



Universiteit
Leiden
The Netherlands

Trans-ethyl methyl ether in space. A new look at a complex molecule in selected hot core regions

Fuchs, G.W.; Fuchs, U.; Giesen, T.F.; Wyrowski, F.

Citation

Fuchs, G. W., Fuchs, U., Giesen, T. F., & Wyrowski, F. (2005). Trans-ethyl methyl ether in space. A new look at a complex molecule in selected hot core regions. *Astronomy And Astrophysics*, 444, 521-530. Retrieved from <https://hdl.handle.net/1887/6986>

Version: Not Applicable (or Unknown)

License: [Leiden University Non-exclusive license](#)

Downloaded from: <https://hdl.handle.net/1887/6986>

Note: To cite this publication please use the final published version (if applicable).

Trans-ethyl methyl ether in space

A new look at a complex molecule in selected hot core regions^{*}

G. W. Fuchs^{1,2}, U. Fuchs³, T. F. Giesen³, and F. Wyrowski⁴

¹ Department of Chemistry, UC Berkeley, Berkeley, CA 94720, USA

² Raymond and Beverly Sackler Laboratory for Astrophysics, Leiden Observatory, Leiden University, Postbus 9513, 2300 RA, Leiden, The Netherlands
e-mail: fuchs@strw.leidenuniv.nl

³ I. Physikalisches Institut, Universität zu Köln, Zùlpicher Str. 77, 50937 Köln, Germany
e-mail: ufuchs@ph1.uni-koeln.de, giesen@ph1.uni-koeln.de

⁴ Max-Planck-Institut für Radioastronomie, Auf dem Hügel 69, 53121 Bonn, Germany
e-mail: wyrowski@mpifr-bonn.mpg.de

Received 9 June 2005 / Accepted 22 August 2005

ABSTRACT

An extensive search for the complex molecule trans-ethyl methyl ether towards several hot core regions has been performed. Using the IRAM 30 m telescope and the SEST 15 m we looked at several frequencies where trans-ethyl methyl ether has strong transitions, as well as lines which are particularly sensitive to the physical conditions in which the molecule can be found. We included G34.26, NGC 6334(I), Orion KL, and W51e2 which have previously been proven to have a rich chemistry of complex molecules. Our observations cannot confirm the tentative Orion KL detection made by Charnley et al. (2001) within their stated column density limits, but we confirm the existence of the trans-ethyl methyl ether towards W51e2 with a column density of $2 \times 10^{14} \text{ cm}^{-2}$. The dimethyl ether/methanol ratio of 0.6 as well as the newly found ethyl methyl ether/ethanol ratio of 0.13 indicate relative high abundances of ethers toward W51e2. Furthermore, the observation of ethyl methyl ether also confirms the importance of ethanol as a grain mantle constituent. We present new upper limits of around $8 \times 10^{13} \text{ cm}^{-2}$ for the column densities of the molecule toward Orion KL, G34.26, NGC 6334(I) and estimate the column density towards SgrB2(N) to be of the same order. The W51e2 observations are discussed in more detail.

Key words. line: identification – astrochemistry – radio lines: ISM – ISM: clouds – ISM: molecules

1. Introduction

Various large and complex molecules of prebiotic importance have been found as constituents of dense interstellar clouds but their detection remains difficult and some of them doubtful (Snyder et al. 2005). Fortunately, space offers many places with a rich chemistry and hot molecular cores (HMCs) are prime targets for such investigations. However, the spectra towards hot cores often reveal a high line density and thus proper line assignments are difficult and have to be viewed with a critical eye. The ethyl methyl ether ($\text{CH}_3\text{OC}_2\text{H}_5$) is an asymmetric top molecule with two methyl groups acting as internal rotors. Ethyl methyl ether (EME) has two conformers where the trans-EME is lower in energy than the gauche-EME by 550 cm^{-1} (Durig et al. 2002) and thus favored as an astrophysically detectable species. We searched for several transitions of trans-EME to confirm the tentative detection of this molecule by

Charnley et al. (2001). A positive result means the detection of one of the biggest molecules in the interstellar medium (ISM). Charnley et al. (2001) observed a single line in W51 e1/e2 and Orion KL at 160.1 GHz and another line in SgrB2(N) at 79.6 GHz which they assigned to trans-ethyl methyl ether. Based on these isolated measurements, Charnley et al. (2001) estimate the column density of EME to be in the range of 10^{14} – 10^{15} cm^{-2} in Sgr B2(N). Thus, assuming a hydrogen column density of 10^{24} cm^{-2} gives a fractional abundance of 10^{-10} – 10^{-9} . For W51, they estimate EME to have an abundance of a few times 10^{-10} . Furthermore, they conclude that the emission of EME closely follows that of methanol, as expected if these molecules are chemically linked. Charnley et al. (1995) suggest the following production mechanism for EME in hot cores where temperatures can rise to 100–300 K. Icy mantles which are thought to contain alcohols (e.g. methanol, ethanol) evaporate from dusty grains, e.g. Millar & Hatchell (1998), the alcohols themselves are thought to be produced via surface reactions starting from the precursor CO. Once the

^{*} Table 5 is only available in electronic form at <http://www.edpsciences.org>

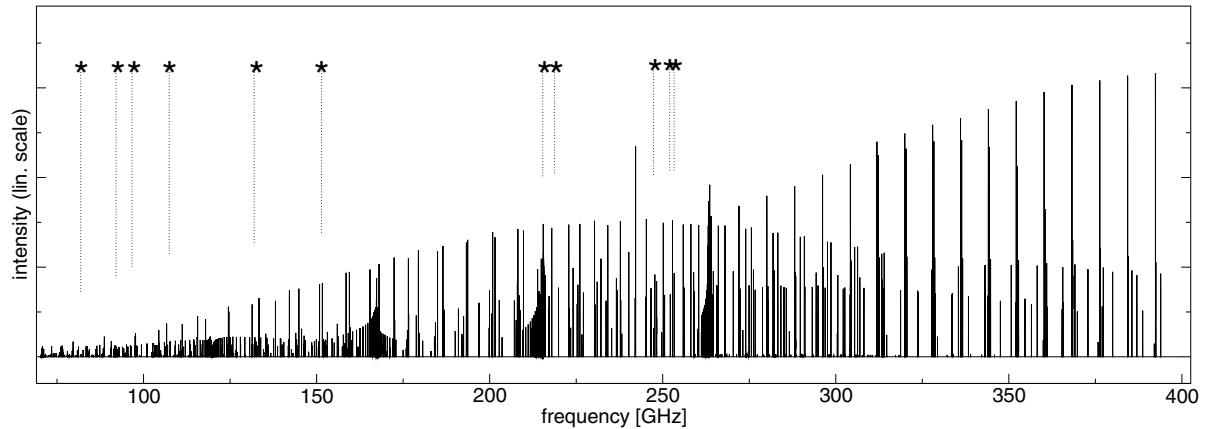
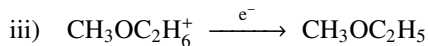
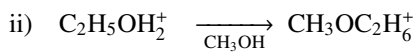
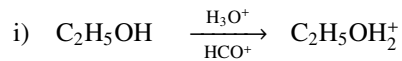


Fig. 1. Stick spectrum of EME at a rotational temperature of 130 K which is typical for complex molecules in hot core regions. EME has a dense spectrum with many characteristic features. The frequencies are taken from Fuchs et al. (2003). The (*) indicate our chosen frequency regions for the W51e2 observations. The intense line at 237 GHz was not selected because of a blending line at the same frequency.

Table 1. Astronomical sources.

Source	RA	Dec	Eq.	<i>LSR</i> (km s ⁻¹)	telescope
G34.26	18 53 18.496	01 14 58.66	2000	+60.0	IRAM 30 m
NGC 6334 I	17 17 32.300	-35 44 04.00	1950	-8.0	SEST 15 m
Orion KL	05 35 14.171	-05 22 23.10	2000	+8.8	IRAM 30 m
W51 e2	19 23 43.959	14 30 34.55	2000	+55.0	IRAM 30 m

mantles evaporate the alcohols can be converted into ethers and other species. The production of ethers is thought to happen via ion-molecule reactions¹ (i), methylation of the protonated species² (ii) and dissociative recombination with electrons (iii):



In hot cores saturated species are the dominant constituents. For instance, methanol CH_3OH is highly abundant and dimethyl ether $(\text{CH}_3)_2\text{O}$ and ethanol $\text{C}_2\text{H}_5\text{OH}$ are seen only in hot cores (Herbst 2001). Charnley et al. (1995) mention in their work that dimethyl ether, diethyl ether $(\text{C}_2\text{H}_5)_2\text{O}$, and trans-EME should be present in detectable quantities within cores rich in ethanol and methanol. Furthermore, a detection of either EME or diethyl ether would confirm the importance of ethanol as a grain mantle constituent.

