Singly and doubly deuterated formaldehyde in massive star-forming regions

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ABSTRACT

Context. Deutered molecules are good tracers of the evolutionary stage of star-forming cores. During the star formation process, deuterated molecules are expected to be enhanced in cold, dense pre-stellar cores and to deplete after protostellar birth.

Aims. In this paper, we study the deuteration fraction of formaldehyde in high-mass star-forming cores at different evolutionary stages to investigate whether the deuteration fraction of formaldehyde can be used as an evolutionary tracer.

Methods. Using the APEX SEPIA Band 5 receiver, we extended our pilot study of the $J = 5\rightarrow 2$ rotational lines of HDCO and D$_2$CO to eleven high-mass star-forming regions that host objects at different evolutionary stages. High-resolution follow-up observations of eight objects in ALMA Band 6 were performed to reveal the size of the H$_2$CO emission and to give an estimate of the deuteration fractions HDCO/H$_2$CO and D$_2$CO/HDCO at scales of $\sim 6''$ (0.04–0.15 pc at the distance of our targets).

Results. Our observations show that singly and doubly deuterated H$_2$CO are detected towards high-mass protostellar objects (HMPOs) and ultracompact H ii regions (UC H ii regions), and the deuteration fraction of H$_2$CO is also found to decrease by an order of magnitude from the earlier HMPO phases to the latest evolutionary stage (UC H ii), from $\sim 0.13$ to $\sim 0.01$. We have not detected HDCO and D$_2$CO emission from the youngest sources (i.e. high-mass starless cores or HMSCs).

Conclusions. Our extended study supports the results of the previous pilot study: the deuteration fraction of formaldehyde decreases with the evolutionary stage, but higher sensitivity observations are needed to provide more stringent constraints on the D/H ratio during the HMSC phase. The calculated upper limits for the HMSC sources are high, so the trend between HMSC and HMPO phases cannot be constrained.

Key words. ISM: molecules – astrochemistry – stars: formation

1. Introduction

Deuterium fractionation, and how it changes as star-formation proceeds, is a powerful tool to establish the evolutionary stage and age of star-forming cores. The formation of deuterated molecules is favoured at low temperatures ($T \leq 20 \, \text{K}$) and at high densities ($n \geq 10^4 \, \text{cm}^{-3}$) in molecular cloud cores, the ideal conditions for CO freeze out (Caselli et al. 1999, 2002). However, deuterated molecules are also observed with relatively high abundances in warmer environments, for example, in circumstellar disks and in hot cores (e.g. Salinas et al. 2017; Sakai et al. 2018). Different chemical processes contribute to the deuteration enrichment at different temperatures. At low ($T \approx 10$–20 K) temperatures, the dominant gas-phase process is $\text{CH}_3^+ + \text{HD} = \text{CH}_3^+ \text{D} + \text{H}_2$. At high temperatures ($T \approx 30$–50 K), however, the $\text{CH}_3^+ + \text{H}_2 = \text{CH}_2^+ \text{D} + \text{H}_2$ and $\text{C}_2\text{H}_5^+ + \text{HD} = \text{C}_2\text{H}_5\text{D} + \text{H}_2$ reactions become important (see detailed gas-phase deuteration models by e.g. Roueff et al. 2007; Roueff & Lique 2013). Reactions on the grain surfaces can also influence the D/H ratio via low-temperature surface reactions during the pre-stellar and starless phase that enrich the D/H ratio of molecules in the solid phase. These molecules can then be desorbed into the gas phase during the warm protostellar phase (e.g. Brown & Millar 1989a,b; Tielens 1983).

We can characterise the deuteration fraction, $D_{\text{frac}}$, as the relative abundance between a species containing a deuterium atom as compared to the same species containing a hydrogen atom. It is thus expected that $D_{\text{frac}}$ is enhanced in cold and dense pre-stellar cores, and it should then decrease after protostellar birth, when the young stellar object heats up the central region of the core (Caselli 2002). Several studies (see e.g. Crapsi et al. 2005; Caselli et al. 2008; Emprechtinger et al. 2009) have confirmed this theoretical scenario for low-mass star-forming cores, with the observations of deuterated species produced by gas-phase reactions (e.g. H$_3^+$ and N$_2$H$^+$). Recent studies have shown that high $D_{\text{frac}}$ values are also typical for high-mass star-forming cores and that the $D_{\text{frac}}$ of some species could also be an evolutionary tracer in the intermediate- and high-mass regime (e.g.}
Busquet et al. 2010; Fontani et al. 2011, 2015; Sakai et al. 2012; Zahorecz et al. 2017).

Recent theoretical studies (e.g. Bovino et al. 2017; Körtgen et al. 2018; Hsu et al. 2021) have shown that the evolution of the deuterium enrichment depends on several factors, such as the initial physical conditions, grain size distribution, the depletion fraction, and the cosmic ray ionisation rate. Observational studies (e.g. Crapsi et al. 2005; Emprechtinger et al. 2009; Fontani et al. 2011, 2015; Treviño-Morales et al. 2014) have confirmed that the deuterium chemistry is a good chemical clock for both low-mass and high-mass star-forming cores. However, in order to have an accurate estimate of the evolutionary stage of the source, it is necessary to use as many different molecular tracers as possible since the derived $D_{\text{frac}}$ may differ due to variations in the chemical evolution of the different species (see e.g. Treviño-Morales et al. 2014).

We can distinguish the following evolutionary stages in high-mass star-forming regions: high-mass starless cores (HMSCs), high-mass protostellar objects (HMPOs), and ultracompact H II regions (UC H II regions) (e.g. Beuther et al. 2007; Tan et al. 2014). Gerner et al. (2014) derived the chemical age of $\sim 6 \times 10^4$, $\sim 4 \times 10^4$, and $\sim 10^5$ yr for the HMSC, HMPO, and UC H II stage, respectively. Their derived total timescale for the high-mass star-formation is consistent with theoretical estimates of $\sim 10^5$ yr. Based on the study of several deuterated species in 27 massive cores in different evolutionary stages from HMSCs to UC H II regions, Fontani et al. (2011, 2015) found that species formed exclusively in the gas ($N_2H^+$) that showed different evolutionary trends from those that formed partially ($NH_3$) or totally (CH$_3$OH) on grain mantles. The $D_{\text{frac}}$(N$_2H^+$) is higher in the HMSC phase and it drops by about an order of magnitude during the HMPO and UC H II stages, while deuterated methanol is detected only towards HMPOs and externally heated HMSCs. Therefore, high $D_{\text{frac}}$(N$_2H^+$) is a good indicator of the initial conditions in starless and pre-stellar cores, while high $D_{\text{frac}}$(CH$_3$OH) values are a good probe of the earliest protostellar phases (Fontani et al. 2015). Furthermore, $D_{\text{frac}}$(NH$_3$) does not show a significant difference across the different evolutionary stages, and thus, there are no dominant formation pathways (gas-phase versus grains) for NH$_3$ and its deuterated forms.

