

## Millimeter observations of organic molecules toward high-mass star formation region G34.26+0.15

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**Abstract** To investigate the chemical origination of organic molecules CH<sub>3</sub>OH, CH<sub>3</sub>OCH<sub>3</sub>, C<sub>2</sub>H<sub>5</sub>OH, CH<sub>3</sub>OCH, CH<sub>3</sub>CN, C<sub>2</sub>H<sub>3</sub>CN and C<sub>2</sub>H<sub>5</sub>CN in the hot core associated with high-mass star formation region G34.26+0.15, Submillimeter Array observations were made with its 230 GHz receiver. The molecular gas distribution has shown that the oxygen- and nitrogen-containing molecules peak at different positions. Comparing the spatial distributions with rotational temperatures and fractional abundances of the observed molecules, we discuss the possible chemical origination of these organic molecules.

**Key words:** astrochemistry — molecular processes — line: identification — ISM: abundances

### 1 INTRODUCTION

Ultracompact HII (UC HII) regions are thought to be indicators of high-mass star formation (Churchwell 2002). Hot molecular cores (HMCs) represent the immediate environments where massive stars were recently born. HMCs are observed to have compact structure, be dense, have a warm molecular gas component and are frequently found around high-mass star formation sites associated with UC HII regions (Kurtz et al. 2000). Their hot core phase also shows rich molecular line emissions at (sub)millimeter wavelengths. Therefore, observations of HMCs are crucial for characterizing physical and chemical structures associated with massive star formation. G34.26+0.15 is one of the well studied high-mass star formation regions, at a distance of 3.7 kpc from the Sun (Kuchar & Bania 1994). Molecular line observations suggested that massive star formation is taking place in this star formation complex (e.g., Liu et al. 2013). Three UC HII regions (“A, B and C”) have been identified by continuum observations at centimeter wavelengths (Wood & Churchwell 1989; Avalos et al. 2006). Single dish observations of multiple organic molecular lines suggested that hot gas components are present in the G34.26+0.15 star formation complex (Hatchell et al. 1998; Fontani et al. 2007), while higher spatial resolution observations with interferometers showed that the hot gas components (which are HMCs) are associated with UC HII region “C” (Mookerjea et al. 2007). However, only a few molecular transitions of C<sub>2</sub>H<sub>5</sub>CN, CH<sub>3</sub>OCH<sub>3</sub> and HCOOCH<sub>3</sub> were observed in previous observations with higher spatial resolution (Mookerjea et al. 2007). To ob-

tain accurate physical parameters, many transitions of the same species spanning wide energy ranges are needed. In this paper, we present Submillimeter Array (SMA)<sup>1</sup> observations of multiple transitions in organic molecules. Using the multiple lines, we derive relatively accurate physical parameters and discuss the physical and chemical environments of the G34.26+0.15 HMC.

### 2 OBSERVATIONS

We obtained the data from the SMA data archive. The SMA observations of G34.26+0.15 were made in April 2011 using its compact array. The phase tracking center of the observations was R.A. (J2000) = 18<sup>h</sup>53<sup>m</sup>18.573<sup>s</sup> and Dec. (J2000) = 01°14′58.3″. The double spectral sideband mode was used and centered at 227.5 GHz for the lower sideband and 239.5 GHz for the upper sideband. The spectral resolution was 0.8125 MHz, which corresponds to a velocity resolution of  $\sim 1.1$  km s<sup>-1</sup>. Quasi-stellar object (QSO) 3C 279 and Saturn were employed for bandpass calibration. QSOs 1911–201 and 1741–038 were observed for phase calibration. Absolute flux density was scaled by observing Saturn, and is estimated to be accurate to within 15%. Data reduction was done in Miriad. The resultant full width at half maximum beam size was 3.1″ × 2.5″ (PA = -58°). The 1 $\sigma$  rms of continuum and line emissions were 0.03 and 0.2 Jy beam<sup>-1</sup>, respectively.