2. EME spectrum and observations

Only two investigations on the laboratory rotational spectrum of EME have been reported in the literature. The more recent measurements by Fuchs et al. (2003) from 55 GHz to 350 GHz also include the previously published data below 40 GHz by Hayashi & Kuwada (1975) in their analysis. Reliable frequency predictions up to 400 GHz are now available. Since EME consists of two methyl groups designated as internal rotors, the

laboratory spectra reveal extra line splittings which are not necessarily seen in space due to larger line widths in the HMC environment. The spectroscopic aspects of ethyl methyl ether in the mm-wave region are described in our earlier publications (see Fuchs et al. 2003; Fuchs 2003, and references therein). EME is an asymmetric top molecule which has weak *a*-type transitions and fairly strong *b*-type transitions above 100 GHz which in theory split into quintets caused by internal torsion. Laboratory measurements can resolve only doublets and quartets with splittings mostly between 0.8 and 2 MHz in our examined frequency ranges. However, none of the astronomically observed lines is torsionally resolved. Figure 1 shows the complexity of the EME spectrum up to 400 GHz with its characteristic features. The IRAM 30 m telescope at Pico Veleta, Spain, has been used during several days in Sept. 2002 and July 2003 and the SEST 15 m telescope at La Silla, Chile, in March 2003. The hot core regions of G34.26, NGC 6334I, Orion KL and W51e2 were observed, see Table 1. Rotational transitions of EME have been searched in the 1 mm-, 2 mm-, and 3 mm-range. Depending on the source, between 5 and 11 frequency settings have been used, see the online table for observational details. Typical frequency resolutions were 0.3, 1 and 1.25 MHz, typical bandwidths between 490 and 1020 MHz and rms noise levels between 0.01 K at 81 GHz and 0.02–0.05 K at 245 GHz. The bands were chosen to allow for the observation of CH_3OH (*a*-type), $\text{C}_2\text{H}_5\text{OH}$ and CH_3OCH_3 transitions as well. At least 5 strong lines have been measured from each of these species, except for CH_3OH in Orion KL where only three transitions have been observed.

The intensities are measured in T_{A}^* the effective antenna temperature corrected for spillover losses and atmospheric

¹ See Mackay et al. (1979); Bohme et al. (1979).

² See Karpas & Meot-Ner (1989).

attenuation, and are then converted to $T_{\text{mb}} = T_{\text{A}}^*/\rho$ using the beam efficiencies ρ of the telescopes at the given frequencies.

All sources were examined using long integration times up to 220 min on+off time, elevations greater 40° and under good atmospheric weather conditions.

3. Source selection

A selection of previous work on the sources is given below. The authors do not intend to give a review of these sources but rather focus on contributions relevant for the EME analysis.

G34.3+0.15 has been surveyed by MacDonald et al. (1996) and Thompson et al. (1999) in the 330–360 GHz region. Nomura & Millar (2004) examined the chemical structure of this region and gave an overview of the observed species and their abundances along with some comparisons to their model predictions. Single molecule investigations like ethanol have been performed by Millar et al. (1995). Nummelin et al. (1998b) examined CH_3CHO and $\text{c-C}_2\text{H}_4\text{O}$ but also gave abundances for methanol, ethanol and dimethyl ether (DME). Heaton et al. (1989) mapped the G34.3 region using the $\text{NH}_3(3,3)$ emission and determined a size of $3'' \times 1''.2$. The column density of H_2 towards G34.26 has been estimated by Millar et al. (1995) to be $N_{\text{I}}(\text{H}_2) = 5.3 \times 10^{23} \text{ cm}^{-2}$. Nummelin et al. (1998b) published a similar result of $N_{\text{I}}(\text{H}_2) = 3.0 \times 10^{23} \text{ cm}^{-2}$ for a $20''$ source size.

NGC 6334I (NGC 6334F, G351.41+0.64) has been mapped in the continuum by Sandell (2000). McCutcheon et al. (2000) took several spectra in the range between 334 and 348 GHz using the JCMT. They also mapped the region in CH_3OCH_3 emission. Nummelin et al. (1998b) estimated the abundances of ethanol, methanol and DME in NGC 6334I and a hydrogen column density of $N_{\text{I}}(\text{H}_2) = 2.0 \times 10^{23} \text{ cm}^{-2}$ within $20''$. Cheung et al. (1978) estimate $N_{\text{I}}(\text{H}_2) \approx 10^{24} \text{ cm}^{-2}$.

Orion KL is a well studied object in the sky. Turner (1989, 1991) performed a line survey between 70 and 115 GHz, Lee & Cho (2002) presented line spectra between 159.7 to 164.7 GHz and Sutton et al. (1985) scanned the spectral region between 215 and 247 GHz. Blake et al. (1987) studied the chemical composition of OMC-1 in the 230 GHz atmospheric window. They estimate the hydrogen column density to be $N_{\text{I}}(\text{H}_2) = 1.0 \times 10^{23} \text{ cm}^{-2}$ for the hot core. The region between 330 to 355 GHz was examined by Sutton et al. (1991), and Schilke et al. (1997) performed a survey from 325 GHz to 360 GHz (see also references therein). The distribution of some key molecules for the EME analysis has been presented by Sutton et al. (1995). Caselli et al. (1993) applied a dynamical-chemical model which included abundances of many observed species.

The **W51** molecular cloud is described in Carpenter & Sanders (1998). The chemistry of the W51e2 region has been investigated by several groups. For example, Zhang et al. (1998) has done CH_3CN and $\text{CS}(3-2)$ observations, Remijan et al. (2004, 2002) investigated CH_3CN and CH_3COOH , Liu et al. (2001) looked for HCOOH in this source and Kuan et al. (2004) searched for several molecules of potential prebiotic importance. The hydrogen column density $3.6 \times 10^{23} \text{ cm}^{-2}$ (Ikeda et al. 2001) is based on C^{18}O data from Schloerb et al. (1987).

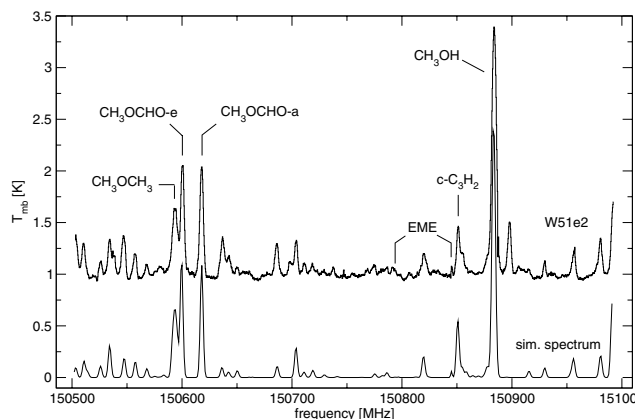


Fig. 2. Observed spectrum of W51e2 at 151 GHz (*upper*) and simulated spectrum using myXCLASS (*lower*) including $\text{C}_2\text{H}_5\text{OCH}_3$, $\text{C}_2\text{H}_5\text{OOCH}$, $\text{CH}_3\text{CH}_3\text{CO}$, $\text{a}-(\text{CH}_2\text{OH})_2$, $\text{c-C}_2\text{H}_4\text{O}$, $\text{C}_2\text{H}_5\text{OH}$, CH_3OCH_3 , $\text{CH}_3\text{CHO-e}$, CH_3OH , $\text{C}_2\text{H}_5\text{OH}$, $\text{CH}_3\text{OCHO-e}$, and $\text{CH}_3\text{OCHO-a}$.

Also the **SgrB2** region has been investigated by many groups. It is not part of our own observations but since Charnley et al. (2001) detected one line at an EME transition frequency we use the published data to estimate the column density of EME in Sect. 6. A spectral line survey from 70 and 150 GHz has been performed by Cummins et al. (1986), and from 70 to 115 GHz by Turner (1989, 1991). A further 3 mm spectral line survey has been carried out by Friedel et al. (2004) using the NRAO12 m and the BIMA array telescopes. Nummelin et al. (1998a) extended these observations to frequencies between 218 and 263 GHz. Selected molecules, like the oxygen bearing CH_3COOH , have been studied by Remijan et al. (2002), and NH_3 by Hüttemeister et al. (1993) with comparisons between several sources including those studied here. Martín-Pintado et al. (1990) estimated $N_{\text{I}}(\text{H}_2) = 5.0 \times 10^{24} \text{ cm}^{-2}$ to be the hydrogen column density³.