As NH$_3$, formaldehyde (H$_2$CO) can also be produced both in the gas phase and on grain surfaces. The two main pathways for the production of H$_2$CO involve CH$_3$OH in the gas phase (see Yamamoto 2017) and multiple hydrogenation of CO on the ices (see e.g. Roberts & Millar 2007). Its gas phase pathway is similar to the one of N$_2H^+$, but it can also occur at warmer temperatures (30–50 K, see Parise et al. 2009). The ice phase formation route of H$_2$CO is similar to that of CH$_3$OH and their deuterated forms. Laboratory work and observational studies of low-mass protostars suggest an important contribution from grain surface chemistry to the production of H$_2$CO (Watanabe 2005; Roberts & Millar 2007; Bergman et al. 2011). However the relative importance of the grain surface versus gas-phase formation routes remains unclear. Furthermore, only a handful D$_2$CO detections in high-mass star-forming regions are available (D$_2$CO has been firmly detected towards the Orion Compact Ridge, NGC7129-FIRS2, AFGL5142, IRAS05358+3543, and tentatively towards the MonR2 UC H II region; Fuente et al. 2005; Treviño-Morales et al. 2014; Turner 1990; Zahorecz et al. 2017).

All of these initial results suggest that H$_2$CO and its deuterated species form mostly on grain surfaces. However, more observations carried out with interferometers (i.e. at high-angular resolution) are needed to separate the HDCO and D$_2$CO emission originating from the small and dense cores and to get a more detailed view of the formaldehyde deuteration in high-mass star-forming regions.

In this paper, we report our findings of a search for HDCO and D$_2$CO emission towards an extended sample of high-mass star-forming regions at different evolutionary stages, using the Atacama Pathfinder EXperiment (APEX\textsuperscript{1}) and the Atacama Large Millimeter/submillimeter Array (ALMA) telescopes. The source sample and the observations are described in Sect. 2, while our results are presented in Sect. 3. In Sect. 4, we discuss our results and put them in context with respect to previous findings in low-mass and high-mass star-forming regions. In Sect. 5, we summarise our conclusions.

2. Observations and target selection

In this section, we describe our selected sources and the single-dish and interferometric observations we have carried out for our study.

2.1. Selected targets

The selected sources are extracted from the sample of Fontani et al. (2011), for which the deuteration fraction of other molecules (e.g. N$_2$H$^+$, CH$_3$OH, and NH$_3$) has been measured (Fontani et al. 2015). We selected the brightest sources from each evolutionary class, which are observable with the APEX telescope from the southern hemisphere. Table 1 reports the observed central coordinates, velocities, distances, kinetic temperatures, and source evolutionary classification for our targets. The sample is divided into three HMSCs, three HMPOs, two sources containing HMSC and HMPO, which would represent an intermediate stage between HMSC and HMPO, and three UC H II regions (see details below). The source coordinates have been centred at interferometric millimeter and centimeter continuum peaks or high-density gas tracer peaks (e.g. NH$_3$ with VLA) that are separated by at least $\sim 10''$ from other cores. The APEX beam is $\sim 34''$ at 190 GHz. Therefore, for AFGL5142 and IRAS05358+3543, multiple cores at different evolutionary stages are covered within the same pointing. We categorise these two sources as a transitional stage between HMSCs and HMPOs in our work.

2.2. IRAM-30m H$_2$CO observations

Previous observations of these objects were performed using the IRAM-30m telescope (details can be found in Fontani et al. 2011, 2015). The frequency setups included the $3_{0,1} - 2_{0,2}$, $3_{2,2} - 2_{2,1}$, and $3_{2,1} - 2_{2,0}$ transitions of H$_2$CO. The spectroscopic information of these transitions is shown in Table 2 (Müller et al. 2005). The spectra were obtained and calibrated in antenna temperature units, $T_A^*$, and then converted to the main beam brightness temperature, $T_{MB}$, with the following equation: $T_A^* = T_{MB}/\eta_{MB}$, where a main beam efficiency, $\eta_{MB}$, of 0.66 was used. The H$_2$CO line emission was detected in all sources and the measured intensities show a wide range between 0.24 K $< T_{MB} < 13$ K (see Fig. 1).

2.3. APEX SEPIA observations

The sources were observed with the APEX SEPIA receiver (Swedish-ESO PI receiver for APEX; Billade et al. 2012;
Table 1. Observed sources and their coordinates, LSR velocities, distances, kinetic temperatures, and \(^{12}\text{C}/^{13}\text{C}\) ratio at the corresponding distance.