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### 3 RESULTS

#### 3.1 Continuum Image

The continuum image of G34.26+0.15 at 1.3 mm is shown in Figure 1. In this image, a compact source structure is revealed and the continuum peak at the 1.3 mm waveband is consistent with the UC HII region “C.” Therefore the emission at 1.3 mm may contain free-free emission. Two dimensional Gaussian fitting to the continuum image gives the peak flux of  $9.5 \pm 0.4 \text{ Jy beam}^{-1}$ , the integrated flux of  $13.6 \pm 0.6 \text{ Jy}$  and deconvolved size of the core of  $2.8'' \times 1.8''$  (P.A. =  $48^\circ$ ). The flux density of 1.3 cm continuum emission is  $\sim 5 \text{ Jy}$  (Mookerjea et al. 2007). Assuming that the 1.3 cm continuum emission is optically thin and the spectral index is 0.2, we estimate the free-free emission contribution of 7.9 Jy at 1.3 mm toward the continuum core of G34.26+0.15. The 1.3 mm continuum flux density corrected for free-free emission is  $5.7 \pm 0.5 \text{ Jy}$ . As done by Qin et al. (2010), taking a mean dust grain size of 0.1  $\mu\text{m}$ , grain emissivity of  $2 \times 10^{-5}$  at 1.3 mm and a dust temperature of 250 K for G34.26+0.15, we derived the  $\text{H}_2$  gas mass and column density of  $182 \pm 16 M_\odot$  and  $6.4 \pm 0.6 \times 10^{24} \text{ cm}^{-2}$ .

#### 3.2 Molecular Lines

The double sideband spectral mode of the SMA observations covers a bandwidth of 8 GHz.

Figure 2 only shows the SMA spectrum in which the 2 GHz spectrum is at the continuum peak position. One can see that many molecular transitions populate there. The identified species include complex oxygen-bearing molecules, nitrogen-bearing molecules and other molecules. Complex organic molecules of  $\text{CH}_3\text{OH}$ ,  $\text{CH}_3\text{OCHO}$ ,  $\text{C}_2\text{H}_5\text{CN}$  and  $\text{HC}_3\text{N } v_7 = 1$  have strong emissions. In this paper, we are only interested in the organic molecules of  $\text{CH}_3\text{OH}$ ,  $\text{CH}_3\text{OCH}_3$ ,  $\text{CH}_3\text{OCHO}$ ,  $\text{C}_2\text{H}_5\text{OH}$ ,  $\text{C}_2\text{H}_3\text{CN}$ ,  $\text{C}_2\text{H}_5\text{CN}$ ,  $\text{CH}_3\text{CN}$  and  $\text{CH}_3\text{CCH}$ , since more than three transitions of each of these molecules are observed, which span a wide energy range and enable us to derive physical parameters accurately. Molecular line parameters are obtained from Gaussian fitting to each transition and are presented in Table 1.

#### 3.3 Gas Distribution

The spatial gas distributions of specific molecules are important for study of the related chemistry, since these distributions indicate the formation routes of specific molecules. The sample images of oxygen-bearing molecules and nitrogen-bearing molecules are shown in Figure 3.

According to Figure 3, there are spatial separations between oxygen-bearing and nitrogen-bearing molecules in G34.26+01.5, and peak positions of nitrogen-bearing molecules coincide with the peaks in continuum emission. The peaks of oxygen-bearing molecules move to the northeast of the continuum emission. The spatial separation of nitrogen- and oxygen-bearing molecules is also observed

in G19.61–0.23, Orion KL and the W3(OH) complex (e.g., Blake et al. 1987; Qin et al. 2010, 2015).

### 4 DATA ANALYSIS

As seen in Table 1, multiple transitions of  $\text{CH}_3\text{OH}$ ,  $\text{CH}_3\text{OCH}_3$ ,  $\text{CH}_3\text{OCHO}$ ,  $\text{C}_2\text{H}_5\text{OH}$ ,  $\text{C}_2\text{H}_3\text{CN}$ ,  $\text{C}_2\text{H}_5\text{CN}$ ,  $\text{CH}_3\text{CN}$  and  $\text{CH}_3\text{CCH}$  are observed, spanning a wide energy range. Following the method used by Qin et al. (2010), we apply the rotational temperature diagram (RTD) method to calculate physical parameters of these molecules. One can use an RTD to derive rotational temperature and column density of a specific species in case multiple transitions, spanning a wide energy range, are observed (Goldsmith & Langer 1999). The energy level population of molecular transitions can be described by a single rotational temperature, if local thermodynamic equilibrium (LTE) holds and the molecular gas is optically thin.

