The molecular condensations ahead of Herbig-Haro objects

II. A theoretical investigation of the HH 2 condensation

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Abstract. Clumps of enhanced molecular emission are present close to a number of Herbig-Haro (HH) objects. These enhancements may be the consequence of an active photochemistry driven by the UV radiation originating from the shock front of the HH object. On the basis of this picture and as a follow up to a molecular line survey toward the quiescent molecular clump ahead of the HH object, HH 2 (Girart et al. 2002), we present a detailed time and depth dependent chemical model of the observed clump. Despite several difficulties in matching the observations, we constrain some of the physical and chemical parameters of the clump ahead of HH 2. In particular, we find that the clump is best described by more than one density component with a peak density of 3×10^5 cm⁻³ and a visual extinction of ≤ 3.5 mag; its lifetime can not be much higher than 100 years and the impinging radiation is enhanced with respect to the ambient one by probably no more than 3 orders of magnitude. Our models also indicate that carbon-bearing species should not completely hydrogenate as methane when freezing out on grains during the formation of the clump.

Key words. ISM: clouds - ISM: molecules - ISM: general

1. Introduction

Herbig-Haro (HH) objects are regions of hot plasma created by stellar jets impacting on interstellar gas. In the last decade, molecular condensations have been found ahead of HH objects. These clumps were detected in enhanced emission from HCO⁺ and NH₃ (e.g. Rudolph & Welch 1988; Torrelles et al. 1992; Girart et al. 1994). The detected clumps are cool, with temperatures around 10 K, and quiescent: they show no evidence of shocks and are therefore not connected dynamically with the stellar jet. The enhanced emission comes from localised regions in the vicinity of the HH objects, and has now been detected from a number of objects (Girart et al. 2002 (hereafter Paper I), and references therein). Torrelles et al. (1992) suggested that the chemical enhancements may be explained by UV radiation from the HH object; Girart et al. (1994) added that the mechanism induced by the UV radiation could be the removal of the ice mantles from grains in the clumps ahead of the jet which then would drive a photochemistry. In this picture, the jet and its HH object constitute a micro-probe of the physical and chemical nature of the molecular cloud. Theoretical studies (Taylor & Williams 1996; Viti & Williams 1999, hereafter VW99) supported this scenario and also showed that the chemistry of these clumps should be rich; the impact of a high radiation field on a dense gas in a clump should yield enhanced abundances not only of HCO⁺ and NH₃ but also of several other species such as CH₃OH, H₂S, H₂CO, SO and SO₂. Also, as the jet is moving at quite high speeds (100– 1000 km s⁻¹), the clumps ahead of these jets must be transient in that they will eventually be reached by the jet and may be destroyed within a timescale on the order of 1000 years or less (Raga & Williams 2000). As a consequence, considering their short lifetime, statistically, these clumps must be very common.

Understanding the nature of the condensations ahead of HH objects can therefore shed light on the nature of the interstellar molecular clouds. The condensations associated with HH objects appear to have the same physical characteristics as transient dense clumps believed to be the birthplace of low-mass stars. The physical and chemical nature of these clumps are usually explored via the detection of emission in several molecules that trace dense gas such as NH₃ and CS. However, low angular resolution contour maps of molecular emission of these two species are puzzling: the CS half-power contour is much more extended than that of ammonia, their peaks are displaced, and the widths of their lines are different (Pastor et al. 1991; Myers et al. 1991; Morata et al. 1997). The NH₃ contour is closely associated with embedded sources in the cloud.

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One interpretation of these differing morphologies is that the cloud is composed of transient clumps, unresolved in these observations, that develop CS as the chemistry evolves, but disperse before appreciable NH_3 is formed (Taylor et al. 1996). Occasionally, a more massive clump will evolve faster – both chemically and dynamically – and so would be responsible for the spatial association of NH_3 with star formation signatures. Observational support for this model is now provided by high resolution BIMA observations in lines of CS, HCO^+ , and N_2H^+ of a portion of the molecular cloud L673 (Morata et al. 2003). These observations show much structure on scales of 10 arcsec, different in the different species, suggesting that the structure is transient and has time-dependent chemistry.

We suggest that the clumps ahead of the HH objects may be of the same population as the clumps observed by Morata et al. (2003) but they happen to have a different chemistry because they are illuminated by – and are photoprocessed by – the HH radiation. Hence, observations of the condensations associated with HH objects give an independent method of determining physical parameters of dense clumps in interstellar molecular clouds.