| Name            | RA (hh:mm:ss.s) | Dec (dd:mm:ss) | \(v_{\text{LSR}}\) (km s\(^{-1}\)) | \(d\) (kpc) | \(T_{\text{kin}}\) (K) | \(^{12}\text{C}/^{13}\text{C}\) |
|-----------------|-----------------|----------------|-----------------------------------|-------------|---------------------|-----------------------------|
|                 |                 |                |                                   |             |                     |                             |
| HMSC            |                 |                |                                   |             |                     |                             |
| G028-C1         | 18:42:46.9      | −04:04:08      | 78.3                              | 5.0         | 17                  | 37.3                        |
| G034-F2         | 18:53:16.5      | +01:26:10      | 57.7                              | 3.7         | −                   | 45.4                        |
| G034-G2         | 18:56:50.0      | +01:23:08      | 43.6                              | 2.9         | −                   | 50.4                        |
|                 |                 |                |                                   |             |                     |                             |
| HMSC/HMPO       |                 |                |                                   |             |                     |                             |
| AFGL5142        | 05:30:48.0      | +33:47:54      | −3.9                              | 1.8         | 25*                 | 57.2                        |
| IRAS05358+3543  | 05:39:13.0      | +35:45:51      | −17.6                            | 1.8         | 35*                 | 57.2                        |
|                 |                 |                |                                   |             |                     |                             |
| HMPO            |                 |                |                                   |             |                     |                             |
| 18089-1732      | 18:11:51.4      | −17:31:28      | 32.7                              | 3.6         | 38                  | 46.0                        |
| 18517+0437      | 18:54:14.2      | +04:41:41      | 43.7                              | 2.9         | −                   | 50.4                        |
| G75-core        | 20:21:44.0      | +37:26:38      | 0.2                               | 3.8         | 96                  | 44.8                        |
|                 |                 |                |                                   |             |                     |                             |
| UC H II         |                 |                |                                   |             |                     |                             |
| G5.89-0.39      | 18:00:30.5      | −24:04:01      | 9.0                               | 1.3         | −                   | 60.4                        |
| I19035-11 VLA1  | 19:06:01.5      | +06:46:35      | 32.4                              | 2.2         | 39                  | 54.7                        |
| I9410+2336      | 19:43:11.4      | +23:44:06      | 22.4                              | 2.1         | 21                  | 55.3                        |

Notes. The distances and kinetic temperatures were adopted from Fontani et al. (2011). The asterisks indicate that we used the average \(T_{\text{kin}}\) values of the HMSCs and HMPOs within our APEX beam.

Belitsky et al. (2018), as part of the ESO E-095.F-9808A, E-096.C-0484A, and E097.C-0897A projects between July 2015 and June 2016. The first results based on the science verification dataset for AFGL5142, IRAS05358+3543, and G5.89-0.39 are presented in Zahorecz et al. (2017). We carried out single-pointing observations of HDCO and D\(_2\)CO emission using the position switching observing mode. The J2000 central coordinates used in our observations are shown in Table 1. The observed transitions of HDCO (at 185 GHz and 193 GHz) and D\(_2\)CO (at 174 GHz) are shown in Table 2. The pointing was checked every 60–90 min, and the typical system temperatures were 150 K. The beam size at 183 GHz was \(\sim 0.3\) arcsec. The spectra were obtained and calibrated in antenna temperature units and then converted to the main beam brightness temperature using \(T_{\text{MB}}\) of 0.68. The XFFTS spectrometer provided a velocity resolution of 0.112 km s\(^{-1}\), 0.059 km s\(^{-1}\), and 0.066 km s\(^{-1}\) for the H\(_2\)\(^{13}\)CO (at 206 GHz), HDCO (at 185–193 GHz), and D\(_2\)CO (at 174 GHz) transitions, respectively. However, for the data analysis, we smoothed the spectra to a uniform velocity resolution of 0.68. At this velocity resolution, we reached an rms of \(\sim 0.01\) K with typical integration times between 32 and 48 min. The PWV was 0.3–1.5 mm. The data were reduced and analysed with the Gildas software\(^2\) (Pety 2005).

2.4. ALMA Cycle 5 observations

We performed high-resolution single pointing observations with ALMA in Cycle 5 as part of the 2017.1.01157.S project (PI: S. Zahorecz). We used the Atacama Compact Array (ACA, Morita Array; Iguchi et al. 2009). Band 6 observations were executed for G028-C1, G34-F2, G34-G2, 18089-1732, 18517+0437, G5.89-0.39, 1935+VLA1, and I941+2336. We used two setups to observe selected molecular lines, including H\(_2\)\(^{13}\)CO, and to observe the continuum emission. Table 2 contains the basic information of the observed H\(_2\)\(^{13}\)CO transition. In this paper, we use the continuum and H\(_2\)\(^{13}\)CO emission information at 213 GHz to derive the source sizes. A detailed analysis of all the detected lines will be presented in a following paper. A continuum sensitivity of at least ~20 mJy beam\(^{-1}\) was achieved for the targets. For the three HMSC cores, a higher sensitivity of ~3 mJy beam\(^{-1}\) was achieved. For the molecular line observations, a sensitivity of 0.1 Jy beam\(^{-1}\) was achieved at a velocity resolution of 0.086 km s\(^{-1}\).

3. Results

In this section, we present the line and continuum emission detected towards our targets. We use the ALMA Band 6 data to estimate the source sizes. Our calculations of the excitation temperature and column density of formaldehyde and its deuterated species are based on the single-dish APEX and IRAM spectra. We also present the column density ratios of the detected molecules.

3.1. Line spectra of deuterated \(H_2CO\) species

In Fig. 1, we present the \(H_2CO\), HDCO, and D\(_2\)CO 3\(_{0,3}\) – 2\(_{0,2}\) lines and the H\(_2\)\(^{13}\)CO 3\(_{1,2}\) – 2\(_{1,1}\) line towards our sources observed with the IRAM-30m and APEX telescopes. Six sources show multi-velocity components; we performed the calculations separately for them. Neither HDCO nor D\(_2\)CO emission were detected towards the HMSC sources. The HDCO lines (upper energy levels of 18.5–50.4 K) are clearly detected in seven sources (all sources except the HMSC sources and I9035+VLA1). The D\(_2\)CO 3\(_{0,3}\) – 2\(_{0,2}\) transition is only detected clearly towards four sources: the two HMSC/HMPO sources (AFGL5142 and IRAS05358+3543), one HMPO object.

\(^2\) See http://www.iram.fr/IRAMFR/GILDAS
Fig. 1. Spectra of the $H_2CO$ $3_0,3-2_0,2$ and $H^{13}CO$ $3_1,2-2_1,1$ lines (top two panels) observed with the IRAM 30 m telescope (Fontani et al. 2011, 2015) and HDCO $3_0,3-2_0,2$ and $D_2CO$ $3_0,3-2_1,2$ lines (bottom two panels) observed with APEX SEPIA Band 5 receiver (Zahorecz et al. 2017, and this work). Red lines indicate the fitted spectra. Horizontal lines indicate the $T_{MB}=0$ K and the 3σ level. The vertical line indicates the $V_{LSR}$ for each source. The sources are sorted by the evolutionary stage; the order is the same as their order in Table 1.

We tentatively detected the same line towards one more HMPO object, G75-core, and one more UC H II region, I19035-VLA1, but further observations are needed to confirm these detections. We used only the clear detections in our calculations and considered the rest as upper limits.