Ikeda et al. (2001, 2002) gave temperatures and abundances of methanol, ethanol and in most cases also for DME for all above mentioned sources. Table 2 summarizes the CH_3OH , $\text{C}_2\text{H}_5\text{OH}$, CH_3OCH_3 and $\text{C}_2\text{H}_5\text{CH}_3$ abundances including our latest results.

4. Data analysis method

For the analysis of the observed lines we used the Grenoble Image and Line Data Analysis Software GILDAS⁴ and in addition the myXCLASS code written by Peter Schilke which is backed by a database of astrophysically important molecules, including the EME data. myXCLASS can simulate the spectra of many molecules at a time, including their line profiles within the GILDAS environment and has restricted fitting capabilities. Figure 2 shows the observed spectrum of W51e2 at 151 GHz and the synthetic spectrum calculated by myXCLASS assuming optically thin lines where for each molecule the excitation

³ Note, Charnley et al. (2001) assume $N_{\text{H}_2} = 1.0 \times 10^{24} \text{ cm}^{-2}$.

⁴ See <http://www.iram.fr/IRAMFR/GILDAS>

Table 2. Rotational temperatures, column densities and fractional abundances X relative to H_2 of CH_3OH , C_2H_5OH , CH_3OCH_3 and $C_2H_5OCH_3$ (EME) from our and other selected works.

source	molecules		T_{rot} [K]	N_t [cm^{-2}]	X^a	Reference
G34.26	CH_3OH	a -type	368 ± 16	$(1.8 \pm 0.2) \times 10^{16}$	4.5×10^{-8}	MacDonald et al. (1996)
			368 ± 16	$(1.6 \pm 0.2) \times 10^{16}$	4×10^{-8}	Thompson et al. (1999) ^b
		b -type	336 ± 14	$(1.9 \pm 0.1) \times 10^{16}$	4.8×10^{-8}	MacDonald et al. (1996)
		$a + b$ type	96 ± 17	$(2.6 \pm 1.4) \times 10^{16}$	6.5×10^{-8}	Ikeda et al. (2002)
		a -type	140 ± 14	$2.2^{+4.7}_{-0.4} \times 10^{16}$	5.5×10^{-8}	this work
	C_2H_5OH		94 ± 27	$(3.5 \pm 2.0) \times 10^{15}$	8.8×10^{-9}	MacDonald et al. (1996)
			75 ± 37	$(1.7 \pm 2.1) \times 10^{15}$	4.3×10^{-9}	Ikeda et al. (2002)
			152 ± 66	$6.0^{+8.4}_{-3.5} \times 10^{14}$	1.5×10^{-9}	this work
	CH_3OCH_3		137 ± 32	$(3.7 \pm 1.8) \times 10^{15}$	9.3×10^{-9}	Ikeda et al. (2002)
			146 ± 38	$3.5^{+1.8}_{-1.2} \times 10^{15}$	8.8×10^{-9}	this work
	$C_2H_5OCH_3$		100–150	$< 7 \times 10^{13}$	$< 1.8 \times 10^{-10}$	this work
NGC 6334 I	CH_3OH		79^{+21}_{-8}	$3.1^{+0.8}_{-1.0} \times 10^{16}$	1.6×10^{-7}	Nummelin et al. (1998b)
			66 ± 11	$1.2^{+1.9}_{-1.2} \times 10^{16}$	6.0×10^{-8}	this work
	C_2H_5OH		94^{+44}_{-20}	$1.4^{+0.5}_{-0.3} \times 10^{15}$	7.0×10^{-9}	Nummelin et al. (1998b)
			120 ± 44	$4.0^{+1.2}_{-0.9} \times 10^{14}$	2.0×10^{-9}	this work
	CH_3OCH_3		20/440	$3.5 \times 10^{15} / 1.3 \times 10^{16}$	4.1×10^{-8}	Nummelin et al. (1998b)
			119 ± 39	$2.3^{+1.8}_{-1.0} \times 10^{15}$	1.2×10^{-8}	this work
	$C_2H_5OCH_3$		50–150	$< 8 \times 10^{13}$	$< 4 \times 10^{-10}$	this work
Orion KL	CH_3OH		120–140	5×10^{16}	5×10^{-7}	Sutton et al. (1985)
			140	4.5×10^{16}	4.5×10^{-7}	Turner (1991) (Onsala)
			192^{+18}_{-16}	$(1.6 \pm 0.2) \times 10^{16}$	1.6×10^{-7}	Turner (1991) (NRAO)
			> 200	$2.5 - 30 \times 10^{16}$	1.6×10^{-6}	Sutton et al. (1995)
			120–200	$4 - 8 \times 10^{16}$	6×10^{-7}	this work
	C_2H_5OH		216^{+3600}_{-105}	$6.6^{+4.1}_{-2.5} \times 10^{15}$	6.6×10^{-8}	Turner (1991)
			289 ± 330	$4.8^{+35}_{-4.2} \times 10^{14}$	4.8×10^{-9}	this work ^c
	CH_3OCH_3		75	2.5×10^{15}	2.5×10^{-8}	Johansson et al. (1984)
			63	3.0×10^{15}	3.0×10^{-8}	Blake et al. (1986)
			91^{+34}_{-19}	$1.3^{+0.3}_{-0.1} \times 10^{15}$	1.3×10^{-8}	Turner (1991)
			125	$\sim 3 \times 10^{15}$	3.0×10^{-8}	Sutton et al. (1995)
			197.4^{+165}_{-62}	$1.1^{+2.5}_{-0.6} \times 10^{16}$	1.1×10^{-7}	Lee & Cho (2002)
	$C_2H_5OCH_3$		126 ± 43	$4.6^{+3.1}_{-1.8} \times 10^{15}$	4.6×10^{-8}	this work
			100–150	$< 7 \times 10^{13}$	$< 7 \times 10^{-10}$	this work
SgrB2(N)	CH_3OH^d	I	49 ± 1	$(2.8 \pm 0.07) \times 10^{16}$	5.6×10^{-9}	Cummins et al. (1986)
			26^{+138}_{-12}	$8.1^{+8}_{-4} \times 10^{15}$	1.6×10^{-9}	Turner (1991)
		II	120 ± 20	$(3.5 \pm 0.5) \times 10^{15}$	7×10^{-10}	Cummins et al. (1986)
			204^{+392}_{-81}	$6.9^{+5.6}_{-3.1} \times 10^{15}$	1.4×10^{-9}	Turner (1991)
	C_2H_5OH		14	5.3×10^{14}	1.1×10^{-10}	Cummins et al. (1986)
			36	$4.8^{+4.5}_{-2.4} \times 10^{15}$	9.6×10^{-10}	Turner (1991)
	CH_3OCH_3		24	5×10^{14}	1×10^{-10}	Cummins et al. (1986)
			80^{+351}_{-37}	$2.3^{+0.9}_{-0.4} \times 10^{15}$	4.6×10^{-10}	Turner (1991) ^e
W51e2	CH_3OH	a -type	208	1.0×10^{17}	2.7×10^{-7}	Ikeda et al. (2002)
			101 ± 4	$2.5^{+0.5}_{-0.3} \times 10^{16}$	6.9×10^{-8}	this work
	C_2H_5OH		169 ± 28	$(3.1 \pm 1.1) \times 10^{15}$	8.6×10^{-9}	Ikeda et al. (2002)
			143 ± 20	$1.6^{+0.4}_{-0.3} \times 10^{15}$	4.4×10^{-9}	this work
	CH_3OCH_3		145 ± 8	$1.6^{+0.4}_{-0.2} \times 10^{16}$	4.4×10^{-8}	this work
	$C_2H_5OCH_3$		69 ± 94	$2^{+29}_{-1.9} \times 10^{14}$	5.6×10^{-10}	this work

^a We used $N(H_2)(G34.26) = 4 \times 10^{23} \text{ cm}^{-2}$, $N(H_2)(NGC 6334I) = 2 \times 10^{23} \text{ cm}^{-2}$, $N(H_2)(Orion KL) = 1 \times 10^{23} \text{ cm}^{-2}$, $N(H_2)(SgrB2(N)) = 5 \times 10^{24} \text{ cm}^{-2}$ and $N(H_2)(W51e2) = 3.6 \times 10^{23} \text{ cm}^{-2}$, see Sect. 4.

^b Observed at center of G34.26, as opposed to the halo.

^c The Onsala and CIT survey (see Turner 1991) did not find any ethanol towards Orion KL and estimated an upper limit between 1.5 and $4 \times 10^{14} \text{ cm}^{-2}$.