Following theoretical studies which indicated that the impact of radiation from an HH object on a dense gas should lead to a rich chemistry (Taylor et al. 1996; VW99) a first detailed line survey of a condensation ahead of an HH object (HH 2: Herbig 1951; Haro 1952) was made (Paper I): the condensation ahead of HH 2 was first established by detections of enhanced HCO⁺ and NH₃ (Davis et al. 1990; Torrelles et al. 1992, 1994; Choi & Zhou 1997). In Paper I we presented CSO and BIMA detections of 14 species and we derived the chemical composition of the clump. We also briefly compared the observational results with the VW99 models and found a qualitative agreement. This comparison helped to constrain some of the parameters of this clump such as its maximum age (for which we found ~1000 yrs). However, the VW99 models were not specific to HH 2: the abundances at any one time depended on several initial parameters such as the strength of the radiation field, the fraction of gas frozen on to the grains as ices, the age and size of the clump, and the initial composition of the gas. As a consequence, we were not able to constrain important physical parameters such as the radiation field strength and whether the observed chemistry was a direct consequence of the vicinity of the HH 2 knot. Moreover, it is now established that the strongest HH 2 knot, which is also the highest excitation knot, has associated X-ray emission (Pravdo et al. 2001). This may affect the chemistry and needs to be taken into consideration. A more detailed model specific to HH 2 is therefore needed.

The aim of this work is to develop a detailed model of the chemistry in the clump associated with HH 2 and to use it to determine the chemical and physical parameters of the HH 2 clump. In Sect.2 we summarize the observational situation, and in Sect. 3 we describe the models used in the calculations. Section 4 gives results of calculations for many parameter variations; these results are discussed in Sect. 5, and our conclusions are listed.

Physical and chemical parameters of the clump ahead of HH 2 – the present status of observational studies

In Paper I we derived column densities from a 0.8–3 mm molecular line survey toward the quiescent molecular clump ahead of the bright HH 2 object. For simplicity, we list again here the column densities observed (Table 1). Note that from more recent observations with the IRAM 30 m telescope (in preparation) we derive a column density for HCO⁺ which is one order of magnitude less than the one derived in Paper I. This difference is probably due to the different beam sizes (CSO ~30"; IRAM ~21") and different pointing positions, in a region where density and chemistry both vary strongly with position. This point will be further discussed in a future paper. Here we only draw the attention on the possible range of values for the abundance of HCO⁺.

In particular, from the morphology of the emission we find three groups of species (Girart et al. 2003, Paper III, in prep.):

- SO, SO₂ and CH₃OH with compact emission just ahead of the HH object;
- HCO⁺, HCN and H₂CO with a strong compact clump just ahead of the HH object surrounded by more diffuse and extended emission north-east of the compact clump;
- CS and C¹⁸O with a peak intensity position clearly offset from that of the other molecules.

Chemically, this may mean that SO, SO₂ and CH₃OH are heavily affected by the HH 2 knot (by the UV flux or the X-ray emission or a combination of both) and they are probably very low in abundance in the interclump medium; HCO⁺, HCN and H₂CO are also affected by the HH 2 object and this effect shows up as a molecular intensity enhancement with respect to the interclump medium, where these species are however already present in detectable quantities. Finally CS and C¹⁸O seem to be largely unaffected by the vicinity of the HH 2 knot, or maybe even depleted (e.g. by photodissociation) just ahead of the HH object.

There is a large gradient of abundances for most of the species which indicate a complex structure. This is clear from Fig. 1 where the column densities deduced from BIMA observations of selected species as a function of distance to the HH 2 knot are shown. By taking into account the column densities listed in Table 1, and the gradient of values from Fig. 1, we now investigate whether we are able to reproduce both the observed column densities and the spatial distribution and chemical differentiation shown in the BIMA maps.

3. The calculations

3.1. The model

The model we adopt is an extension of the model employed in VW99. The model calculation is carried out in two phases; both are time- and depth-dependent. In Phase I, the clump is represented by a region that collapses gravitationally within the molecular cloud. The collapse is treated as a free-fall, as described by Rawlings et al. (1992). The effects of a retarded

 Table 1. Extract of Table 3 from Paper I: column densities of observed species.