We checked the $H^{13}CO$ $3_0,3-2_0,2$ and continuum emission of our sources in our Band 6 ALMA data. The continuum emission was detected for six objects out of the eight sources covered in our observations. We observed, but did not detect, compact continuum sources above 3σ towards two sources in the youngest, HMSC evolutionary stage (G034-F2 and G034-G2) at our sensitivity and resolution. We note that Kong et al. (2017) observed these objects and detected 1.3 mm continuum emission of $\sim$1–1 mmJy beam$^{-1}$ with a beam size of $1.5'' \times 1.0''$. The $H^{13}CO$ emission was not detected towards any of the HMSC sources. For the objects with detection, the $H^{13}CO$ emission is associated with the same region as the continuum emission. Figure 2 shows the continuum maps at 213 GHz and the $H^{13}CO$ contours overlaid. We derived the effective radius of the sources based on the 2D Gaussians fitted to the continuum maps. The derived source sizes are between $3''$ and $5.8''$. We note that in Zahorecz et al. (2017), we used $6''$ as the source size for AFGL5142, IRAS05358+3543, and G5.89-0.39. In our actual calculations, we updated the size of G5.89-0.39 based on its ALMA continuum emission. AFGL5142 and IRAS05358+3543 have no ALMA data and G028-C1, G034-F2, and G034-G2 have no $H^{13}CO$ detection and/or continuum detection. We kept the assumption of a $6''$ source size for these five objects in our calculations.

3.2. Molecular column densities and column density ratios

To estimate the molecular column densities from the APEX and IRAM spectra of $H_2CO$, $H^{13}CO$, HDCO, and $D_2CO$, we used the MADCUBA software that assumes LTE conditions, thus neglecting the effects of temperature and magnetic field. The software provides the column density of each molecular line, which can then be used to calculate the ratio of the column densities of different molecules. These ratios can provide insights into the physical conditions in the source, such as the pressure, temperature, and density of the gas. The column density ratios can be used to infer the evolutionary stage of the source, as different molecules are expected to be more abundant in different stages of formation. Additionally, the column density ratios can be used to study the chemical composition of the source, as the abundance of different molecules can provide information on the chemical processes occurring in the source.

The column density ratios can be calculated using the following equation:

$$\frac{N_{H^{13}CO}}{N_{H_2CO}} = \frac{A_{H^{13}CO}}{A_{H_2CO}} \frac{T_{H^{13}CO}}{T_{H_2CO}} \frac{\lambda_{H^{13}CO}}{\lambda_{H_2CO}}$$

where $N$ is the column density, $A$ is the transition probability, $T$ is the temperature, and $\lambda$ is the wavelength. The equation assumes LTE conditions and neglects the effects of temperature and magnetic field. The column density ratios can be used to study the chemical composition of the source, as the abundance of different molecules can provide information on the chemical processes occurring in the source.
Table 2. Targeted transitions and beam sizes, beam coupling efficiencies, and spectral resolution of our IRAM 30 m, APEX SEPIA, and ALMA Band 6 observations.

| Transition    | Frequency (MHz) | $E_u$ (K) | Telescope | $B_{eff}$ (Km s$^{-1}$) | $\Delta v$ (Km s$^{-1}$) | FWHM ($''$) |
|---------------|----------------|-----------|-----------|-------------------------|--------------------------|--------------|
| $^{13}$CO | 3$_0$ - 2$_2$  | 218222.192  | 10.5  | IRAM | 0.62 | 0.26 | 11 |
| 3$_1$ - 2$_0$ | 218760.066 | 57.6 | IRAM | 0.62 | 0.26 | 11 |
| 3$_2$ - 2$_1$ | 218475.632  | 57.6 | IRAM | 0.62 | 0.26 | 11 |
| HD$^{13}$CO | 3$_0$ - 2$_2$ | 212811.184 | 10.2 | ALMA | 0.75 | 0.09 | 5 - 8 |
| 3$_1$ - 2$_1$ | 219908.525  | 22.4  | IRAM | 0.62 | 0.26 | 11 |
| 3$_2$ - 2$_2$ | 206131.626  | 21.7  | APEX | 0.72 | 0.06 | 30 |
| HDCO | 3$_0$ - 2$_2$ | 192893.256 | 9.3  | APEX | 0.72 | 0.06 | 32 |
| 3$_1$ - 2$_1$ | 185307.113  | 16.9  | APEX | 0.72 | 0.06 | 34 |
| 3$_2$ - 2$_0$ | 193907.460  | 41.1  | APEX | 0.72 | 0.06 | 32 |
| 3$_2$ - 2$_1$ | 193391.605  | 41.1  | APEX | 0.72 | 0.06 | 32 |
| D$_2$CO | 3$_0$ - 2$_2$ | 174413.115 | 8.4  | APEX | 0.72 | 0.07 | 36 |

Fig. 1. continued.
Fig. 2. 213 GHz continuum maps towards G028-C1, G034-F2, G034-G2, 18089-1732, 18517+0437, G5.89-0.39, I19035-VLA1, and 19410+2336 (sorted by evolutionary stage). The black contours indicate the 3, 5, 10, and 15 $\sigma$ levels of $^{13}$CO 3–2, integrated intensity, where $\sigma$ is 0.1 Jy beam$^{-1}$. No $^{13}$CO 3–2 emission was detected towards G028-C1. Neither continuum nor $^{13}$CO 3–2 emission was detected towards the cores in the G034 region.
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4. Discussion

In this section we compare our results with those inferred from previous observational studies (Sect. 4.1). We also put our results in context with theoretical studies of formaldehyde deuteration via grain-surface and gas-phase processes (Sect. 4.2).

4.1. Comparison with previous observational studies

Fontani et al. (2011) presented the kinetic temperatures based on ammonia data. Nine sources from our sample have available ammonia temperatures with values lying between \( \sim 20–40 \) K, except for the G75-core, which has a higher kinetic temperature of 96 K. The HDCO-based temperature estimates are in good agreement with the ammonia temperatures for most of the sources: they are in the 15–40 K range. G5.89-0.39 and I19035-VLA1 show higher HDCO excitation temperatures, that is \( \sim 60–70 \) K. For the youngest HMSC sources, we only detected H\(_2\)CO. The excitation temperatures fall into the 20-30 K range. For some of the sources, the HDCO derived temperatures were too low to fit the line intensity ratios of the observed H\(_2\)CO
transitions. Also in these cases, we derived the H$_2$CO excitation temperature, finding values a factor of 2–3 higher than the HDCO values in the range of ~70–130 K.