^d Cummins et al. (1986) and Turner (1991) used two rotational temperature diagrams (I and II) to fit the observed CH_3OH lines. Cummins et al. (1986) uses the transitions with E_u below 70 K for the fit I whereas Turner (1991) splits the two components I and II at $E_u = 40$ K.

^e Turner (1991) gives two fits of T_{rot} and N_t , the values used here correspond to the most reliable fit.

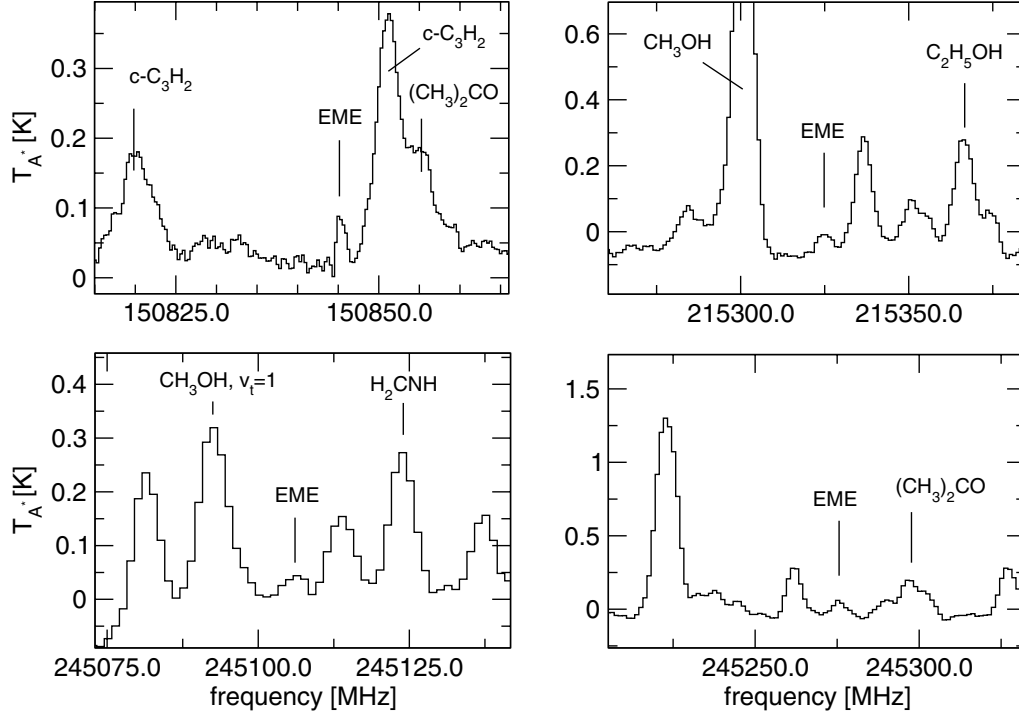


Fig. 3. Spectra recorded towards W51e2. The EME transitions $20_{0,20}-19_{1,19}$ at 150 845.5 MHz, $19_{5,14}-19_{4,15}$ at 215 325.4 MHz, $16_{3,14}-15_{2,13}$ at 245 106.4 MHz, and $17_{3,15}-16_{2,14}$ at 245 190.7 MHz are indicated.

is characterized by a rotational thermal equilibrium temperature. For each source a simulated spectrum has been created which covered all measured frequency bands. This enables an estimate of the abundances and temperatures of the molecules in these sources and can help to reveal blendings of several lines at a given frequency position. myXCLASS is specially suited to deal with many convoluting line profiles with different line widths and intensities as can regularly be observed in hot core regions where lines of different molecules blend with each other.

The spatial extent of the molecule can influence the value of the inferred column density. This is because the circular Gaussian beam size diameter θ_B of the telescope varies with frequency ν , i.e. $\theta_B \sim 1/\nu$ (see telescope beam *HPBW* in the online table). If the telescope beam points towards a source of circular Gaussian type with size θ_S the resulting convolution beam filling factor b is given by $b = \theta_S^2 / (\theta_B^2 + \theta_S^2)$, see Snyder et al. (2005) and the work by Ulich & Haas (1976). In the case of $\theta_S \gg \theta_B$ the beam filling factor is close to unity, for $\theta_S \sim \theta_B$ the beam dilution is 0.5 and for $\theta_S \ll \theta_B$ it follows $b \approx \theta_S^2 / \theta_B^2$. Thus, assuming that the intrinsic intensity distribution T_S of a source is Gaussian, a beam filling factor can be introduced to calculate the measured intensity $T_{mb} = b \cdot T_S$. HMCs are compact objects with diameters $< 10''$. Using the IRAM 30 m telescope with *HPBW* around $30''$ and $10''$ at 80 and 250 GHz respectively and assuming a $10''$ source size the beam filling factor is $b(@80 \text{ GHz}) = 0.1$ and $b(@250 \text{ GHz}) = 0.5$. Hence, an intensity difference of a factor 5 can occur by not correcting for beam dilution. The program myXCLASS has the source size as an input parameter and can thus simulate the effect of the assumed source size on the line intensities.

For the EME lines we can expect small optical depths so that the rotational temperature T_{rot} and the total molecular column density $N_T [\text{cm}^{-2}]$ can be determined by a least-squares fit of

$$\log_{10} \left(\frac{3k_B}{8\pi^3} \frac{\int T_{mb} dv}{\nu \mu^2 S} \right) = \log_{10} \left(\frac{N_T}{Q(T_{rot})} \right) - \frac{\log_{10}(e) E_u}{T_{rot}} \quad (1)$$

with $\int T_{mb} dv$ the observed line integral, $\mu^2 S$ the dipole and line strength value derived from Pickett et al. (1998) (see their Eq. (2)), Q the partition function, and E_u the energy of the upper level. The resulting fit can be plotted in a rotational-temperature-diagram (RTD) or Boltzmann plot. For detections at the confusion limit Snyder et al. (2005) has shown that Boltzmann plots cannot necessarily be used as independent tools to verify interstellar spectral line identifications. Any false assignment does not automatically reveal itself in the RTD by large displacements from the fitted line and thus can only be avoided by careful line check in catalogs and the literature. As a standard procedure any candidate line of EME has to be checked against competing lines, i.e. the expected or known intensities of these other lines at the EME frequencies have to be determined. Therefore, a consistent and comprehensive list of molecule abundances in the source is important. Line surveys are invaluable aids in determining the chemical composition of a source.

5. Results

5.1. W51e2

Several lines at EME transitions have been observed towards W51e2, see Fig. 3. We looked for strong EME b -type lines

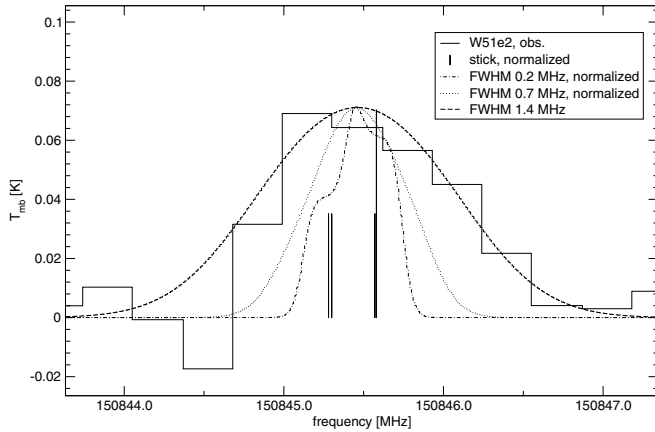


Fig. 4. The $20_{0,20}-19_{1,19}$ transition of EME at 150 845.5 MHz. The line profile observed towards W51e2 (bold) is compared to three simulated line profiles (dashed) of 0.2, 0.7, and 1.4 MHz line widths ($FWHM$). For a line width of 0.2 MHz ($FWHM$) the splitting due to internal torsional motion of the molecule is partly resolved. The stick diagram indicates the position and relative intensities of the split lines based on our recent high resolution laboratory measurements Fuchs et al. (2003). For comparison with the observed spectrum all simulated spectra are normalized to have the same peak intensity.

and at the same time tried to exclude as many blendings with other molecules. By using myXCLASS we first estimated a column density of $\sim 10^{14} \text{ cm}^{-2}$ and a rotational temperature of roughly 100 K. We checked every frequency band for consistency by comparing a simulated spectrum of EME and other known molecules with the observed spectrum. Figure 4 shows an observed line with simulated EME lines superimposed. The laboratory measurements of the EME line positions are at a precision level unmatched by any astronomical observations. The figure illustrates the effect of the linewidth on the total observable line shape and shows that the unique EME feature which is due to the torsional splitting remains unresolved in the observed spectrum. The observed lines⁵ at transition frequencies of EME are shown in Table 3. The more stronger lines can be found above 150 GHz whereas many lines at the 1σ – 2σ level have been observed at lower frequencies. Hot cores reveal a great richness and density in molecular emission lines, see Fig. 8, and it is not astonishing that 77% of the EME lines in the observed bands are blended by strong lines of other known molecules. In our case by far the most blendings occur in the 215 GHz band which is therefore not further detailed in Table 3. 12% of the blendings are at line positions of yet unidentified species. However, from intensity considerations it is clear that these latter lines cannot be dominated by EME because in this case EME should have also strong lines at other frequencies where we have not observed any strong feature. Note that many transition in Table 3 have labels with possible blendings. By carefully checking the databases and literature but also by fitting many other species to our spectra we could estimate the abundances of several species which could be potential competitors to the EME lines. To further clarify our detection we made two Boltzmann fits, as shown in Fig. 5. Here,

⁵ The $5_{3,2}-4_{2,3}$ transition line at 160.1 GHz from Charnley et al. (2001) is not listed in Table 3 and has not been considered in our fits.