Molecule	Column density	
СО	3.8(17)	
HCO ⁺	1.0(13)-1.0(14)*	
CH ₃ OH	6.4(13)	
H_2CO	5.7(13)	
SO	3.2(13)	
SO_2	1.7(13)	
HCN	4.1(12)	
CN	3.4(12)	
CS	2.8(12)	

* The low value is a lower limit derived from IRAM observations. See text.

collapse have already been discussed (VW99). During the collapse phase, gas–phase chemistry and freeze–out on to dust grains with subsequent processing are assumed to occur. The initial density of the clump is taken to be 1000 cm⁻³, and the final density is treated as a free parameter. In Phase II we simulate the presence of the HH object by subjecting the clump to the HH radiation field (which evaporates all the ices and promotes a photochemistry in what was formerly dark cloud material) and also to the X-ray flux. The chemical network is taken from the UMIST database (Millar et al. 1997) and is described in VW99. We refer the reader to VW99 for a detailed explanation of the model, to which the following changes have been made for the present study:

- i) In Phase II, we simulate crudely the presence of X-ray ionization by enhancing our standard cosmic ray ionization rate, $\zeta (1\zeta = 1.3 \times 10^{-17} \text{ s}^{-1})$, as in Nguyen et al. (2002).
- ii) In Phase II, the enhanced radiation field is now depthdependent ($\propto 1/r^2$, where *r* is the distance from the HH shock). The constant of proportionality is decided by imposing a value (variable) for the radiation field at the edge of the clump.

3.2. The model parameter grids

We have computed two separate grids of models (Grids A and B), differing in the assumption of the age of the clump at the end of Phase I. The results from these two grids will be discussed separately in Sects. 4.1 and 4.2. For Grid A Models, we assumed that once the final density was reached (between 3×10^5 yrs and 4.2×10^5 yrs), the clump survived for a substantial amount of time (~2 Myrs) before being affected by the HH object. This is different from our assumption in VW99, and from Grid B (see below), where we assumed that once the final density was reached the clump was *immediately* irradiated.

In order to determine the parameter space needed to be investigated, we recall the observational results from Papers I and III: the temperature of the clump is estimated to be ~13 K and the upper limit to the hydrogen molecule number density is $\sim 3 \times 10^5$ cm⁻³. The size of the clump is derived from the SO contour, giving a clump diameter of ~0.01 pc. This corresponds to a visual extinction of about 6 mag with respect



Fig. 1. Slices of the rotational temperature (top panel), column density (central panel) and relative abundance with respect to the CO (bottom panel) as a function of distance to HH 2. The left vertical dashed line shows the position of HH 2 knot L, while the right dashed line shows approximately the position where the CSO spectrum was taken (Paper I). The slice has a 0 position RA(J2000) = 05:36:26.55, Dec(J2000) = -06:47:06.0 and the position angle of the slice is PA = 169.

to the position of the HH object. However, this inferred extinction across the clump assumes that the SO contour corresponds to the true size of the clump and the gas around the clump is completely deficient in SO; and that the material between us and the clump along the line of sight is diffuse. All these assumptions are uncertain, and we therefore treat the clump as a one-dimensional multi-point slab extending up to 6.4 mag and examine the time-dependent chemistry throughout the clump. We consider lower and upper limits to the density of 3×10^4 and 3×10^5 cm⁻³, respectively.

We assume that, during Phase I, a fraction of the gas phase material is frozen on to the grains, and undergoes hydrogenation, where this is chemically possible. This fraction is arranged by adjusting the grain surface area per unit volume, and assumes a sticking probability of unity for all species. The fraction of material on grains is then dependent on the product of the sticking probability and the amount of cross section provided per unit volume by the adopted grain size distribution.



Fig. 2. Column density of HCO⁺ as a function of time at $A_v = 3.4$ mag for selected models from Grid A. The two horizontal lines encompass the area of possible observed values for HCO⁺.

This product was varied so that at the end of Phase I, we would have a low, medium or high abundance of ices (see Col. 3 of Table 2). Note that as the chemistry is time-dependent, different species form at different times. As a consequence the material frozen out on the grains at any time is *not* representative of the whole gas but of selected species. In this case, we have determined the fraction of material frozen out by monitoring mainly the amount of frozen CO, and H_2O at the edge and centre of the clump.