Fontani et al. (2015) have proposed that in high-mass star-forming regions, the different trend for the deuteration pattern of different molecules, such as N$_2$H$^+$, HNC, NH$_3$, and CH$_3$OH as a function of the evolutionary stage, is likely due to the way deuteration occurs for the different species: for N$_2$H$^+$, deuteration occurs in the gas phase, while for CH$_3$OH it occurs on grain surfaces; and for NH$_3$, deuteration likely occurs via a mixture of these two processes. In the following, we discuss the likelihood of each of these two chemical routes to drive the formation of the deuterated forms of H$_2$CO during the massive star formation process.

### 4.2. Grain-surface deuteration of H$_2$CO

Traditionally, it has been assumed that the deuterated forms of H$_2$CO are a product of grain-surface chemistry. Indeed, Taquet et al. (2012) showed that the gas-phase formation timescales of D$_2$CO are longer than the depletion timescales in the prestellar phase and, therefore, one would expect D$_2$CO to be a product of material processed on solid ices. Consequently, the N(D$_2$CO)/N(H$_2$CO) ratio should show an evolutionary trend similar to that derived for D$_{\text{tot}}$(CH$_3$OH), which peaks at the HMPO phase (Fontani et al. 2015). If HDCO showed a similar behaviour as D$_2$CO, we could conclude that HDCO also forms mostly on dust grains. This hypothesis would be supported by the fact that HDCO is detected in hot cores and hot corinos while N$_2$D$^+$ is not (see e.g. Fuente et al. 2005).

Figure 3 shows that the column density ratios of N(HDCO)/N(H$_2$CO) and N(D$_2$CO)/N(H$_2$CO) decrease from the HMSC-HMPO phase to the UC H II phase by factors of $\leq$10. We note that for the two sources in the HMSC/HMPO transitional category (AFGL5142 and IRAS05358+3543), we cannot disentangle the contribution of the HMSC cores from the HMPO ones. However, from our observations it is clear that they show higher column density ratios than those found for the UC H II regions, confirming the decrease of the N(HDCO)/N(H$_2$CO) and N(D$_2$CO)/N(H$_2$CO) trend. Based on the available upper limits for the HMSC cores, we cannot exclude the possibility that the D$_2$CO/H$_2$CO column density ratio peaks in the early phases. However, the upper limits obtained for the HMSC cores suggest lower column density ratios than those of the HMPO objects, or at most equal HDCO/H$_2$CO column density ratios. This behaviour is similar to the trend observed for D$_{\text{tot}}$(CH$_3$OH), but with a larger scatter. This suggests that grain surface chemistry may be the main formation mechanism for HDCO and D$_2$CO.

Rodgers & Charnley (2002) proposed a grain surface formation scenario for these molecules in which H$_2$CO and its deuterated forms represent intermediate steps in the formation of CH$_3$OH and deuterated-CH$_3$OH via hydrogenation and D-addition reactions. They defined the F parameter (see Eq. (1)) to investigate the extent by which surface chemistry is responsible for the production of HDCO and D$_2$CO. Rodgers & Charnley (2002) found that gas-phase chemistry should produce F values between 1.6 and 2.3, and grain surface chemistry should result in F values $\geq$1. Previous observations, however, have reported F values lower than 1, which are not consistent with the cases presented above. Alternatively, low F values could be obtained through a grain surface chemistry over a long time period (Rodgers & Charnley 2002), because longer timescales allow the conversion of HDCO into D$_2$CO on the surface of dust grains. We derived the F parameter based on our calculated HDCO/H$_2$CO and D$_2$CO/H$_2$CO ratios using the definition of Rodgers & Charnley (2002):

$$F = \frac{[\text{HDCO}/\text{H}_2\text{CO}]}{[\text{D}_2\text{CO}/\text{H}_2\text{CO}]}.$$  

In our sample sources with D$_2$CO detections, F lies between 0.1 and 1.1 (except 18517+0437, where the two fitted components show F values ~3 and ~0.001). For the sources with only upper limits for the detection of D$_2$CO, the lower limits for the F values are within 0.1 and 0.6. These values cannot be explained by gas phase chemistry, but they are more consistent with grain surface chemistry models.

Table 3 also shows that the D$_2$CO/HDCO ratio remains constant at ~0.2, within the uncertainties, regardless of the source or evolutionary stage (except for 18517+0437). As for the low-mass regime, the ratio HDCO/D$_2$CO lies well below the statistical value given as D-species/D$_2$-species $= 4 \times$ (D-species/H-species$^{-1}$, which is consistent with the grain surface formation scenario (see Ceccarelli et al. 2014). Ceccarelli et al. (2014) calculated the column density ratios for HDCO/H$_2$CO and D$_2$CO/H$_2$CO for a sample of low-mass star-forming cores (see their Fig. 5). The derived ratios lie close to 1 for those objects. The average derived HDCO/H$_2$CO ratio is $\sim$5.3 with a standard deviation of 3.4 for our high-mass...
star-forming regions, and 1.1 if we exclude the two velocity components of the 18517+0437 region, showing a large difference from the other sources. If we compare these findings with the results of Ceccarelli et al. (2014), it is possible that the higher values measured towards our sample are the result of the shorter timescales available in the high-mass regime for the formation of \( \text{D}_2\text{CO} \) in the ices (Taquet et al. 2012; Rodgers & Charnley 2002). In fact, the \( \text{HDCO}/\text{H}_2\text{CO} \) ratios are factors \(~10\) or more larger than \( \text{D}_{\text{HDCO}}(\text{CH}_3\text{OH}) \) in high-mass star-forming regions, which is also consistent with the shorter timescales available for deuteration in high-mass star formation with respect to the low-mass case, and with the formation of these species via hydrogenation and D-addition reactions on grain surfaces (Turner 1990; Rodgers & Charnley 2002).