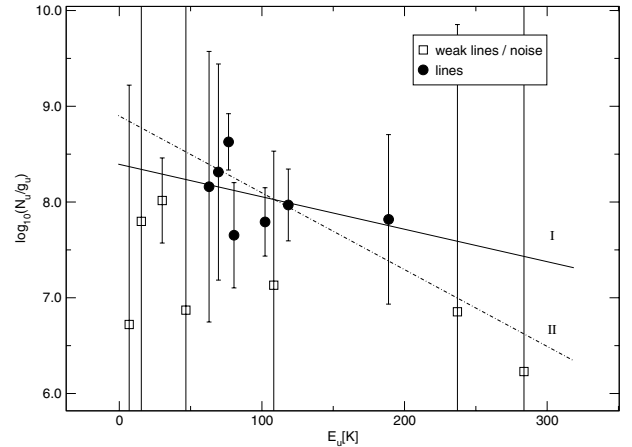


Fig. 5. Boltzmann plot for the EME transitions found towards W51e2 based on 14 observed lines, assuming no beam dilution. Due to difficulties in assigning the lines to EME two fits have been performed. **I)** Fit including all lines marked with $\square$ and $\bullet$ in Table 3. Here $T_{\text{rot}} = 126 \pm 115 \text{ K}$ and $\log(N_l) = 14.9 \pm 0.7$. **II)** Fit using only lines with intensities $\geq 2\sigma$ ($\bullet$) resulting in $T_{\text{rot}} = 54 \pm 38 \text{ K}$ and $\log(N_l) = 14.3 \pm 1.2$. Lines which are less than 2σ intense are marked with an $\square$.

every line in Table 3 which is marked with an ($\bullet$) or ($\square$) is depicted. If we use all these 14 lines in the Boltzmann fit (I) we get $T_{\text{rot}} = 126 \pm 115 \text{ K}$ and $N_l = 7.9^{+32}_{-6} \times 10^{14} \text{ cm}^{-2}$ as the beam averaged column density. Since we have many weak lines (i.e. around 1σ level) the reliability of the fit can be checked by excluding the weaker or doubtful lines marked with ($\square$ = noisy-like). Fit (II) only includes lines ($\bullet$) with intensities larger than 2σ . As a result we obtain $T_{\text{rot}} = 54 \pm 38 \text{ K}$ and $N_l = 6.3^{+25}_{-5} \times 10^{14} \text{ cm}^{-2}$. This fit II should be regarded as more reliable than fit I. Furthermore, it has been shown that the excluded weak and noisy lines ($\square$) are not in contradiction to fit II. Due to the large errors of the weak lines in the Boltzmann plot this is indeed not the case, but we would still expect slightly higher intensities at the ($\square$) frequency positions.

W51e2 is known to have a size $< 10''$ so that the effect of beam dilution should be considered. For each fit (I and II) the above mentioned temperatures and abundances of EME can be recalculated by assuming an $5''$ source size. This results in source averaged column densities and temperatures: fit I gives $T_{\text{rot}} = 74 \pm 50 \text{ K}$ and $N_l = 2^{+8}_{-1.5} \times 10^{14} \text{ cm}^{-2}$, fit II $T_{\text{rot}} = 69 \pm 94 \text{ K}$ and $N_l = 2^{+29}_{-1.9} \times 10^{14} \text{ cm}^{-2}$. Hence, the column densities and temperatures of the diluted and non-diluted beam yield comparable results within their error limits and we will use the source averaged value of fit (II) for further investigations. As has already been stated, RTDs cannot necessarily be used as reliable independent evidence to support the detection of a molecule close to the confusion limit, as was shown by Snyder et al. (2005) in the case of glycine. Our estimated column density is also consistent with observations in less favored frequency regions as can be seen in Fig. 6 where an EME line is expected. For our fitted $2 \times 10^{14} \text{ cm}^{-2}$ column density no contradiction within the 1σ level can be found, but for a $N_l = 4\text{--}8 \times 10^{14} \text{ cm}^{-2}$ the simulated spectrum is in clear conflict with observation.

Table 3. EME transitions observed in W51e2.

Transition	Observed frequency ^a (MHz)	Theor. frequency ^c (MHz)	Obs-calc (MHz)	line components	E_{upper} (K)	$\int T_{\text{mb}} dv$ (K km s ⁻¹)		
11 _{2,10} –11 _{1,11}	80 882.3(5) ^b	80 882.7(10) ^d	–0.4	5	30.1	0.09(4)	□	n
24 _{1,23} –24 _{0,24}	81 198.4(1)	81 198.7(5)	–0.3	5	118.5	0.16(6)	•	c ₁
35 _{3,32} –35 _{2,33}	91 439.6(5)	91 440.4(10)	–0.8	5	255.2	1.0(44)		b
36 _{3,33} –36 _{2,34}	–	91 475.6(9)		5	269.3			b
34 _{2,32} –34 _{1,33}	91 630.1 ^e	91 630.8(5)	–0.7	5	237.1	0.03(9)	□	c ₂ , n
34 _{3,31} –34 _{2,32}	–	91 688.3(10)	–	5	241.5	–		n
37 _{3,34} –37 _{2,35}	91 812.7(4)	91 812.6(9)	0.1	5	283.7	0.1(5)	□	n
35 _{2,33} –35 _{1,34}	–	96 207.0(9)	–	5	250.8	–		b
29 _{3,26} –29 _{2,27}	96 390.8(3)	96 391.4(12)	–0.6	5	179.1	1.85(37)		b
3 _{2,1} –2 _{1,2}	96 464.1 ^e	96 464.3(6)	–0.2	5	6.9	0.002(5)	□	n
22 _{3,19} –22 _{2,20}	107 654.8 ^e	107 656.8(14)	–2.0	5	108.3	0.05(7)	□	n
7 _{2,5} –6 _{1,6}	131 350.2(10)	131 350.7(8)	–0.5	5	15.4	0.02(6)	□	c ₃ , n
15 _{1,15} –14 _{0,14}	131 372.68(80)	131 372.8(5)	–0.12	5	46.7	0.02(10)	□	c ₄ , n
36 _{1,35} –36 _{0,36}	–	150 502.4(19)		5	260.0 6	–		b
34 _{4,30} –34 _{3,31}	150 661.3(6)	150 662.9(16)	–1.6	5	248.7	0.12(2)		b
13 _{6,x} –14 _{5,y}	150 794.1 ^e	150 794.1(18)	0.0	10	76.6	0.17(0.05)	•	c ₅
20 _{0,20} –19 _{1,19}	150 845.52(30)	150 845.5(2)	0.02	5	80.4	0.20(11)	•	c ₆
[...] ^f								b
19 _{5,x} –19 _{4,15}	215 325.4 ^e	215 326.3(13)	–0.9	6	102.2	0.28(10)	•	b
[...] ^f								b
28 _{0,28} –27 _{1,27}	217 939.6(3)	217 940.7(1)	–1.1	5	154.7	3.66(19)		b
16 _{3,14} –15 _{2,13}	245 106.4(32)	245 104.7(29)	1.43	5	62.9	0.46(65)	•	c ₇
31 _{1,31} –30 _{0,30}	245 275.0(13)	245 274.1(1)	0.86	5	188.8	0.87(77)	•	
17 _{3,15} –16 _{2,14}	252 190.7(11)	252 189.4(29)	1.3	5	69.5	0.70(79)	•	c ₈
28 _{2,27} –27 _{1,26}	253 309.3(3)	253 308.2(12)	1.09	5	161.0	16.01(126)		b

^a Rest frequencies (assuming a source LSR velocity of +55 km s⁻¹) in W51e2.^b The numbers in parenthesis are the estimated uncertainties (1 σ).^c Frequencies are taken from Fuchs et al. (2003).^d The deviation from the mean frequency of the transitions to its extreme values, e.g. due to rotational and internal rotational splitting the transition 11_{2,10}–11_{1,11} is split in 5 transitions which occur between 80 881.7–80 883.6 GHz, thus the frequency denoted in the table is 80 882.7(10).^e Manually determined (or in case of noise the closest peak). Here the transition 13_{6,x}–15_{5,y} are with $x-y = 7-9, 7-10, 8-9, 8-10$ and the x in 19_{5,x}–19_{4,15} denotes either 14 or 15.^f Region with many blends of EME with other (dominating) molecules.