During Phase II, we investigated the chemical consequences of changes in the following parameters: firstly, the strength of the X-ray emission by enhancing the cosmic ray ionisation rate up to 1000 times its canonical value; and, secondly, the strength of the radiation field impinging on the edge of the clump (χ_{edge}) by varying it from 20 to 10⁴ times the mean interstellar radiation intensity, here called Habing units (Habing 1968). This latter parameter is of paramount importance as it is completely unconstrained by direct observations of the HH 2 object from which only the H₂ line flux can be derived. Table 2 gives the parameters defining most of the models computed.

4. Results

4.1. Long-lived clumps: Models A

In order to investigate whether it is feasible for these condensations to be long-lived before they are illuminated by the HH radiation we compute a grid of models (Grid A) where the chemistry is allowed to continue after the clump has reached the desirable final density. The column densities estimated from Grid A Models are shown in Table 3 and the HCO⁺ abundance as a function of time for selected Grid A models is shown in Fig. 2.

Briefly, we discuss the trends evident in the table and figure. Note that from the BIMA maps we believe that the geometry of the clump we observed is such that our line of sight is almost perpendicular to the direction of the radiation coming from the HH object (Paper III) – this means that our

Table 2. Model parameters. The notation $a(b)$ signifies $a \times 10^{b}$. The
model number is listed in Col. 1; Col. 2 shows the density of the gas;
Cols. 3, 4 and 5 show respectively the percentage of the gas depleted
onto grain at the end of Phase I, the enhancement factor of the radia-
tion field in units of Habing and enhancement factor of the ionization
rate, in units of ζ .

Model	$n_f ({\rm cm}^{-3})$	FR (%)	χ (Habing)	ζ (s ⁻¹)	
		Grid A			
A1	3(5)	15	20	1	
A2	3(5)	15	1(3)	1	
A3	3(5)	15	1(4)	1	
A4	3(5)	60	20	1	
A5	3(5)	60	1(3)	1	
A6	3(5)	60	1(4)	1	
A7	3(5)	80	20	1	
A8	3(5)	80	1(3)	1	
A9	3(5)	80	1(4)	1	
A10	3(5)	60	20	100	
A11	3(5)	60	20	1(3)	
A12	3(4)	20	20	1	
A13	3(4)	20	1(3)	1	
A14	3(4)	20	1(4)	1	
A15	3(4)	40	20	1	
A16	3(4)	40	1(3)	1	
A17	3(4)	40	1(4)	1	
A18	3(4)	40	20	100	
A19	3(4)	40	20	1(3)	
A20	3(5)	60	1(3)	100	
		Grid B			
B1	3(5)	7%	20	1	
B2	3(5)	20%	20	1	
B3	3(5)	55%	20	1	
B4	3(5)	0%	20	1	
B5	3(5)	0%	1	1	
B6	3(5)	7%	1(3)	1	
B7	3(5)	20%	1(3)	1	
B8	3(5)	55%	1(3)	1	
B9	3(5)	20%	1(4)	1	
B10	3(5)	20%	1	1	

column densities need to be estimated as we did in VW99 (i.e. $10^{15}(X_i/10^{-6})$ cm⁻² for $A_v \sim 1$) rather than calculated by integrating along the depth points (which represent a slab perpendicular to our line of sight, along the radiation field line). From Fig. 2, we note already that most Grid A models fail to reproduce the high abundance of HCO⁺ for long enough time, although models A2, A5, A8 and A20 do succeed in matching the HCO⁺ lower limit. From Table 3, we note that most of the species are affected by an enhanced radiation field. At early times HCO⁺ increases when we increase the impinging radiation field. However, we note that at late times (not shown), it is lower when the radiation field is higher. This effect is more pronounced at low-to-intermediate visual extinctions ($A_v \leq 4$ mag). CS, on the other hand, seems to decrease slightly with an increase in radiation when the freeze-out is low but increases when the latter is high. SO behaves in a similar way to CS but at low freeze out, when we increase the