Figure 4 presents RTDs of the observed molecules. The linear least-squares fittings to the data are made for molecules that are observed in this work. The derived rotational temperatures and column densities of the studied molecules are presented in Table 2. The fractional abundances of specific molecules relative to  $\text{H}_2$  are derived by use of the ratio of molecular column densities to the  $\text{H}_2$  column density. The results presented in Table 2 show that complex organic oxygen-containing molecules  $\text{CH}_3\text{OH}$ ,  $\text{CH}_3\text{OCH}_3$ ,  $\text{CH}_3\text{OCHO}$  and  $\text{C}_2\text{H}_5\text{OH}$  have high fractional abundances relative to  $\text{H}_2$  of  $0.06 - 1.4 \times 10^{-7}$ , with gas temperatures  $< 200 \text{ K}$ . The nitrogen-containing molecules ( $\text{C}_2\text{H}_3\text{CN}$ ,  $\text{C}_2\text{H}_5\text{CN}$ , and  $\text{CH}_3\text{CN}$ ) have high temperatures of  $> 290 \text{ K}$  and low fractional abundances ranging from  $1.14 \times 10^{-10}$  to  $1.2 \times 10^{-18}$ . The situation is similar in the massive star forming regions Orion KL, G19.61–0.23 and W3(OH) in that the oxygen-containing molecules have higher fractional abundances but lower gas temperatures. Higher gas temperatures and lower fractional abundances are derived towards nitrogen-bearing molecules (Blake et al. 1987; Qin et al. 2010, 2015). In addition,  $\text{CH}_3\text{CCH}$  has an abundance of  $\sim 4.8 \times 10^{-9}$  and a lower rotational temperature of 63 K. Note that single dish observations of  $\text{C}_2\text{H}_3\text{CN}$ ,  $\text{C}_2\text{H}_5\text{CN}$ ,  $\text{CH}_3\text{OCH}_3$  and  $\text{CH}_3\text{OCHO}$  (Fontani et al. 2007) gave lower gas temperatures when compared to our results, and probably single dish observations contained cold gas components which provided averaged gas temperatures.

### 5 DISCUSSION AND CONCLUSIONS

Multiple transitions spanning a wide energy range are observed toward G34.16–0.25, which enable us to accurately derive physical parameters of gas temperature and column density. The peak offsets of oxygen- and nitrogen-containing molecules with respect to the peak position of the continuum source were observed toward G34.26+0.15 (see Fig. 3). The calculations have shown that the oxygen-containing and nitrogen-containing molecules have different molecular abundances and rotational temperatures. The