• Lines included in the Boltzmann fit II, see Fig. 5.

□ Lines included in the Boltzmann fit I and II, see Fig. 5.

n “Noisy” line, i.e. a line which has either less than 2 σ intensity or where the intensity is difficult to obtain due to baseline variations.

b Denotes a blend with an other molecular line where the other line contributes most to the total intensity.

c_i Denotes a contribution of another molecule to the total intensity where the blended line is estimated to contribute less than 20% (if not otherwise mentioned) compared to EME. Assuming $T_{\text{rot}} \sim 100$ K:c₁ C₂H₅OOCH with $N \leq 5 \times 10^{15}$ cm⁻² (could be up to 50%), c₂ HOONO₂ with $N \leq 5 \times 10^{14}$ cm⁻², c₃ c-H₂C₃O with $N \leq 2 \times 10^{12}$ cm⁻², c₄ C₂H₃NH₂ with $N \leq 8 \times 10^{15}$ cm⁻², c₅ C₃H₇CN with $N \leq 2 \times 10^{14}$ cm⁻², c₆ C₂H₅C¹⁵N with $N \leq 3 \times 10^{13}$ cm⁻² based on our observation that C₂H₅CN has $N \sim 3 \times 10^{14}$ cm⁻², c₇ CH₂(OH)CHO with $N \leq 8 \times 10^{14}$ cm⁻², c₈ a-H₂C = CHOH with $N \leq 2 \times 10^{14}$ cm⁻².

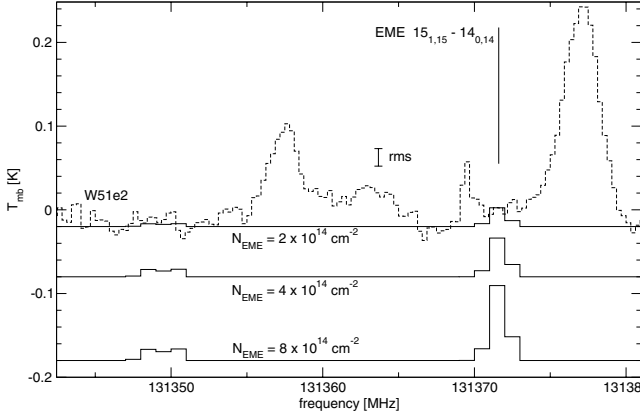


Fig. 6. Observed spectrum at W51e2 (top, dashed) and calculated spectra for different column densities of EME at 70 K.

It is commonly believed that most complex molecules exist within a small area around the forming star, e.g. around 5'' for W51e2. On the other hand Hollis et al. (2001) have shown that complex molecules such as glycolaldehyde can be found on spatial scales of $\geq 60''$ in the case of SgrB2. With our data it is not possible to determine the size of the emitting region but the source size has surely an influence on the detected signal strength which we tried to estimate using the myXCLASS program. Table 4 lists the result of possible column densities consistent with the observations if we assume certain temperatures between 30 K and 150 K. Charnley et al. (2001) presented a map of a line at an EME transition frequency towards SgrB2 of the order of 20''. If we assume W51e2 to have this same source size the column density would have to be corrected to lower values by nearly an order of magnitude. More realistic, our best fit value (fit II) assumes an 5'' size of W51e2.

Furthermore, it can be seen from Fig. 7 (see simulated spectrum for $N = 1 \times 10^{15} \text{ cm}^{-2}$) that in some frequency regions strong EME lines, i.e. lines well above 3σ level, should reveal their internal line splitting. Due to the weakness of the observed lines no line was detected where the torsional splitting could be resolved so that there is no unique proof that these lines indeed belong to EME.

5.2. The other sources

Sample spectra showing all examined sources at 245 GHz are shown in Fig. 8. At this frequency one of the strongest observed EME candidate lines appears in W51e2. However, no line above 1σ appears in these spectra for any other examined source. Also for the other frequency bands we get similar results. For example, the well pronounced feature at 150 845 MHz in W51e2 is only at the $1-2\sigma$ level in Orion KL and is completely invisible in G34.26 and NGC 6334I. An upper limit for the column density of G34.26, NGC 6334I and Orion KL has been estimated by the use of myXCLASS and independently by RTDs using integrated noise at EME transition frequencies and assuming an appropriate line width. Figure 7 compares the 215 GHz region of W51e2 with Orion KL and also shows simulated spectra of EME for $N = 2 \times 10^{14} \text{ cm}^{-2}$ and $N = 1 \times 10^{15} \text{ cm}^{-2}$ at 70 K (assuming an 5'' source size). It

Table 4. Estimated column density for various source sizes and rotational temperatures for W51e2.

source size [arcsec]	T_{rot} [K]	column density [cm^{-2}]
20''	30	$\sim 5 \times 10^{13}$
	50	$\sim 4 \times 10^{13}$
	70	$\sim 4 \times 10^{13}$
10''	50	$\sim 8 \times 10^{13}$
	70	$\sim 9 \times 10^{13}$
	150	1.3×10^{14}
5''	70	2.0×10^{14}
	150	4.5×10^{14}

is immediately clear that the column densities of these sources must be of the order of 10^{14} cm^{-2} or less rather than 10^{15} cm^{-2} as estimated by Charnley et al. (2001).

Our results are summarized in Table 2. The previously published line at the EME $5_{3,2}-4_{2,3}$ transition frequency at 160.1 GHz in Orion KL by Charnley et al. (2001) has not been considered in the fit.

6. Discussion

Charnley et al. (2001) estimate the column density of EME in Orion KL and SgrB2(N) to be between $10^{14}-10^{15} \text{ cm}^{-2}$ and in W51 to be a few times 10^{14} cm^{-2} . EME column densities at around 10^{14} cm^{-2} correspond to signal strengths which are already close to the confusion limit in the HMC spectra. Nevertheless, long integration times and the availability of spectra at many frequencies enable us to estimate the column density of Orion KL to be below $1 \times 10^{14} \text{ cm}^{-2}$ and those of G34.26 and NGC 6334I to be of the same order.

We can now use our values in Table 2 of W51e2 to draw some conclusions using the model of Charnley et al. (1995). According to Figs. 2 and 3a in their paper, which assumes a 10^{-6} fractional abundance X_{Eol} of ethanol as a starting value for the chemical evolution in their model, the age of W51e2 should be either $\geq 10^5$ yr if we use CH_3OH or $\text{C}_2\text{H}_5\text{OH}$ as a probe molecule or 10^4-10^5 if we look at CH_3OCH_3 . However, that would imply a fractional abundance between $\leq 10^{-10}$ and $\sim 3 \times 10^{-9}$ for EME. On the other hand, if we use Fig. 4a of Charnley et al. (1995) with ethanol starting at $X_{\text{Eol}} = 10^{-7}$ this translates to $X_{\text{EME}} \leq 10^{-12}$ and $X_{\text{EME}} \sim 4 \times 10^{-10}$. Assuming $\text{C}_2\text{H}_5\text{OH}$ to be the best molecule to determine the chemical age of the HMC it appears that in both cases, i.e. with ethanol starting values or $X_{\text{Eol}} = 10^{-6}$ and $X_{\text{Eol}} = 10^{-7}$, the fractional abundance of EME is nearly the same, namely $X_{\text{EME}} \sim 2 \times 10^{-10}$. This value is close to our estimated abundance 5.6×10^{-10} for EME in W51e2. According to Table 2 and the model by Charnley et al. (1995) Orion KL and NGC 6334F should be roughly of the same age as W51e2. Although we have only upper limits on the EME abundance in these sources it is clear that as a consequence the $\text{C}_2\text{H}_5\text{OH}$ fractional abundance at 0 yr must be 10^{-7} rather than 10^{-6} . G34.26 is slightly older and SgrB2 is the oldest of the here discussed sources and should have an EME abundance X_{EME} of less than 10^{-10} (corresponding to less than 10^{14} cm^{-2} column density as we see it for Orion KL, G34.26 and NGC 6334I). This however is in