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	A1	A2	A3	A4	A5	A6	A7	A8	A9	A20
СО	9.1(17)	9.2(17)	9.4(17)	6.0(17)	6.1(17)	6.4(17)	4.5(17)	4.5(17)	5.0(17)	6.1(17)
HCO^+	1.4(11)	5.0(12)	1.8(13)	6.7(10)	3.7(12)	3.6(13)	3.9(10)	2.5(12)	4.2(13)	5.1(12)
H_2CO	9.4(13)	1.1(15)	2.5(15)	1.2(14)	1.4(15)	4.8(15)	1.5(14)	1.3(15)	5.8(15)	1.4(15)
CS	3.5(13)	3.4(13)	2.3(13)	6.3(12)	7.7(12)	1.3(13)	1.5(12)	2.4(12)	1.1(13)	7.7(12)
SO	3.1(12)	2.9(13)	2.2(13)	5.8(12)	2.4(13)	3.7(13)	4.6(12)	1.8(13)	4.2(13)	2.4(13)
SO_2	3.7(12)	3.3(12)	1.4(12)	4.5(12)	4.2(12)	3.5(12)	3.4(12)	3.3(12)	4.4(12)	4.2(12)
CN	1.4(12)	4.7(13)	5.6(14)	1.0(12)	4.2(13)	1.2(15)	9.8(11)	4.0(13)	1.5(15)	4.1(13)
HCN	7.3(13)	9.5(13)	1.6(14)	6.1(13)	8.4(13)	2.4(14)	5.8(13)	8.0(13)	2.8(14)	8.2(13)
CH ₃ OH	3.3(15)	2.7(15)	5.6(14)	4.1(15)	3.4(15)	1.3(15)	3.6(15)	3.0(15)	1.6(15)	3.4(15)

Table 3. Column densities for selected species for Models A1–A9 and A20 at 3 years after the clump has been irradiated, at $A_V = 3.4$ mag.

impinging radiation from 20 to 1000 it actually slightly increases. SO₂ decreases in abundance with an increase in radiation at low and intermediate freeze out but increases when we increase χ from 20 to 1000 at high freeze out (but decreases again when χ is 10 000). At early times, CN always seems to increase slightly with radiation while CH₃OH decreases with increasing radiation, regardless of the visual extinction or age; HCN shows the same behaviour of early time HCO⁺, but at all ages. At first sight, it seems that the enhanced radiation field in the vicinity of a HH object cannot be directly responsible for a molecular enhancement which survives for a long enough time to be observed.

One of the most significant parameters affecting the results is the fraction of gas frozen-out and hydrogenated on the grains by the end of Phase I. We find that many of the species, including HCO⁺, always decrease in abundance with an increase of freeze out.

These two trends are apparently inconsistent with our previous models (VW99) and with the explanation that the radiation generated in the HH shock affects the clumps ahead of it by evaporating the icy mantles. However, a close examination of the HCO⁺ formation network shows that the main characteristic in this set of Models A is the lack of C⁺: although H₂O ice is abundant (due to the hydrogenation of oxygen bearing species when they freeze out), C^+ is very low. This is due to a much lower abundance of neutral carbon at the end of Phase I since this phase is of a long duration. When C freezes out, it hydrogenates into methane. This effect is also more pronounced for models where the density is high $(3 \times 10^5 \text{ cm}^{-3})$. For Grid A Models, once the collapse is stopped and the final density is reached the clump continues to chemically evolve for a long time at high density, and the freeze-out and hydrogenation efficiency adopted assures that most of the carbon is locked in frozen methane.

This effect is partly reduced by simulating the effect of X-rays. The primary effect of the increase in the ionisation rate is to raise the abundances of various ionised species (mostly single atoms) compared to their neutral abundances. This effect is most obvious at high A_v , where an enhanced ionisation rate is the only significant contributor to ionisation, since the radiation field is heavily suppressed. However, the increase in the ionisation rate also causes more dissociation of molecules. The increased dissociation of CO causes a significant increase in C at high A_v , where the abundance of C is lowest, and this in

turn allows higher values of C⁺ via ionization of C. The C⁺ may then readily react with H₂O, which at early times is abundant at all depths due to evaporation of icy mantles, enhancing HCO⁺. After 1000 years, ion-neutral reactions and photodissociation have greatly reduced the abundances of most molecules at lower A_v , for all values of the ionisation rate. In the case where the ionisation rate is 100 times the canonical value, at high A_V the increased ionization causes considerable losses in the most abundant molecules. CO, which is already very abundant, is the main beneficiary of the destruction of H₂O and CH₄. Increasing the ionisation rate to 1000 times the canonical value makes the increases in CO at large A_v much greater, and abundances of H₂O and CH₄ fall correspondingly.

