Molecular jet emission and a spectroscopic survey of S235AB

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ABSTRACT

Context. The S235AB star forming region houses a massive young stellar object which has recently been reported to exhibit possible evidence of jet rotation – an illusive yet crucial component of disk aided star formation theories.

Aims. We confirm the presence of a molecular counterpart to the jet and we also study the molecular environment in S235AB.

Methods. We search for velocity wings in the line emission of thermal SiO (J = 2–1, v = 0), a tracer of shocked gas, which would indicate the presence of jet activity. Utilising other lines detected in our survey we use the relative intensities of intra species transitions, isotopes, and hyperfine transitions to derive opacities, temperatures, column densities, and abundances of various molecular species in S235AB.

Results. The SiO (J = 2–1, v = 0) emission exhibits velocity wings of up to 75 km s⁻¹ above and below the velocity of the star, indicating the presence of a jet. The molecular environment describes an evolutionary stage resembling a hot molecular core.

Key words. stars: massive – ISM: jets and outflows

1. Introduction

The target of our study, S235AB, is a star forming region that houses a massive young stellar object (MYSO) called S235AB-MIR, which was recently reported to exhibit fast (~50 km s⁻¹) jet-tracing water masers with a signature of rotation (Burns et al. 2015). The confirmation of a rotating jet in an MYSO would have a great impact on the theories of star formation as it would corroborate a magneto-centrifugal launching mechanism for jets from massive young stars (see Konigl & Pudritz 2000).

S235AB-MIR is known to have slow molecular outflows (Felli et al. 2004); however, aside from the fast water maser velocities, there is no evidence in the literature of a fast molecular jet in S235AB-MIR. In Felli et al. (2004) the authors claimed to have found a continuum jet aligned with the water maser jet; however, they later retracted the claim in Felli et al. (2006). As a follow-up to the maser jet results of Burns et al. (2015) the primary aim of our spectral line survey was to find the molecular counterpart to the water maser jet in S235AB, which should be readily identifiable from doppler broadened line “wings” seen in the thermal gas jet tracer SiO (J = 2–1, v = 0) (e.g. López-Sepulcre et al. 2011). The secondary aim of our observations was to further investigate the physical gas conditions in S235AB via basic astrochemical study using auxiliary spectral lines.

S235AB is known to have a dense molecular core centered on S235AB-MIR and molecular outflows mapped in HCO⁺ (1–0) and C⁴S (5–4) (Felli et al. 2004), CS (7–6) (Wu et al. 2010), and ¹³CO (Felli et al. 1997). The Spitzer colours indicate that S235AB-MIR has a mass of 11 M☉, making it the only MYSO in the region (Dewangan & Anandarao 2011), and non-detection of centimetre emission in S235AB attests to its youth (Tofani et al. 1995; Felli et al. 2006).

2. Observations and data reduction

We conducted molecular line observations of S235AB in March and April of 2015 with the 45 m radio telescope of the Nobeyama Radio Observatory (NRO), a branch of the National Astronomical Observatory of Japan.

We observed in two-sideband mode using the TZ receiver and we observed horizontal and vertical linear polarisations simultaneously. The upper sideband (USB) and lower sideband (LSB) central frequencies were 85.85 and 97.85 GHz, respectively, with band widths of 4 GHz each. The beam size was about 18″. The SAM45 spectrometer provided 16 independent frequency arrays (8 per polarisation), which we organised to provide continuous coverage of a wide frequency range, and to include lines of particular interest. The frequency resolution was 244 kHz, providing a bandwidth of 1000 MHz per array.

Observing coordinates were centred on the position of S235AB-MIR at (α,δ)J2000.0 = (05h40m53.384 +35°41′48″.447). Sky-level subtraction was performed using a region of empty sky at coordinates (α,δ)J2000.0 = (05h40m58.30 +35°41′48″.60). Absolute flux calibration was performed using the chopper wheel method, empty sky, and hot load. Pointing accuracy...
Table 1. Summary of detected molecular lines and their measured parameters.