Turner (1990) reported the first interstellar detection of a multiply deuterated molecule, \( \text{D}_2\text{CO} \), in the Orion KL ridge component with a \( \text{D}_2\text{CO}/\text{H}_2\text{CO} \) column density ratio of \(~0.02\). The calculated \( \text{HDCO}/\text{H}_2\text{CO} \) column density ratio was \(~0.1\). These ratios could not be reproduced by ion-molecule gas phase reactions, but they can be fitted by the model of grain surface reactions of Tielen (1983). Additional studies towards dark clouds TMC-1 and L183 were performed by Turner (2001) and they reported \( \text{HDCO}/\text{H}_2\text{CO} \) column density ratios of \(~0.059\) and \(~0.068\), respectively.

In addition, Fontani et al. (2014) successfully detected \( \text{HDCO} \) towards the protostellar bow-shock L1157-B1 where material from icy mantles is thought to be freshly released due to mantle evaporation. They have found a column density ratio of \( \text{HDCO}/\text{H}_2\text{CO} \) value of \(~0.1\), which is similar to the values found in our study. Bianchi et al. (2017) studied the deuteration in Sun-like protostars. Their Fig. 9 shows measured column density ratios of \( \text{HDCO}/\text{H}_2\text{CO} \) as a function of the bolometric luminosity for both low-mass and high-mass star-forming regions from the protostellar phase to the Class I phase. They have also found that the average column density ratio measured in Class 0 sources for \( \text{HDCO} \) is \(~0.12\). However, \( \text{D}_2\text{CO} \) shows an increase going from pre-stellar cores, with an average value of \(~0.045\), to Class 0 sources \( (~0.15) \) and then a strong decrease in the Class I protostar phase (see the case of SVS13-A; Bianchi et al. 2017). Our results indicate a similar, but not identical, evolutionary behaviour since no clear detection of deuterated formaldehyde is available for the HMSC sources.

As part of the ALMA-PILS survey, Persson et al. (2018) and Manigand et al. (2020) have reported the deuteration fraction of formaldehyde for IRAS 16293-2422 A and B. Their relative evolutionary stages are under debate, although recent studies suggest that the B component is at an early evolutionary stage (van der Wiel et al. 2019). Formaldehyde abundance is lower towards the A component with a \( \text{HDCO}/\text{H}_2\text{CO} \) column density ratio of around \(~0.049\) and \(~0.065\) for the A and B components, respectively. They agree well with our results for the HMPO sources. Persson et al. (2018) and Manigand et al. (2020) have also found that the doubly deuterated isotopologue \( \text{D}_2\text{CO} \) is significantly enhanced compared to the singly deuterated isotopologue towards the B component. The calculated \( \text{D}_2\text{CO}/\text{H}_2\text{CO} \) column density ratios are \(~0.041\) and \(~0.006\), which are similar to the values obtained for our HMPO sources. For the B source, the column density ratio of \( \text{D}_2\text{CO}/\text{HDCO} \) is slightly larger than the column density ratio \( \text{HDCO}/\text{H}_2\text{CO} \), which is consistent with formaldehyde forming in the ice as soon as \( \text{CO} \) is frozen onto grains. Figure 4 shows the comparison of the observed \( \text{H}_2\text{CO} \), \( \text{HDCO} \), and \( \text{D}_2\text{CO} \) column density ratios in different environments based on the works mentioned in this section.

### 4.3. Gas-phase deuteration of \( \text{H}_2\text{CO} \)

Alternatively to the grain-surface formation of \( \text{HDCO} \) and \( \text{D}_2\text{CO} \), the deuterated forms of \( \text{H}_2\text{CO} \) could be produced by gas-phase chemistry. Roberts & Millar (2007) and Roueff et al. (2007) proposed that deuterium fractionation of \( \text{H}_2\text{CO} \) may occur via the fractionation of \( \text{CH}_3\text{D}^+ \) \( \rightarrow \text{CH}_2\text{D}^+ \rightarrow \text{CH}_2\text{D} \rightarrow \text{HDCO} \) or of \( \text{H}_2\text{CO} \) directly via the reaction with \( \text{H}_2\text{D}^+ \). In particular, in the gas-phase steady-state models of Bergman et al. (2011) based on the reaction network of Roueff et al. (2007), these authors also predict a decrease in the deuteration of \( \text{H}_2\text{CO} \) from the HMSC/HMPO stage to the UC H II region phase as the temperature exceeds \(~40\) K. The predicted values for the \( \text{D}/\text{H} \) ratio inferred from the \( \text{HDCO}/\text{H}_2\text{CO} \) and \( \text{D}_2\text{CO}/\text{HDCO} \) ratios are indeed consistent with our observations (see their Fig. 12). Indeed, while the \( \text{HDCO}/\text{H}_2\text{CO} \) ratio gives \(~0.1\) for \( T \sim 10–30\) K and \(~0.04\) for \( T \sim 40\) K, the \( \text{D}_2\text{CO}/\text{H}_2\text{CO} \) ratio remains relatively flat at \(~0.2\) for \( T \sim 10–40\) K, as observed in our sample. The only difference between our observations and the predictions by Bergman et al. (2011) is that, while in our observations the \( \text{HDCO}/\text{H}_2\text{CO} \) ratio drops by more than one order of magnitude from the HMPO to the UC H II region stage, the chemical modelling predicts a smaller drop by only factors from \(~3\)–\(~4\). Parise et al. (2009) found high deuteron fractionation towards the Orion Bar PDR region. The calculated deuteron fractions are consistent with predictions of pure gas-phase chemistry models at warm temperatures. They have found an \( (\text{HDCO})/(\text{H}_2\text{CO}) \) ratio of \(~0.0057\). This value is in the range found for the UC H II regions in our sample.

In any case, to distinguish between the grain-surface from the gas-phase scenarios, we would need to obtain stringent upper limits to the \( \text{HDCO}/\text{H}_2\text{CO} \) ratios at the HMSC stage. From our observations, these limits can get as low as \(~\leq0.1\) when considering the rms measured over the whole linewidth of the lines, which is still consistent with the gas-phase results from the models of Bergman et al. (2011) (grain-surface deuteration is expected to give much lower upper limits as a consequence of freeze-out in the cold conditions of HMSC). Therefore, although our results suggest that grain surface deuteration is the most likely mechanism for the formation of \( \text{HDCO} \) and \( \text{D}_2\text{CO} \) in star-forming regions, higher sensitivity observations are needed to provide more stringent constraints to the column density ratio of \( \text{HDCO}/\text{D}_2\text{CO} \) during the HMSC phase.

