\text{obs}}/10 < X_{\text{model}} < 10X_{\text{obs}}$), where $X_{\text{obs}}$ is the observed abundance and $X_{\text{model}}$ the computed abundance. For those species for which only upper limits to their abundances are known and for the case of atomic carbon, for which only a lower limit is known (Stark et al. 1996), we check that the predicted abundance is in agreement with the observed limit; when this is the case, we consider the species reproduced.

6.2. Results

Figures 10 and 11 show the maximum percentage of species reproduced in each cloud as a function of the parameters chosen. Column (a) shows the level of agreement as a function of PAH
size and abundance, with each point optimized for the best elemental abundance of sulfur and the best time. Column (b) shows the level of agreement as a function of the elemental abundance of sulfur and the time optimized at each point for the best PAH parameters. Finally, column (c) shows the agreement as a function of the elemental abundance of sulfur and the time for models without PAHs. These results are shown for each set of elemental abundances (1: EA1, 2: EA2, and 3: EA3). Best agreement, with or without PAHs, is most often achieved at so-called early times of $10^3 - 10^6$ yr. Moreover, high levels of agreement cannot be achieved with high abundances of PAH clusters, as shown in the upper-right corners in column (a) for both figures.

The maximum percentage of molecules reproduced depends on all the parameters, and several solutions exist. PAHs do not significantly improve the best agreement for L134N (already above 70% without PAHs), but they do expand the range of parameters (time and elemental abundance of sulfur) for which we have an agreement larger than 70%, especially if elemental abundances EA1 or EA2 are utilized. Note particularly the expanded range of agreement with the use of elemental abundances EA2 (the high-metal ones). The optimum PAH parameters corresponding to the yellow zones in column (b) do not depend on the optimum sulfur abundance. As an example, high agreement is obtained with $3 \times 10^{-10}$ abundance of 4 Å PAHs both for low $(3 \times 10^{-8})$ and high $(1 \times 10^{-6})$ elemental abundances of sulfur. The chemistry in TMC-1, on the other hand, is better reproduced if PAHs are present, as can be seen immediately by comparison of columns (b) and (c) in Figure 11. Without PAHs, agreement rarely extends to more than 60%, while with PAHs, agreement in excess of 70% exists for all three sets of elemental abundances. For elemental abundances EA1, the PAH parameters are relatively unconstrained, while for the high-metal abundances (EA2), high abundances of small PAHs are needed. The situation is less clear for EA3. For both clouds, we find isolated areas of agreement larger than 70% at very early times, <1000 yr, for large abundances of sulfur $(>8 \times 10^{-6})$ and big (>25 Å) PAHs $(>10^{-8})$ if EA2 and/or EA3 is utilized. Such agreement is of mathematical interest only.

7. CONCLUSION

Most chemical studies of the dense interstellar medium rely purely on gas-phase processes. A few contain, in addition, a varying amount of surface chemistry on interstellar grains, but the inclusion of PAHs, which occupy a middle ground in size between small gas-phase species and dust particles, is quite rare. Part of the reason for this neglect is that to include PAHs properly, one should view them as individual species, but the actual individual species that make up the PAH content of interstellar clouds are not well constrained from spectral observations, and even their chemistry has not been studied in great detail. Therefore, the inclusion of PAHs means that one must treat them as small grains and follow only their charge state. In this study, as in previous ones, we lump all PAHs into just three groups, depending on whether they are neutral, positively charged, or negatively charged. Positively charged PAHs were initially assumed to be unimportant in cold dense sources, an assumption that was confirmed upon the request of the referee. They were not included in the chemical network used for the results presented here. By including an assortment of chemical processes involving negatively charged and neutral PAHs into our standard gas-phase network of reactions, we are able to determine the effect of PAHs on the time-dependent gas-phase chemistry of cold and dense interstellar cloud cores. The gas-phase model employed is known as osu-01-2007 and can be downloaded online. To this model, we add electron attachment to neutral PAHs, neutralization between negatively charged PAHs and positively charged smaller ions, and photodetachment of negatively charged PAHs by external photons. Our standard model contains a fractional abundance of $3.07 \times 10^{-7}$ PAHs of size 4 Å, corresponding to 30 carbon atoms. In addition to this model, we vary the size and abundance of the PAHs to study the impact of these parameters on the chemistry. We also use three different sets of elemental abundances with different abundances of heavy atoms.

In general, the dominant effect in our calculations is that electrons are replaced by PAH$^-$ as the dominant negative charge carrier, an effect that weakens and eventually vanishes as the size of the PAHs and their fractional abundance decrease, probably due to clustering. When the abundance of PAH$^-$ exceeds that of electrons, the overall ionization fraction is reduced, because atomic positive ions are far more rapidly neutralized by reactions with PAH$^-$ than by radiative recombination with electrons. These changes lead to many secondary effects. Our calculations show that many of the molecular species have abundances that are strongly affected by the presence of PAHs. In particular, molecular ions tend to have increased abundances at most times, and large unsaturated neutrals such as the polyynes, as well as saturated species such as methanol, show very large increases in abundance. The sensitivity of our results to variations in the size of PAHs depends on assumptions made in the rate of neutralization of PAH$^-$ and positive ions. The differences among results obtained with different elemental abundances are much smaller in models containing PAHs than in models without them.

By comparing observations in the two dense cloud cores L134N and TMC-1 with grids of models, we showed only that the chemistry in such sources is not consistent with large amounts of large PAH clusters. We are not able to give more severe unambiguous constraints on the PAH parameters, since high levels of agreement can exist for large parameter variations. The situation is similar for the elemental abundance of sulfur, where values over several orders of magnitude are consistent with maximum agreement with observations. As regards time, both the standard early time and very early times can lead to optimum agreement, although the latter must be rejected on astronomical grounds. The situations for L134N and for TMC-1 show differences. For L134N, the best agreement is not improved when PAHs are included, although the range of optimum sulfur elemental abundances and times are extended, most significantly so for the high-metal abundances (EA2). For TMC-1, on the other hand, the best level of agreement is significantly better when PAHs are included, using any of the three elemental abundances. This difference may indicate a difference in PAH properties between the two regions. In particular, for EA2, optimum agreement in TMC-1 is obtained with smaller and more numerous PAHs, while for L134N, such parameters make matters worse. Finally, unlike the case of gas-phase models without PAHs, models that include PAHs are able to accommodate high-metal elemental abundances, similar to those observed in the diffuse medium, if the effects of PAHs are included and optimized.

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4 See http://www.physics.ohio-state.edu/~eric/research.html.
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