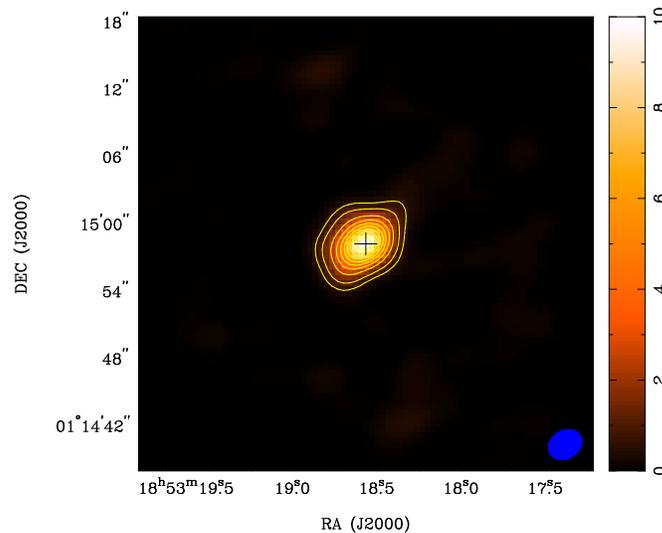
Table 1 Line Parameters

Molecule	Frequency (MHz)	Transition	$S_{ij}\mu^2$ (debye <sup>2</sup> )	$E_U$ (K)	$I$ (Jy beam <sup>-1</sup> )	$\Delta V$ (km s <sup>-1</sup> )	
CH <sub>3</sub> OH	229589.073	15 <sub>(4,11)</sub> -16 <sub>(3,13)</sub>	4.59198	374.43849	13.2± 0.1	7.2± 0.1	
	239731.344	5 <sub>(1,5)</sub> -4 <sub>(1,4)</sub>	3.88508	49.05924	15.1± 0.1	8.1± 0.1	
	240241.502	5 <sub>(3,2)</sub> -6 <sub>(2,4)</sub>	0.98197	82.5315	11.8± 0.1	6.3± 0.1	
	240740.621	26 <sub>(3,23)</sub> -26 <sub>(2,24)</sub>	26.79785	863.97661	4.4± 0.1	5± 0.1	
	241042.683	22 <sub>(-6,16)</sub> -23 <sub>(-5,18)</sub>	6.39588	775.56583	2.3± 0.1	4± 0.1	
	239397.838	16 <sub>(-3,14)</sub> -17 <sub>(0,17)</sub>	0.14009	378.27451	2.2± 0.1	3.5± 0.1	
	227094.601	21 <sub>(1,20)</sub> -21 <sub>(0,21)</sub>	11.59311	557.07117	8.9± 0.1	6.2± 0.1	
	CH <sub>3</sub> OCH <sub>3</sub>	227648.12	26 <sub>(5,21)</sub> -26 <sub>(4,22)</sub> EA	84.89383	355.76623	3.1± 0.1	5.9± 0.1
		227657.0248	24 <sub>(3,22)</sub> -24 <sub>(2,23)</sub> EE	210.14176	285.56309	3.3± 0.1	8.8± 0.2
		226346.0514	14 <sub>(1,13)</sub> -13 <sub>(2,12)</sub> AA	42.91412	98.86426	5.4± 0.1	6.1± 0.1
226495.4549		22 <sub>(1,21)</sub> -22 <sub>(0,22)</sub> EE	107.91955	232.58339	2.1± 0.1	8.9± 0.2	
239326.2099		23 <sub>(1,22)</sub> -23 <sub>(0,23)</sub> EE	107.96363	253.09072	2.2± 0.1	10.6± 0.2	
238958.2129		32 <sub>(1,31)</sub> -32 <sub>(4,28)</sub> AA	126.48193	501.84005	1.8± 0.1	5.6± 0.2	
238016.1043		27 <sub>(2,25)</sub> -27 <sub>(3,25)</sub> EE	218.16053	354.96978	1.8± 0.1	6.3± 0.2	
240985.0778		5 <sub>(3,3)</sub> -4 <sub>(2,2)</sub> EE	54.77959	26.30846	4.1± 0.1	7.8± 0.1	
228424.4096		26 <sub>(3,24)</sub> -25 <sub>(4,21)</sub> AA	29.05868	331.64699	0.8± 0.1	4.3± 0.4	
CH <sub>3</sub> OCHO		228211.291	18 <sub>(3,15)</sub> -17 <sub>(3,14)</sub> E	46.25056	297.16473	1.9± 0.1	4.4± 0.2
	228628.876	18 <sub>(5,13)</sub> -17 <sub>(5,12)</sub> E	44.23729	118.78854	5.0± 0.1	5.3± 0.1	
	228651.404	18 <sub>(5,13)</sub> -17 <sub>(5,12)</sub> A	44.24797	118.78242	5± 0.1	5.5± 0.1	
	229405.021	18 <sub>(3,15)</sub> -17 <sub>(3,14)</sub> E	46.11741	110.73904	5.8± 0.1	6.1± 0.1	
	229420.342	18 <sub>(3,15)</sub> -17 <sub>(3,14)</sub> A	46.12064	110.73143	6.3± 0.1	5.8± 0.1	
	240021.14	19 <sub>(3,16)</sub> -18 <sub>(3,15)</sub> E	48.61567	122.25813	7.2± 0.1	7.5± 0.1	
	240034.673	19 <sub>(3,16)</sub> -18 <sub>(3,15)</sub> A	48.63142	122.2513	5.2± 0.1	8.8± 0.1	
	237807.626	19 <sub>(6,13)</sub> -18 <sub>(6,12)</sub> E	45.4611	136.79123	4.6± 0.1	5.4± 0.1	
	237829.831	19 <sub>(6,13)</sub> -18 <sub>(6,12)</sub> A	45.55653	136.78611	4.8± 0.1	5.4± 0.1	
	238156.8625	22 <sub>(1,22)</sub> -21 <sub>(1,21)</sub> A	58.16589	133.83147	11.2± 0.1	8± 0.1	
	238190.191	7 <sub>(6,1)</sub> -6 <sub>(5,2)</sub> A	2.52123	40.41583	0.9± 0.1	6.7± 0.5	
	238368.799	19 <sub>(3,16)</sub> -18 <sub>(3,15)</sub> A	48.61357	308.9218	1.6± 0.1	8.9± 0.3	
	225756.199	18 <sub>(5,13)</sub> -17 <sub>(5,12)</sub> E	43.47338	305.4027	1.5± 0.1	5± 0.2	