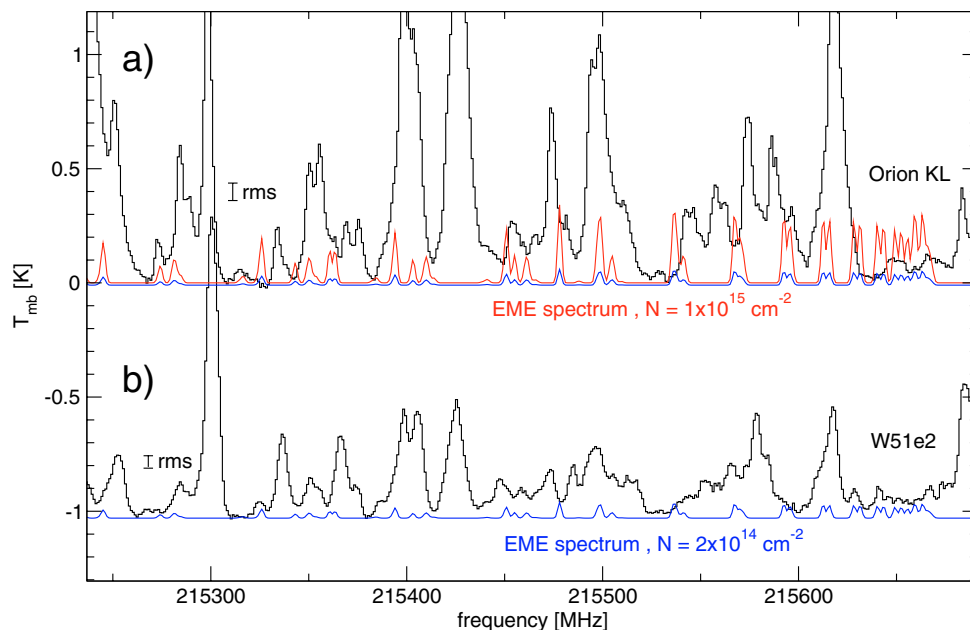


Fig. 7. Observed spectra of Orion KL and W51e2 at 215 GHz. This frequency region has the highest line density of all our observed bands but even here gaps can be found where the presence of EME can be checked. Two simulated spectra of EME are plotted, i.e. $N = 2 \times 10^{14} \text{ cm}^{-2}$ (blue) and $N = 1 \times 10^{15} \text{ cm}^{-2}$ (red) with $T_{\text{rot}} = 70 \text{ K}$. The splitting of the lines which can best be seen above 215 600 MHz is mainly due to the internal rotation of the methyl groups of EME with an additional small asymmetry contribution. a) For Orion KL the previously estimated limits of 10^{14} – 10^{15} cm^{-2} by Charnley et al. (2001) seem questionable as can be seen from the superimposed simulated spectra on the observed spectra. b) In the case of W51e2 the simulated spectrum is in agreement with the observation.

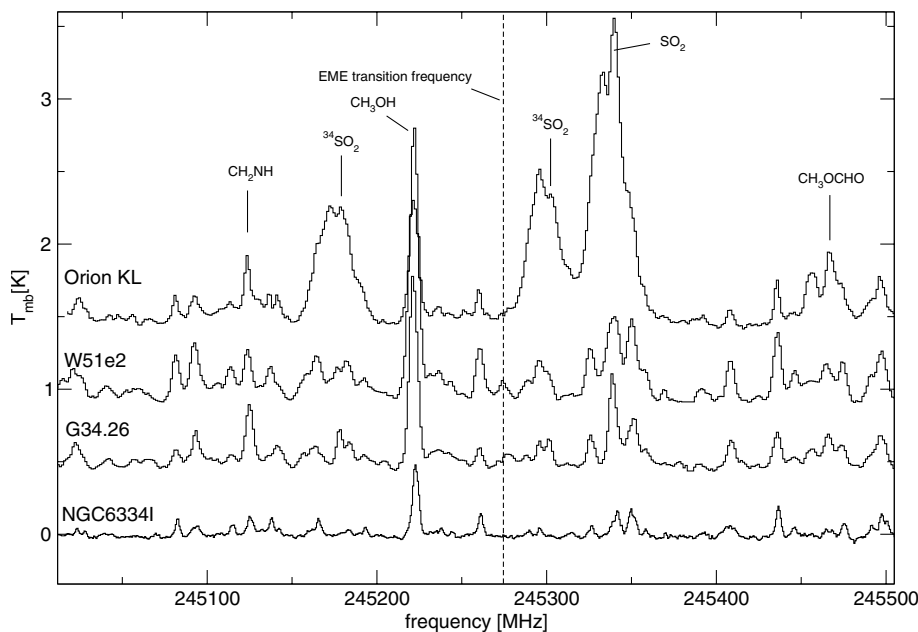


Fig. 8. Spectra of the observed sources at 245 GHz. Only W51e2 reveals a line at an EME transition frequency, whereas all other sources do not show any significant feature above 1σ level. However, W51e2, G34.26 and NGC 6334I have similar spectra indicating the same chemical composition and equivalent dynamics.

contradiction to the claimed observations of EME towards SgrB2 by Charnley et al. (2001) when they estimate an abundance of 10^{-10} – 10^{-9} .

Looking at the observed values only, we notice that Orion KL has the highest fractional abundance of ethanol relative to H_2 followed by NGC 6334I, G34.26, W51e2 and then SgrB2(N). However, no EME has been detected in Orion KL.

The relative abundances of DME with respect to methanol and ethanol are shown in Fig. 9. Here, W51e2 shows a high dimethyl ether content relative to methanol and ethanol compared to the other sources, which may be a general indication of a higher ether abundance relative to the alcohols. Assuming that we indeed have detected EME in W51e2 the ratio between EME and ethanol is 0.13 which is 5 times less than

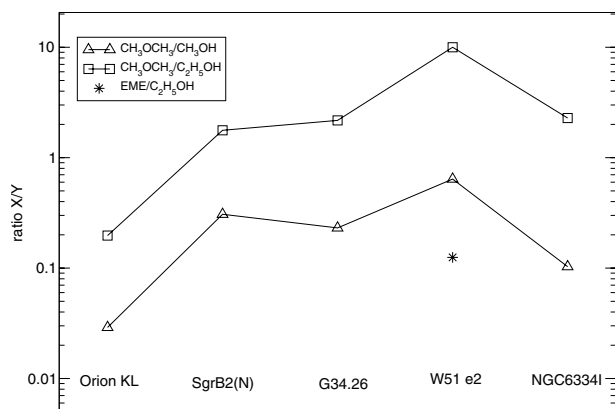


Fig. 9. Relative abundances of CH_3OH , $\text{C}_2\text{H}_5\text{OH}$, CH_3OCH_3 derived from Table 2 by using the values from Turner (1991) for Orion KL, Turner (1991) for SgrB2(N), Thompson et al. (1999) and Ikeda et al. (2002) for G34.26, this work for W51e2 and Nummelin et al. (1998b) for NGC 6334I. The ratio of EME/ $\text{C}_2\text{H}_5\text{OH}$ in W51e2 is 0.13 which is 5 times smaller than the ratio of $\text{CH}_3\text{OCH}_3/\text{CH}_3\text{OH}$.

the ratio of DME and methanol. If all sources would have a 0.13 EME/ethanol ratio the theoretical column densities would be around $2 \times 10^{14} \text{ cm}^{-2}$ or higher⁶ which at least for G34.26, NGC 6334I and Orion KL has been shown to be not the case. Thus, W51e2 is indeed particularly rich in EME.

We conclude that only the tentative detection of EME by Charnley et al. (2001) towards W51e2 can be confirmed within the following caveats 1) the EME Boltzmann fits include a few lines close to the noise level, 2) the obtained temperature of 70 K for EME is lower than that of other observed molecules ($\gtrsim 140$ K), 3) no torsional splitting of the EME lines could be observed and also other characteristic spectral fingerprints have not been observed. However, there are lines at EME transition frequencies which until now can best be explained by the existence of EME in W51e2.

Acknowledgements. We are grateful to the IRAM 30 m and SEST 15 m staff for their hospitality and assistance. Many thanks to Peter Schilke (MPIfR) for providing us with the myXCLASS program code. Many thanks to Eric Herbst for his invaluable suggestions and discussions. This work was supported by the *Deutsche Forschungsgemeinschaft*, DFG within SFB 494, IRAM, and ESO.