| Molecular formula | Transition | $T_A^{sys}$ [K] | Obs frequency [MHz] | $\Delta\nu$ [MHz] | $\nu_{peak}$ [km s$^{-1}$] | rms [K] |
|-------------------|------------|-----------------|--------------------|-----------------|-----------------|-----------|
| CH$_3$OH | 2(1, 2)–1(1, 1) A++ | 0.230 | 95,914,406 | 0.819 | –17.22 | 0.012 |
| C$^{34}$S | 2–1 | 1.037 | 96,413,016 | 0.852 | –17.13 | 0.008 |
| CH$_3$OH | 2(–1, 2)–1(–1, 1) E | 0.962 | 96,739,445 | 0.778 | –17.18 | 0.006 |
| CH$_3$OH | 2(0, 2)–1(0, 1) A++ | 1.330 | 96,741,422 | 0.778 | –17.14 | 0.012 |
| CH$_3$OH | 2(0, 2)–1(0, 1) E | 0.461 | 96,744,617 | 0.836 | –17.21 | 0.016 |
| CH$_3$OH | 2(1, 1)–1(1, 0) E | 0.210 | 96,755,602 | 0.770 | –17.29 | 0.007 |
| C$^{33}$S | 2–1 1/2–1/2 | 0.030 | 97,169,438 | 0.811 | –16.96 | 0.002 |
| C$^{33}$S | 2–1 1/2–1/2 | 0.151 | 97,171,828 | 0.811 | –16.96 | 0.002 |
| C$^{33}$S | 2–1 1/2–1/2 | 0.038 | 97,175,188 | 1.155 | –16.74 | 0.003 |
| OCS | 8–7 | 0.048 | 97,301,203 | 0.786 | –16.98 | 0.005 |
| CH$_3$OH | 2(1, 1)–1(1, 0) A– | 0.191 | 97,382,859 | 0.803 | –17.16 | 0.005 |
| $^{34}$SO | $N, J = 2, 3, 1$, 2 | 0.094 | 97,715,461 | 0.868 | –17.18 | 0.002 |
| CS | 2–1 | 3.639 | 97,981,016 | 2.146 | –17.19 | 0.019 |
| CH$_3$CHO | 5(1, 4)–4(1, 3) E | 0.030 | 98,863,484 | 1.262 | –17.16 | 0.004 |
| SO | $N, J = 2, 3, 1$, 2 | 1.531 | 99,299,953 | 0.901 | –17.14 | 0.013 |
| CH$_3$OH | 5(−1, 5)–4(0, 4) E | 2.471 | 84,521,188 | 3.170 | –16.94 | 0.020 |
| OCS | 7–6 | 0.075 | 85,139,250 | 0.606 | –17.51 | 0.004 |
| HC$^{18}$O$^+$ | 1–0 | 0.035 | 85,162,375 | 0.959 | –17.53 | 0.005 |
| c-C$_2$H$_2$ | 2(1, 2)–1(0, 1) | 0.093 | 85,338,969 | 0.614 | –17.22 | 0.005 |
| CH$_3$CH | 5(2)–(4) | 0.129 | 85,437,859 | 0.721 | –16.96 | 0.004 |
| CH$_3$CH | 5(1)–(4) | 0.058 | 85,455,727 | 0.688 | –17.22 | 0.003 |
| CH$_3$CH | 5(0)–(4) | 0.085 | 85,457,219 | 0.287 | –17.72 | 0.001 |
| HC$^{15}$N | 1–0 | 0.157 | 86,054,992 | 0.795 | –17.09 | 0.005 |
| SO | $N, J = 2, 3, 1$, 2 | 0.376 | 86,093,984 | 0.721 | –17.00 | 0.005 |
| H$^{13}$CN | 1–0 $F = 1$ | 0.322 | 86,338,750 | 0.786 | –17.05 | 0.005 |
| H$^{13}$CN | 1–0 $F = 2$ | 0.593 | 86,340,180 | 0.811 | –17.01 | 0.004 |
| H$^{13}$CN | 1–0 $F = 0$ | 0.129 | 86,342,242 | 0.688 | –16.95 | 0.004 |
| HCO | 1(0, 1)–0(0, 0) 3/2–1/2 $F = 2$–1 | 0.044 | 86,670,648 | 0.664 | –16.40 | 0.005 |
| H$^{15}$CO$^+$ | 1–0 | 0.282 | 86,754,352 | 0.746 | –17.22 | 0.006 |
| SiO | 2–1 $v = 0$ | 0.056 | 86,847,023 | 1.901 | –17.04 | 0.001 |
| HN$^{13}$C | 1–0 $F = 2$ | 0.124 | 87,090,812 | 0.705 | –16.84 | 0.005 |
| C$_2$H | 1–0 3/2–1/2 $F = 1$–1 | 0.137 | 87,284,180 | 0.352 | –17.08 | 0.003 |
| C$_2$H | 1–0 3/2–1/2 $F = 2$–1 | 0.890 | 87,316,969 | 0.770 | –17.15 | 0.008 |
| C$_2$H | 1–0 3/2–1/2 $F = 1$–0 | 0.479 | 87,328,641 | 0.754 | –17.06 | 0.006 |
| C$_2$H | 1–0 1/2–1/2 $F = 1$–1 | 0.637 | 87,402,102 | 0.778 | –17.34 | 0.008 |
| C$_2$H | 1–0 1/2–1/2 $F = 0$–1 | 0.296 | 87,407,227 | 0.696 | –17.21 | 0.010 |
| C$_2$H | 1–0 1/2–1/2 $F = 1$–0 | 0.149 | 87,446,523 | 0.819 | –17.03 | 0.008 |

was checked using a nearby SiO maser source, RU Aur, every ~1–2 h.

During the March observations the atmospheric conditions were good with a typical temperature of $T_{sys} \approx 140$ K; however, strong wind conditions resulted in poor pointing accuracy and caused much of the on-source scans to suffer from beam warping and pointing offsets. Only 1 h of observing session provided usable data. During the April observations wind conditions were much better, providing good pointing accuracy (usually better than 3") and $T_{sys}$ was between 250 and 300 K. Data from the full four-hour observing session was used.