	225928.659	6 <sub>(6,0)</sub> -5 <sub>(5,1)</sub> A	2.51986	36.29203	2.4± 0.1	4.2± 0.1	
	226435.521	21 <sub>(0,21)</sub> -20 <sub>(0,20)</sub> E	55.61866	308.8777	3.6± 0.1	6.4± 0.1	
	226713.06	20 <sub>(2,19)</sub> -19 <sub>(2,18)</sub> E	52.04002	120.22039	4.9± 0.1	7.7± 0.1	
	226718.688	20 <sub>(2,19)</sub> -19 <sub>(2,18)</sub> A	52.04707	120.20699	5.4± 0.1	7.4± 0.1	
	226773.13	20 <sub>(1,19)</sub> -19 <sub>(1,18)</sub> E	52.03546	120.21637	4.7± 0.1	7.7± 0.1	
	226778.786	20 <sub>(1,19)</sub> -19 <sub>(1,18)</sub> A	52.0425	120.20297	4.9± 0.1	9.3± 0.1	
	227019.55	19 <sub>(2,17)</sub> -18 <sub>(2,16)</sub> E	48.76106	116.573	5.5± 0.1	6.1± 0.1	
227028.121	19 <sub>(2,17)</sub> -18 <sub>(2,16)</sub> A	48.76691	116.56277	4.6± 0.1	9.1± 0.1		
227599.261	18 <sub>(3,15)</sub> -17 <sub>(3,14)</sub> A	46.0925	297.48196	1.5± 0.1	4.7± 0.2		
C <sub>2</sub> H <sub>5</sub> OH	227891.911	13 <sub>(1,12)</sub> -12 <sub>(1,11)</sub> , vt= 1-1	20.6452	140.01443	2.2± 0.1	5.1± 0.2	
	228886.6501	9 <sub>(3,6)</sub> -9 <sub>(2,8)</sub> , vt= 1-0	5.3747	110.07161	0.7± 0.1	4.8± 0.6	
	229491.5539	46 <sub>(7,39)</sub> -46 <sub>(6,40)</sub> , vt= 0-0	0.24656	1020.70466	1.9± 0.1	4.7± 0.2	
	240110.238	27 <sub>(2,25)</sub> -27 <sub>(1,26)</sub>	29.31314	327.23803	0.8± 0.1	4.4± 0.5	
	240838.7471	14 <sub>(1,13)</sub> -13 <sub>(0,13)</sub> , vt= 0-1	9.09134	147.09916	1.6± 0.1	4.1± 0.2	
	238568.1891	6 <sub>(1,5)</sub> -5 <sub>(0,5)</sub> , vt= 1-0	4.24277	80.72611	1.1± 0.1	5.3± 0.4	
	239478.079	14 <sub>(2,13)</sub> -13 <sub>(2,12)</sub> , vt= 0-0	21.90963	148.43824	2.2± 0.1	5± 0.2	
	239551.366	14 <sub>(2,13)</sub> -13 <sub>(2,12)</sub> , vt= 1-1	21.90948	153.11156	2.2± 0.1	4.9± 0.2	
	225796.891	13 <sub>(4,9)</sub> -12 <sub>(4,8)</sub> , vt= 0-0	18.80219	152.09896	2± 0.1	6± 0.2	
	226581.339	26 <sub>(2,24)</sub> -26 <sub>(1,25)</sub>	28.24034	304.69227	0.9± 0.1	4.2± 0.4	
	227294.752	13 <sub>(3,10)</sub> -12 <sub>(3,9)</sub> , vt= 1-1	19.66545	148.56961	1.7± 0.1	5.2± 0.2	
	C <sub>2</sub> H <sub>3</sub> CN	226256.8803	24 <sub>(2,23)</sub> -23 <sub>(2,22)</sub>	1040.33362	144.79326	0.7± 0.1	5.3± 0.5
238726.8113		26 <sub>(1,26)</sub> -25 <sub>(1,25)</sub>	1133.0669	157.37752	2.4± 0.1	4.8± 0.2	
238796.2925		25 <sub>(3,22)</sub> -24 <sub>(3,21)</sub>	1075.85721	167.9148	1.2± 0.1	6.2± 0.3	
227897.6076		24 <sub>(7,17)</sub> -23 <sub>(7,16)</sub>	958.82619	242.46929	0.7± 0.1	7.6± 0.7	
227906.7091		24 <sub>(6,18)</sub> -23 <sub>(6,17)</sub>	982.32004	214.45704	0.9± 0.1	7.8± 0.5	
228087.2448		24 <sub>(11,14)</sub> -23 <sub>(11,13)</sub>	827.73655	396.9138	0.7± 0.1	4.3± 0.5	
228104.6144		24 <sub>(4,21)</sub> -23 <sub>(4,20)</sub>	1018.75591	171.33744	0.9± 0.1	3.9± 0.4	
228160.3049		24 <sub>(4,20)</sub> -23 <sub>(4,19)</sub>	1018.75515	171.34687	1.5± 0.1	5.6± 0.3	
229087.0477		24 <sub>(3,21)</sub> -23 <sub>(3,20)</sub>	1031.47236	156.45448	0.7± 0.1	5.7± 0.6	
229647.8388		25 <sub>(1,25)</sub> -24 <sub>(1,24)</sub>	1089.34365	145.92055	1.9± 0.1	4.7± 0.2	