In general, HCO^+ is seen to be enhanced with the simulated X-ray field at large A_v , while other species such as CH₃OH are not strongly affected, although at very high ionisation rates their abundances are reduced. Sulphur-bearing species are not affected at low A_V but they increase, as a consequence of the faster destruction of H₂S, at high A_v . In summary, simulating an X-ray field by enhancing the ionization rate has some effect on the chemistry of the clump. However, the consequences are not significant when compared to other effects.

From this first set of models we conclude that if the quiescent clump is indeed long-lived before the jet arrives, then the suggestion that these regions of enhanced emission arose because of the influence of HH radiation on the dense clump is not correct. A requirement for this theory to be correct is that we have a reasonable amount of free carbon in the gas phase when the jet gets close enough to irradiate the clump; we find that if the clump is long-lived, then either most of the carbon is locked in frozen CH4 or it is locked in CO (assuming a very low freeze out parameter). We therefore assume in what follows that - in general - high density clumps in interstellar clouds are transient. This is indeed consistent with the picture proposed by Taylor et al. (1996), with the observational evidence of Girart et al. (2002), and with the possible origin of such transient clumps in slow-mode MHD waves (Falle & Hartquist (2002) - see Sect. 1.

Although Models A have not provided a match to the observations of HH 2 they have been instructive. Their results have demonstrated that the chemistry is extremely sensitive to variations in the density, to the fraction of gas frozen out as ices, and to lifetime of the clumps. We decided therefore to perform a second (reduced) grid of models (grid B) where Phase I is ended and Phase II is started as soon as the collapse is finished.

4.2. Transient clumps: Models B

Details of the grid of Models B are listed in Table 2. We note that for Grid B we have also evaluated a model in which species do not freeze out on to dust grains (Model B4), a model like Model B4 but where the radiation field in Phase II is kept equal to the ambient one (Model B5), and a model where freeze-out occurs but in which there is no enhancement in the ambient radiation field (Model B10). Other models were also computed, and we shall discuss them later.

HCO⁺ is one of the molecules most enhanced in abundance in this clump; yet it is difficult to make in large amounts in the conditions found in clumps ahead of HH objects. Therefore, we first concentrate on this species. In Fig. 3 we show the column density of HCO⁺ as a function of time for $A_v \sim 3$ (left plot) and ~6 mag (right plot). From Table 1 and Fig. 1 we derive a column density for HCO⁺ between 3×10^{12} cm⁻² (indicated by the HCO⁺ 1–0 emission not corrected by optical depth from BIMA (see Fig. 1) and IRAM observations) and 10^{14} cm⁻² (obtained from the multi-transition analysis from BIMA and CSO observations) depending on where around the HH object we look. These values are shown in Fig. 3 (left and right panels) as the area encompassed between the two horizontal lines. From both plots we can see that it is quite difficult to reproduce a high abundance of HCO⁺. It certainly seems that, at high A_V , HCO⁺ is never abundant. We can also definitely exclude models where freeze-out is not included but there is enhanced radiation (B4), models in which freeze-out is not included and there is no enhancement of the radiation field (B5), and models in which freeze-out is included but there is no enhancement of the radiation field (B10). These conclusions are consistent with the assumption that the enhanced chemistry of the condensation ahead of the HH 2 object is a direct consequence of evaporation of ices together with a photochemistry caused by the enhanced radiation field due to the proximity of the HH object. These conclusions confirm that there seems to be no alternative way of producing a high abundance of HCO⁺.

Another result from Fig. 3 is that high abundances of HCO⁺ are not easily maintained beyond about one hundred years. This period is, in fact, comparable to the expected survival time of the clump ahead of HH 2 (see Sect. 1). From these plots, the best model candidates are Models B6, B7 and B8 (see Table 2), for $A_V \leq 3.5$ mag. They are favoured over Models A2, A5, and A8 in that they succeed in maintaining a higher HCO⁺ abundance for a longer period.

Having found the models that match reasonably well the HCO^+ observations, we now investigate whether other species are also reasonably accounted for by these same models. While most of the species are well represented by B7 and B8, we find that some of the species, in particular H₂CO, CN, and HCN are always *overabundant* (up to 2 orders of magnitude) in all our models. We now consider the chemistry of these species in an attempt to resolve these problems.