### 5. Summary

We have observed \(~213\) GHz continuum, \( \text{H}_3\text{C}_2\text{O} \), \( \text{HDCO} \), and \( \text{D}_2\text{CO} \) emission with APEX and ALMA towards a sample of high-mass-star-forming regions at different evolutionary stages. HDCO transitions were detected for all of them except the three youngest sources in the HMSC stage, while the \( \text{D}_2\text{CO} \) line was detected only for four sources from the HMSC-HMPO stage to the UC H II stage. The \( \text{H}_3\text{C}_2\text{O} \) emitting regions are similar to the continuum source sizes; they are in the range of \(~3''–5.8''\) corresponding to \(~0.02–0.07\) pc at the distance of the investigated regions. The column density ratio of \( \text{HDCO}/\text{H}_2\text{CO} \) shows the highest values of \(~0.13\) in the earlier evolutionary stages (HMSC-HMPO) and then decreases by an order of magnitude towards the UC H II phase to \(~0.01\). This study confirms the idea that \( \text{H}_2\text{CO} \), and its deuterated species, form mostly on grain surfaces although some gas-phase contribution is expected at the warm HMPO stage. The calculated upper limits for the HMPO sources are high, around \(~0.5\), so we cannot constrain the trend between HMSC and HMPO phases. More sensitive follow-up
Fig. 4. Comparison of the observed $N$(D$_2$CO)/$N$(H$_2$CO) (top panel) and $N$(D$_2$CO)/$N$(H$_2$CO) (bottom panel) column density ratios for our sources and for prestellar cores (Turner 2001; Bacmann et al. 2003; Chacón-Tanarro et al. 2019), low-mass protostars (Parise et al. 2006; Ceccarelli et al. 2014; Bianchi et al. 2017; Fontani et al. 2014; Manigand et al. 2020), and sources in the Orion region (Turner 1990; Parise et al. 2009).

observations are needed for the HMSC sources to better constrain the upper limits. The observed column density ratios are in good agreement with the values previously reported in the literature for low-mass prestellar cores, protostellar objects, and PDR regions.

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## Appendix A: Observed H$_2$CO, HDCO, and D$_2$CO lines

In Table A.1 and we report the parameters derived fitting single or multiple velocity components to the detected lines with MADCUBA.

### Table A.1. Fitted H$_2$CO, H$_2$CO, HDCO, and D$_2$CO excitation temperatures and column densities.

| Source     | Size $\prime\prime$ (pc) | Species | $N$ (10$^{13}$ cm$^{-2}$) | $T_{ex}$ (K) | $v_{LSR}$ (km s$^{-1}$) | $T_{N_{H_2}}$ (K) |
|------------|--------------------------|---------|---------------------------|-------------|--------------------------|------------------|
| **HMSC**   |                          |         |                           |             |                          |                  |
| G028-C1    | 6.0 (0.15)               | H$_2$CO | 2.72 ± 0.34               | 30.2 ± 4.9  | 80.1 ± 0.1               | 17               |
|            | 6.0 (0.15)               | HDCCO   | <0.81                     | 30.2 ± 4.9  | 80.1 ± 0.1               | 17               |
|            | 6.0 (0.15)               | D$_2$CO | <0.99                     | 30.2 ± 4.9  | 80.1 ± 0.1               | 17               |
| G034-F2    | 6.0 (0.11)               | H$_2$CO | 1.23 ± 0.30               | 25.2 ± 8.2  | 58.9 ± 0.5               |                  |
|            | 6.0 (0.11)               | HDCCO   | <0.63                     | 31.2 ± 10.8 | 56.8 ± 0.5               |                  |
|            | 6.0 (0.11)               | D$_2$CO | <0.50                     | 30.5        |                         |                  |
| G034-G2    | 6.0 (0.08)               | H$_2$CO | 1.48 ± 0.12               | 19.0 ± 4.8  | 42.0 ± 0.2               |                  |
|            | 6.0 (0.08)               | HDCCO   | <0.50                     | 29.6 ± 12.8 | 43.8 ± 0.2               |                  |
| **HMSC/HMPO** |                     |         |                           |             |                          |                  |
| AFGL5142   | 6.0 (0.05)               | H$_2$CO | 876.51 ± 57.11            | 13.6 ± 0.9  | −1.7 ± 0.1               | 25               |
|            | 6.0 (0.05)               | H$_2$CO | 401.61 ± 14.18            | 22.4 ± 6.4  | −1.8 ± 0.1               | 25               |
|            | 6.0 (0.05)               | H$_2$CO | 2.03 ± 0.14               | 13.6 ± 0.9  | −1.2 ± 0.2               | 25               |
|            | 6.0 (0.05)               | H$_2$CO | 1.17 ± 0.10               | 22.4 ± 6.4  | −1.3 ± 0.2               | 25               |
|            | 6.0 (0.05)               | HDCCO   | 18.79 ± 3.82              | 13.6 ± 0.9  | −1.2 ± 0.2               | 25               |
|            | 6.0 (0.05)               | D$_2$CO | 7.03 ± 0.99               | 22.4 ± 6.4  | −1.3 ± 0.2               | 25               |
|            | 6.0 (0.05)               | D$_2$CO | 2.88 ± 0.32               | 13.6 ± 0.9  | −1.2 ± 0.2               | 25               |
|            | 6.0 (0.05)               | D$_2$CO | 1.89 ± 2.42               | 22.4 ± 6.4  | −1.3 ± 0.2               | 25               |
| IRAS05358+3543 |           |         |                           |             |                          |                  |
|            | 6.0 (0.05)               | H$_2$CO | 100.51 ± 3.68             | 71.9 ± 3.3  | −1.6 ± 0.2               | 35               |
|            | 6.0 (0.05)               | H$_2$CO | 1.96 ± 0.14               | 27.2 ± 6.0  | −1.6 ± 0.2               | 35               |
|            | 6.0 (0.05)               | HDCCO   | 13.69 ± 1.75              | 27.2 ± 6.0  | −1.6 ± 0.2               | 35               |
|            | 6.0 (0.05)               | D$_2$CO | 2.46 ± 0.28               | 27.2 ± 6.0  | −1.6 ± 0.2               | 35               |
| **HMPO**   |                          |         |                           |             |                          |                  |
| 18089-1732 | 3.6 (0.06)               | H$_2$CO | 360.46 ± 7.49             | 84.7 ± 1.5  | 35.0 ± 0.5               | 38               |
|            | 3.6 (0.06)               | H$_2$CO | 184.09 ± 5.28             | 80.8 ± 2.4  | 32.3 ± 0.5               | 38               |
|            | 3.6 (0.06)               | H$_2$CO | 8.91 ± 0.65               | 18.7 ± 11.8 | 34.2 ± 0.2               | 38               |
|            | 3.6 (0.06)               | H$_2$CO | 4.42 ± 0.49               | 22.5 ± 17.4 | 31.9 ± 0.2               | 38               |
|            | 3.6 (0.06)               | HDCCO   | 7.28 ± 1.76               | 18.7 ± 11.8 | 34.2 ± 0.2               | 38               |
|            | 3.6 (0.06)               | D$_2$CO | 6.04 ± 2.63               | 22.5 ± 17.4 | 31.9 ± 0.2               | 38               |
|            | 3.6 (0.06)               | D$_2$CO | <0.39                    | 20          |                          | 38               |
| 18517+0437 | 5.3 (0.07)               | H$_2$CO | 137.19 ± 16.22            | 22.6 ± 6.5  | 44.2 ± 0.5               |                  |
|            | 5.3 (0.07)               | H$_2$CO | 20.59 ± 3.03              | 23.8 ± 9.2  | 45.1 ± 0.5               |                  |
|            | 5.3 (0.07)               | H$_2$CO | 1.03 ± 0.48               | 22.6 ± 6.5  | 44.0 ± 0.1               |                  |
|            | 5.3 (0.07)               | H$_2$CO | 2.66 ± 0.48               | 23.8 ± 9.2  | 44.6 ± 0.2               |                  |
|            | 5.3 (0.07)               | HDCCO   | 15.73 ± 2.00              | 22.6 ± 6.5  | 43.9 ± 0.1               |                  |
|            | 5.3 (0.07)               | HDCCO   | 0.13 ± 0.02               | 23.8 ± 9.2  | 45.7 ± 0.5               |                  |
|            | 5.3 (0.07)               | D$_2$CO | 1.53 ± 0.47               | 22.6 ± 6.5  | 43.8 ± 0.5               |                  |
|            | 5.3 (0.07)               | D$_2$CO | 0.72 ± 0.33               | 23.8 ± 9.2  | 44.9 ± 0.5               |                  |
| G75-core   | 6.0 (0.11)               | H$_2$CO | 152.4 ± 4.79              | 34.9 ± 12.3 | 0.5 ± 0.1                | 96               |
|            | 6.0 (0.11)               | D$_2$CO | 16.79 ± 2.18              | 34.9 ± 12.3 | −0.4 ± 0.2               | 96               |
|            | 6.0 (0.11)               | D$_2$CO | 5.12 ± 0.11               | 34.9 ± 12.3 | −0.1 ± 0.1               | 96               |
|            | 6.0 (0.11)               | HDCCO   | 12.18 ± 2.89              | 34.9 ± 12.3 | −0.1 ± 0.2               | 96               |
|            | 6.0 (0.11)               | D$_2$CO | <1.26                    | 34.9 ± 12.3 |                      | 96               |