C <sub>2</sub> H <sub>5</sub> CN	227780.9718	25 <sub>(3,22)</sub> –24 <sub>(3,21)</sub>	365.32437	150.84922	3.8 ± 0.1	5.8 ± 0.1
	228483.1359	25 <sub>(2,23)</sub> –24 <sub>(2,22)</sub>	368.05881	146.59165	4.4 ± 0.1	5.3 ± 0.1
	229265.16	26 <sub>(2,25)</sub> –25 <sub>(2,24)</sub>	382.45273	154.02143	4.4 ± 0.1	5.3 ± 0.1
	240319.3368	28 <sub>(1,28)</sub> –27 <sub>(1,27)</sub>	414.0537	169.2645	5.1 ± 0.1	5.3 ± 0.1
	240429.1835	28 <sub>(0,28)</sub> –27 <sub>(0,27)</sub>	414.0245	169.24387	4.1 ± 0.1	5.6 ± 0.1
CH <sub>3</sub> CN	2238766.0489	13 <sub>9</sub> –12 <sub>9</sub>	286.97085	658.1578	4.1 ± 0.1	5.8 ± 0.1
	238843.926	13 <sub>8</sub> –12 <sub>8</sub>	171.20193	537.03504	5 ± 0.1	6.5 ± 0.1
	238912.7154	13 <sub>7</sub> –12 <sub>7</sub>	195.68053	430.09712	8.1 ± 0.1	6 ± 0.1
	238972.3895	13 <sub>6</sub> –12 <sub>6</sub>	433.70052	337.36833	13.6 ± 0.1	7.1 ± 0.1
	239022.9241	13 <sub>5</sub> –12 <sub>5</sub>	234.7829	258.86994	13.6 ± 0.1	7 ± 0.1
	239064.2988	13 <sub>4</sub> –12 <sub>4</sub>	249.46102	194.61954	16.3 ± 0.1	6.7 ± 0.1
	239096.4966	13 <sub>3</sub> –12 <sub>3</sub>	521.81905	144.6318	17.1 ± 0.1	8.4 ± 0.1
	239119.5044	13 <sub>2</sub> –12 <sub>2</sub>	269.05112	108.91819	18.5 ± 0.1	6.6 ± 0.1
	239133.3129	13 <sub>1</sub> –12 <sub>1</sub>	273.94228	87.48682	17.7 ± 0.1	12.8 ± 0.1
CH <sub>3</sub> CCH	239179.2812	14 <sub>4</sub> –13 <sub>4</sub>	4.81196	201.69776	1.4 ± 0.1	4.6 ± 0.2
	239211.2149	14 <sub>3</sub> –13 <sub>3</sub>	9.99811	151.14227	4.8 ± 0.1	5.4 ± 0.1
	239234.0336	14 <sub>2</sub> –13 <sub>2</sub>	5.1327	115.02278	3.4 ± 0.1	5.5 ± 0.1
	239252.2938	14 <sub>0</sub> –13 <sub>0</sub>	5.24012	86.1221	3.7 ± 0.1	10.7 ± 0.1