The chemistry of H_2CO , and to a lesser extent CN and HCN, is indirectly governed by the abundance of CH_4 . The main reaction forming H_2CO is:

$$CH_3 + O \rightarrow H_2CO + H$$
 (1)

where CH_3 comes from the photodissociation of CH_4 . We note that CH_4 is abundant in most of our models where freeze-out occurs – this is because we assume that every neutral and ionized carbon atom that hits the grains hydrogenates to methane which is then released to the gas phase in the second phase of our model. The efficiency of hydrogenation of carbon on grain surfaces is entirely unknown. In the models discussed so far, this efficiency has been taken to be 100%. In order to consider the sensitivity of the results to this assumption we investigate a series of models in which we vary the efficiency with which neutral and ionized carbon, on hitting the grains, hydrogenate to methane.

We also investigate several models where the initial relative elemental abundance of oxygen is halved (still maintaining the same C/O ratio). The justification for reducing the initial relative oxygen abundance comes from our rather high choice for the initial relative abundance of cosmic oxygen (4.6×10^{-4}). This value is close to the upper limit of the interstellar oxygen abundances derived from 14 lines of sight by Meyer et al. (1998, see Table 2).

The new models are B11 (as B2 but with only 10% of carbon freezing as methane), B12 (as B7 with only 10% of carbon freezing as methane and half of the original oxygen), B13 (as B9 with only 10% of carbon freezing as methane and half of the original oxygen), B14 (as B11 with half of the original oxygen), B15 (as B12 but with no carbon freezing as methane).

Figures 4 and 5 show the results from our models, including the new ones, for H_2CO and HCO^+ . Again, for all the figures, we also encompass the area of possible observational values for the column densities (lower and upper limit are derived as for HCO⁺). None of the models succeed in reproducing the low abundance of H₂CO for long enough: B13 lies within the observational band for ~ 15 yrs which may be too short to be significant; B15 on the other hand is still overabundant for the first 100 yrs but only by a factor that varies between ~ 1.5 and ~5. In the context of modelling interstellar chemistry, and the uncertainty in the observations, this is a reasonable agreement. Moreover, if the observed H₂CO transitions has an opacity of 2-3, then the fitting with B15 would be much better. We therefore favour Model B15. HCO+ is also well reproduced by Model B15 (Fig. 5) as are SO (Fig. 6) and methanol (Fig. 7). Note that the abundance of SO is slightly underabundant in Model B15. However, the initial relative sulphur abundance is very uncertain and we chose it to be a factor of hundred lower than the solar value - a conventional assumption for dark clouds (Oppenheimer & Dalgarno 1974; Ruffle et al. 1999). This value may well be in error by a factor of a few, so that we may regard Model B15 as well within the uncertainty for SO.

However, Model B15 does not reproduce the abundances of HCN and CN. We have examined the chemistry for these two species; we find that at the end of Phase I their abundances are consistent with those observed towards other dense cores (cf. van Dishoeck 1998), although already somewhat too high



Fig. 3. Column density of HCO⁺ as a function of time at $A_v = 3.4$ mag (left) and 6.2 mag (right) for selected models from Grid B. The two horizontal lines encompass the area of possible observed values for HCO⁺.

with respect to the observed values for HH 2. This suggests that the (quite conventional) chemical network that produces CN and HCN is probably correct. In Phase II, however, CN quickly increases due to the enhancement of the radiation field, via the reaction:

$$NO + C \rightarrow CN + O$$
 (2)

CN is then efficiently destroyed by the neutral reactions:

 $N + CN \rightarrow N_2 + C \tag{3}$

$$NH_3 + CN \rightarrow HCN + NH_2 \tag{4}$$

therefore enhancing HCN as well as CN. The first reaction is dominant at early times while the second reaction is dominant after the first 40 years or so. HCN is therefore a direct consequence of CN and its destruction (via photodissociation) leads back to CN. Photodissociation of CN does not appear to be very important. This may be due to the rate adopted for this reaction (taken from the UMIST database). We therefore investigate, arbitrarily, the possibility that a faster CN photodissociation rate would decrease both CN and HCN, and do this by artificially increasing the rate. We find no substantial difference for the first 100 years. Finally, a possible reason for the discrepancy of predicted abundances of CN and HCN versus observed could be that the observed column densities are underabundant because they are optically thick. This will be further investigated by future BIMA observations for HCN.