**Notes.** We note that to estimate the upper limits of the HDCO and/or D$_2$CO column densities for G034-F2, G034-G2, 18089-1732, and G5.89-0.39, we used one component only and we used an average excitation temperature based on the successful fits of H$_2$CO/HDCO. No error bar is provided for $T_{ex}$ in these cases. Source velocities and sizes are also indicated. The last column shows the kinetic temperatures adopted from Fontani et al. (2011).
### Table A.1. continued.

| Source          | Size (arcsec) | Species | $N$ (10$^{13}$ cm$^{-2}$) | $T_{ex}$ (K) | $v_{LSR}$ (km s$^{-1}$) | $T_{NH_3}$ (K) |
|-----------------|---------------|---------|---------------------------|--------------|------------------------|----------------|
|                 | (pc)          |         |                           |              |                        |                |
|                 |               |         | UC H II                   |              |                        |                |
| G5.89-0.39      | 3.0 (0.02)    | H$_2$CO | 8062 ± 732                | 132.6 ± 10.5 | 11.5 ± 0.2             |                |
|                 | 3.0 (0.02)    |         | 2667 ± 567                | 86.0 ± 7.6   | 8.4 ± 0.2              |                |
|                 | 3.0 (0.02)    | H$^{13}$CO | 130.30 ± 12.32            | 70.0 ± 7.5   | 11.6 ± 0.2             |                |
|                 | 3.0 (0.02)    | HDCO    | 31.32 ± 3.00              | 70.0 ± 7.5   | 11.7 ± 0.2             |                |
|                 | 3.0 (0.02)    |         | 182.26 ± 29.73            | 37.1 ± 3.4   | 8.4 ± 0.2              |                |
|                 | 3.0 (0.02)    | D$_2$CO | 89.08 ± 9.00              | 37.1 ± 3.4   | 8.4 ± 0.2              |                |
|                 | 4.8 (0.05)    | H$_2$CO | 92.65 ± 2.92              | 60.3 ± 2.0   | 33.1 ± 0.1             | 39             |
|                 | 4.8 (0.05)    | H$^{13}$CO | 4.10 ± 0.52              | 60.3 ± 2.0   | 32.9 ± 0.2             | 39             |
|                 | 4.8 (0.05)    | HDCO    | <4.79                     | 60.3 ± 2.0   |                        | 39             |
|                 | 4.8 (0.05)    | D$_2$CO | <3.63                     | 60.3 ± 2.0   |                        | 39             |
| I19035-VLA1     | 5.8 (0.06)    | H$_2$CO | 16.27 ± 1.34              | 60.1 ± 4.8   | 22.7 ± 0.1             | 21             |
|                 | 5.8 (0.06)    | H$^{13}$CO | 5.59 ± 0.19              | 25.4 ± 11.1  | 22.5 ± 0.1             | 21             |
|                 | 5.8 (0.06)    | HDCO    | 6.43 ± 1.82               | 25.4 ± 11.1  | 22.4 ± 0.2             | 21             |
|                 | 5.8 (0.06)    | D$_2$CO | 1.32 ± 0.28               | 25.4 ± 11.1  | 22.6 ± 0.2             | 21             |

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