**Table 2** Rotational Temperature, Column Density and Abundance

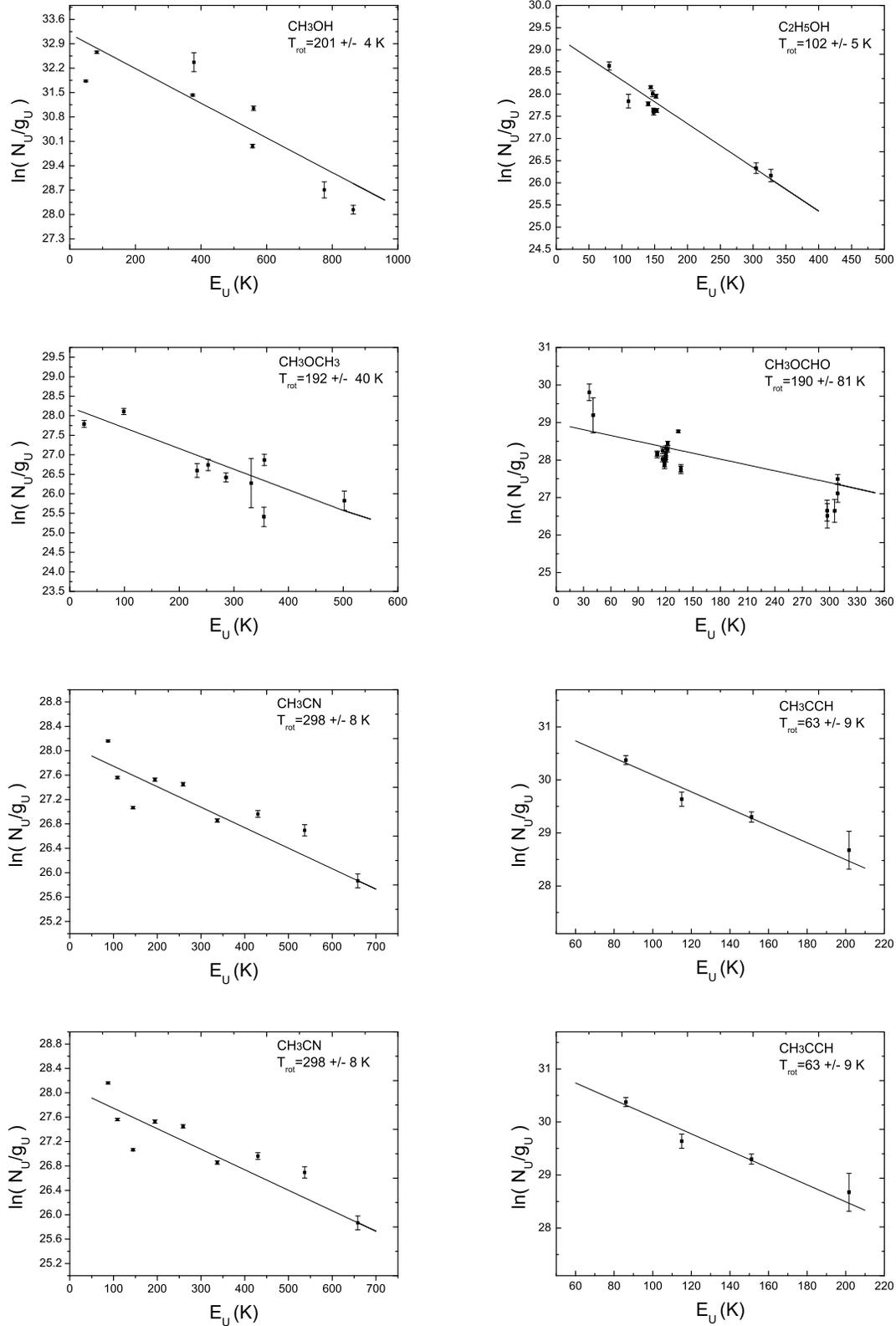
Molecule	Temperature (K)	Column Density (cm <sup>-2</sup> )	Abundance
CH <sub>3</sub> OH	201 ± 4	$(9.1 ± 0.4) × 10^{17}$	$(1.4 ± 0.1) × 10^{-7}$
CH <sub>3</sub> OCH <sub>3</sub>	192 ± 40	$(4.6 ± 1.8) × 10^{17}$	$(7.2 ± 3.9) × 10^{-8}$
CH <sub>3</sub> OCHO	190 ± 81	$(1.2 ± 0.9) × 10^{17}$	$(1.9 ± 1.4) × 10^{-8}$
C <sub>2</sub> H <sub>5</sub> OH	102 ± 5	$(3.6 ± 0.4) × 10^{16}$	$(5.6 ± 0.6) × 10^{-9}$
CH <sub>3</sub> CCH	63 ± 9	$(3.1 ± 1.1) × 10^{16}$	$(4.8 ± 1.7) × 10^{-9}$
C <sub>2</sub> H <sub>3</sub> CN	290 ± 137	$(7.3 ± 5.6) × 10^{14}$	$(1.1 ± 0.9) × 10^{-10}$
C <sub>2</sub> H <sub>5</sub> CN	300 ± 130	$(8.1 ± 6.5) × 10^{15}$	$(1.3 ± 1) × 10^{-9}$
CH <sub>3</sub> CN	298 ± 8	$(7.9 ± 0.4) × 10^{15}$	$(1.3 ± 0.1) × 10^{-9}$