References

- Blake, G., Sutton, E., Masson, C., & Phillips, T. 1987, *ApJ*, 315, 621
 Blake, G. A., Sutton, E., Masson, C., & Phillips, T. 1986, *ApJS*, 60, 357
 Bohme, D. K., Mackay, G. I., & Tanner, S. 1979, *Am. Chem. Soc.*, 101, 3724
 Carpenter, J. H., & Sanders, D. 1998, *AJ*, 116, 1856
 Caselli, P., Hasegawa, T., & Herbst, E. 1993, *ApJ*, 408, 548
 Charnley, S. B., Kress, M., Tielens, A., & Millar, T. 1995, *ApJ*, 448, 232
 Charnley, S. B., Ehrenfreund, P., & Kuan, Y.-J. 2001, *Spectrochim. Acta. Part A*, 57, 685
 Cheung, L., Frogel, J., Gezari, D., & Hauser, M. 1978, *ApJ*, 226, L149
 Cummins, S. E., Linke, R., & Thaddeus, P. 1986, *ApJS*, 60, 819

- Durig, J., Jin, Y., Phan, H., Liu, J., & Durig, D. 2002, *Struct. Chem.*, 13, 1
 Friedel, D. N., Snyder, L. E., Turner, B. E., & Remijan, A. 2004, *ApJ*, 600, 234
 Fuchs, U. 2003, *High Resolution Spectroscopy of Ethyl Methyl Ether and Hydrogen Cyanide Isotopomers* (Göttingen: Cuvillier Verlag)
 Fuchs, U., Winnewisser, G., Groner, P., De Lucia, F. C., & Herbst, E. 2003, *ApJS*, 144, 277
 Hayashi, M., & Kuwada, K. 1975, *J. Molec. Struct.*, 28, 147
 Heaton, B. D., Little, L. T., & Bishop, I. S. 1989, *A&A*, 213, 148
 Herbst, E. 2001, *Chem. Soc. Rev.*, 30, 168
 Hollis, J. M., Vogel, S. N., Snyder, L. E., Jewell, P. R., & Lovas, F. J. 2001, *ApJ*, 554, L81
 Hüttemeister, S., Wilson, T., Henkel, C., & Mauersberger, R. 1993, *A&A*, 276, 445
 Ikeda, M., Ohishi, M., Nummelin, A., et al. 2001, *ApJ*, 560, 792
 Ikeda, M., Ohishi, M., Nummelin, A., et al. 2002, *ApJ*, 571, 560
 Johansson, L., Andersson, C., Ellmér, J., et al. 1984, *A&A*, 130, 227
 Karpas, Z., & Meot-Ner (Mautner), M. 1989, *J. Phys. Chem.*, 93, 1859
 Kuan, Y.-J., Charnley, S. B., Huang, H.-C., et al. 2004, *Adv. Space Res.*, 33, 31
 Lee, C. W., & Cho, S.-H. 2002, *J. Korean. Astron. Soc.*, 35, 187
 Liu, S.-Y., Mehringer, D. M., & Snyder, L. E. 2001, *ApJ*, 552, 654
 MacDonald, G. H., Gibb, A. G., Habing, R. J., & Millar, T. J. 1996, *A&AS*, 119, 333
 Mackay, G., Tanner, S., Hopkinson, A., & Bohme, D. 1979, *Can. J. Chem.*, 57, 1518
 Martín-Pintado, J., de Vicente, P., Wilson, T. L., & Johnston, K. J. 1990, *A&A*, 236, 193
 McCutcheon, W., Sandell, G., Matthews, H., et al. 2000, *MNRAS*, 316, 152
 Millar, T. & Hatchell, J. 1998, *Chemistry and Physics of Molecules and Grains in Space*, Faraday Discussions, 109, 15
 Millar, T., Macdonald, G., & Habing, R. 1995, *MNRAS*, 273, 25
 Nomura, H., & Millar, T. 2004, *A&A*, 414, 409
 Nummelin, A., Bergman, P., Friberg, P., et al. 1998a, *ApJS*, 117, 427
 Nummelin, A., Dickens, J., Bergman, P., et al. 1998b, *A&A*, 337, 275
 Pickett, H., Poynter, R., Cohen, E., et al. 1998, *J. Quant. Spectrosc. Radiat. Transfer*, 60, 883
 Remijan, A., Snyder, L., Liu, S.-Y., Mehringer, D., & Kuan, Y.-J. 2002, *ApJ*, 576, 264
 Remijan, A., Sutton, E., Snyder, L., et al. 2004, *ApJ*, 606, 917
 Sandell, G. 2000, *A&A*, 358, 242
 Schilke, P., Groesbeck, T., Blake, G., & Phillips, T. 1997, *ApJS*, 108, 301
 Schloerb, F., Snell, R., & Schwartz, P. 1987, *ApJ*, 319, 426
 Snyder, L., Lovas, F., Hollis, J., et al. 2005, *ApJ*, 619, 914
 Sutton, E., Blake, G. A., Masson, C., & Phillips, T. 1985, *ApJS*, 58, 341
 Sutton, E., Jaminet, P., Danchi, W., & Blake, G. A. 1991, *ApJS*, 77, 255
 Sutton, E., Peng, R., Danchi, W., et al. 1995, *ApJS*, 97, 455
 Thompson, M., G. H. Macdonald, G., & Millar, T. 1999, *A&A*, 342, 809
 Turner, B. 1989, *ApJS*, 70, 539
 Turner, B. E. 1991, *ApJS*, 76, 617
 Ulich, B., & Haas, R. 1976, *ApJS*, 30, 247
 Zhang, Q., Ho, P. T. P., & Ohashi, N. 1998, *ApJ*, 494, 636

⁶ Except for the SgrB2(N) value by Cummins et al. (1986).

Online Material

Table 5. Observational parameters.

Receiver frequency (MHz)	T_{sys}^a (K)	τ	Tot. integr. time (on + off) (min)	ρ^b	Telescope beam <i>HPBW</i> ($''$)	1σ (rms) noise level (K as T_A^*)	Band width (MHz)	Frequency resolution (MHz)
G34.26								
81 041	144	0.03	85	0.79	30	0.012	490	0.3125
91 626	122	0.03	115	0.77	26	0.010	490	0.3125
96 371	137	0.02	145	0.77	25	0.014	490	0.3125
131 372	200	0.06	115	0.71	18	0.018	490	0.3125
150 750	307	0.04	220	0.68	16	0.017	490	0.3125
215 500	388	0.22	196	0.56	11	0.032	1020	1.0
217 900	350	0.14	147	0.55	11	0.023	1020	1.0
245 260	511	1.90	160	0.50	10	0.034	510	1.25
252 227	848	0.37	85	0.48	10	0.059	510	1.25
NGC 6334 I								
81 198	327	0.41	38	0.76	59	0.009	994	0.6911
91 450	195	0.26	37	0.74	55	0.007	994	0.6911
91 630.7	185	–	87	0.74	55	0.005	994	0.6911
91 811.7	212	0.06	57	0.74	55	0.007	994	0.6911
96 380	252	0.21	55	0.73	53	0.009	994	0.6914
150 662	263	0.34	95	0.65	33	0.012	994	0.6989
245 105	513	0.96	77	0.47	22	0.024	994	0.6989
245 274	534	0.50	127	0.47	22	0.017	994	0.6989
Orion KL								
81 041	138	0.03	125	0.79	30	0.012	490	0.3125
91 626	182	0.07	95	0.77	26	0.014	490	0.3125
150 750	316	0.14	220	0.68	16	0.029	490	0.3125
215 500	539	0.33	160	0.56	11	0.078	1020	1.0
245 275	740	0.39	190	0.50	10	0.053	510	1.25
W51e2								
81 041	113	0.04	95	0.79	30	0.012	490	0.3125
91 626	116	–	60	0.77	26	0.012	490	0.3125
96 371	119	0.07	95	0.77	25	0.013	490	0.3125
107 650	223	0.14	52	0.75	22	0.018	490	0.3125
131 372	179	–	60	0.71	18	0.019	490	0.3125
150 750	249	0.26	215	0.68	16	0.020	490	0.3125
215 500	221	0.415	195	0.56	11	0.063	1020	1.0
217 900	301	0.634	112	0.55	11	0.044	1020	1.0
245 260	371	0.75	75	0.50	10	0.051	510	1.25
252 227	384	0.51	95	0.48	10	0.062	510	1.25
253 300	330	–	85	0.48	10	0.045	510	1.25

^a Average system temperature.^b $T_A^* = \rho T_{\text{mb}}$.