In conclusion, we can find no reason why the abundances of HCN and CN predicted by our models should be much larger than apparently observed. The values computed at the end of Phase I are reasonably consistent with observations of dark clouds, and the introduction of the HH radiation field will certainly initiate a chemistry that should *increase* CN and HCN production. We conclude that the models suggest the presence of large amounts of CN and HCN not currently observed, or



Fig. 4. Fractional abundance of H_2CO as a function of time for selected models from Grid B at $A_V \sim 3$ mag.

alternatively that the HCN and CN observed column densities are underestimated due to optical depth effects.

5. Discussion and conclusions

As a follow up to a molecular line survey toward the quiescent molecular clump ahead of the HH 2 object (Paper I), we have presented here a detailed time-dependent chemical model of the observed clump. There are still considerable uncertainties remaining in the observational results, but we have been able to use these data to restrict the possible chemical models and to identify physical parameters of the clump. Our general conclusions concerning the model are:

 our chemical analysis suggests that the clump ahead of HH 2 was young when it was first irradiated by the HH object. A long period (~2 My) at high density is definitely excluded by this study. If future observations of a large



Fig. 5. Fractional abundance of HCO⁺ as a function of time for selected models from Grid B at $A_V \sim 3$ mag.



Fig. 6. Fractional abundance of SO as a function of time for selected models from Grid B at $A_V \sim 3$ mag.

sample of clumps ahead of HH objects also indicate young material, this may mean that in general clumps in molecular clouds are transient, consistent with current interpretations of molecular clouds by both single dish and array telescopes (see Sect. 1);

- BIMA array observations show a much more complex structure than revealed by single dish CSO observations (see also Hester et al. 1998 where a much more complex structure is suggested). As a consequence, our theoretical clump, modelled on the basis of the SO contour from the CSO, is probably only representative of the small, high density gas just ahead of the HH object. We find that it is not possible for the clump ahead of HH 2 to have a uniform high density is $\sim 3 \times 10^5$ cm⁻³ closest to HH 2 and falling with distance from it, is probably more realistic;
- the characteristic chemical signature seen in HH 2 is of short duration, probably no more than a few hundred years, depending on the density in the clump and the penetration of HH radiation into it. Thus, secular changes may occur



Fig. 7. Fractional abundance of CH₃OH as a function of time for selected models from Grid B at $A_V \sim 3$ mag.

in clumps ahead of HH objects on astronomically short timescales, possibly short enough to be detectable;

- the peculiar chemistry found ahead of HH objects is achieved for very particular combinations of ice fraction on grains (at the end of Phase I) and enhanced radiation field caused by the proximity of the HH object;
- it is difficult to reconcile theoretical predictions with observations if a large quantity of gas phase CH₄ is present in Phase II (expected when depletion and hydrogenation occur). We infer that carbon bearing species may not completely hydrogenate as methane when freezing out on grains in Phase I. Note that if this is the case, then Grid A models may be more successful in reproducing the observations; therefore our result on the lifetime of the clump prior to the irradiation of the HH objects, would be less conclusive.

Despite the difficulties we have found in modelling the chemistry in the clump ahead of HH 2, we believe that we are able to constrain some of its physical and chemical parameters. In particular:

- the "clump" is probably better described by a two or more density components, with an average density of $1-5 \times 10^4$ cm⁻³ and a peak density of $\sim 3 \times 10^5$ cm⁻³;
- the visual extinction along the high density component cannot be higher than ~3.5 mag;
- the lifetime of the clump *after* it starts being irradiated cannot be much longer than ~100 years;
- the radiation field impinging at edge of the high density component of the clump is expected to be in the range of 20 and 1000 Habing, and certainly substantially less than 10⁴ Habing;
- the composition of the clump before it is irradiated by the HH object is probably such that not more than half of the molecular material is depleted on the grains;
- X-ray emission from the HH 2 object does not appear to affect substantially the chemistry of the clump.

Our aim in this work is to reproduce the abundances observed towards the clump ahead of HH 2. Other HH objects may provide a different irradiating field, in both intensity and spectrum, from that of HH 2, and some clumps may have lower/higher density and therefore probably a lower/higher degree of freeze-out than the clump we observe ahead of HH 2. Therefore, the chemistry of clumps ahead of other HH objects may be diverse.

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