**Fig. 1** 1.3 mm continuum image of G34.26+0.15, with both a color scale and contours. The contours are 5%, 10%, 20%, 30%, 40%, 50%, 60%, 70%, 80%, 90% and 100% of the peak value. The plus sign indicates the peak position of the continuum.

spatial distributions, gas temperatures and fractional abundances imply there are physical and chemical differences among various species. We discuss the chemical origin of the observed molecules in the following:

CH<sub>3</sub>OH has the highest fractional abundance of  $1.4 × 10^{-7}$  when compared with other organic molecules in our observations. Higher CH<sub>3</sub>OH abundance is also observed in the massive star-forming regions W51N, G19.61 and W3(OH) (Rong et al. 2016; Qin et al. 2010,





**Fig. 4** RTDs. The linear least-squares fittings to the observed transitions are shown as straight lines. The filled symbols represent the observed transitions and  $3\sigma$  errors are indicated by vertical bars.

butions of  $\text{CH}_3\text{CN}$ ,  $\text{C}_2\text{H}_5\text{CN}$  and  $\text{C}_2\text{H}_3\text{CN}$  indicate that these nitrogen-containing species may have similar chemical originations and be formed in high temperature gas-phase reactions.  $\text{CH}_3\text{OH}$  and  $\text{CH}_3\text{CN}$  are representative oxygen- and nitrogen-containing molecules and the abundance ratio of  $\text{CH}_3\text{OH}$  to  $\text{CH}_3\text{CN}$ ,  $115 \pm 8$ , is appropriate. This value is roughly consistent with that in the UC HII region source W3(OH) ( $\sim 138$ ) and larger than that in high mass protostar W3( $\text{H}_2\text{O}$ ) ( $\sim 16$ ) (see Qin et al. 2015) and other high-mass young stellar objects (Bisschop et al. 2007; Gibb et al. 2000). The larger  $\text{CH}_3\text{OH}$  to  $\text{CH}_3\text{CN}$  abundance ratio in G34.26+0.15 may imply that the surrounding gas can be effectively heated by the UC HII region and then  $\text{CH}_3\text{OH}$  is evaporated from the grain surface as the cloud warms up. The derived  $\text{CH}_3\text{CCH}$  abundance of  $4.8 \times 10^{-9}$  is in good agreement with that in other high mass star-forming regions and larger than that in cold cores (Miettinen et al. 2006).  $\text{CH}_3\text{CCH}$  can be formed on dust grains through the reaction between CH and  $\text{C}_2\text{H}_4$  (Canosa et al. 1997).  $\text{CH}_3\text{CCH}$  peaks at the same position as the nitrogen-bearing molecules and has lower gas temperature, which may indicate that  $\text{CH}_3\text{CCH}$  and nitrogen-bearing molecules are synthesized on the same site but are distributed in different layers of the cloud as temperature increases.

In summary, the SMA observations showed that there is a spatial separation between nitrogen- and oxygen-containing molecules. The nitrogen-bearing molecules have higher rotational temperatures but lower molecular abundances compared to the oxygen-containing species. These two groups of molecules have different chemical originations.

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