To detect as many molecular lines as possible we alternated between two frequency array setups, each of which covered different frequency ranges. The only frequency array common to both setups was that containing the SiO line, which was our transition of highest priority. Subsequently, the frequency array containing the SiO line was observed for the full observing time, while other frequency arrays were observed for roughly half of this time.

Data reduction was performed using the NEWSTAR software developed by the NRO. To reduce the aforementioned effect of the wind in the March observations, we inspected all scans individually. Scans with bad pointing were determined by the failure in detecting C$^{34}$S, a rarefied gas whose region of peak emission is known to be compact from the maps of Felli et al. (2004; their Fig. 9). This greatly improved the quality of integrated spectra.

After flagging bad scans, second-order polynomial baselines were fit to individual scans. Finally, all scans were integrated and polarisations were merged. Good scans on the target integrated to a total of 1 h for the March data and 2 h for the April data. The achieved sensitivity was typically better than 0.006 K for most frequency arrays, and reached 0.0025 K for the array containing the SiO emission. The antenna gain was roughly 4.4 Jy/K.
3. Results

Gaussian profiles were fit to spectral lines detected above 5 times rms the noise. The detections and line parameters are listed in Table 1. Detected molecular transitions were identified by cross-referencing with the Cologne Database for Molecular Spectroscopy (CDMS) catalogue (Müller et al. 2005).

3.1. SiO – jet tracer

Our survey detected the common jet tracer transition line of SiO (2–1). In low-temperature environments SiO molecules are confined to dust grains, resulting in a low gas abundance. In a shocked gas environment these molecules are released from dust grains and become abundant in the gas phase (Martin-Pintado et al. 1992). As such SiO is a useful tracer of shocked gas, and velocity wings in its spectrum indicate the presence of jets and outflows (López-Sepulcre et al. 2011). The spectrum of SiO (2–1) gas in S235AB is displayed in Fig. 1, showing wide velocity wings.

3.2. Rotation temperature

We detected multiple rotational transitions of sulphur monoxide (SO) such that it was possible to estimate its rotation temperature, $T_{\text{rot}}$, by using the “population diagram” method detailed in Goldsmith & Langer (1999). Emission was optically thin and the derived rotation temperature was $T_{\text{rot}} = 26 \pm 4$ K, which is similar to the value obtained by Felli et al. (2004) of $\sim 30$ K using transitions of CH$_3$CN (5–4).

We also measured the rotation temperature of methanol using transitions of CH$_3$OH (2–1), using lines of both E- and A-species. The population diagram is shown in Fig. 2, the slope gives $T_{\text{rot}} = 10 \pm 1$ K and the gas is optically thin. The line fit ignores the CH$_3$OH (5–4) E transition at $E_{\text{upper}} = 40$ K, which we discuss below. Methanol in this frequency regime is thought to be subthermally excited (Menten et al. 1988). Our estimate can therefore be considered the lower limit of the kinematic temperature of the methanol cloud.

Multiple transitions of OCS were detected, but were too weak to attain a meaningful temperature estimate.

3.3. Detection of a class I methanol maser

The CH$_3$OH (5–4) E transition (84.521188 GHz) deviates from its expected excitation conditions, as can be seen in the population diagram. This transition is known to exhibit maser behaviour (Menten et al. 1988); therefore, it seems that we have detected maser emission at this transition in S235AB (Fig. 3). The full line width of the emission is 11 km s$^{-1}$, which is far too large to emanate from a single maser feature and thus indicates a likely blending of multiple features. The central velocity of this class I methanol maser is close to that of the molecular core, suggesting that it is not directly tracing the jet/outflow system and could instead be produced in the dense ambient gas somewhere near the outflow.

3.4. Optical depth using hyperfine emission lines

Intensity ratios of the main line ($F = 2–1$) to the hyperfine satellite line ($F = 1–0$) of the C$_2$H molecule can be used to derive the gas optical depth via

\begin{equation}
\text{Line intensity ratio} = \frac{1 \text{ e}^{-aT}}{1 \text{ e}^{-aT}}.
\end{equation}

where $a$ is the intrinsic line intensity ratio determined by laboratory experiment. For C$_2$H ($F = 1–0$):($F = 2–1$), $a = 0.5$ in the optically thin limit (Tucker et al. 1974). Therefore, using Eq. (1) our observed line intensity ratio of $T_\lambda(1–0)/T_\lambda(2–1) = 0.55$ gives $T_{\text{rot}} = 0.39$. This value is typical of young, massive, star forming clumps (Sakai et al. 2010).

For optically thin H$^{13}$CN the intrinsic intensity ratios of the hyperfine transitions ($F = 1–1$):($F = 2–1$):($F = 1–0$) are 3:5:1 (Irvine & Schloerb 1984). Taking the transitions ($F = 2–1$):($F = 1–0$) = 5, our observed intensity ratio of $T_\lambda(F = 2–1)/T_\lambda(F = 1–0) = 4.36$ gives $T_{\text{H}^{13}\text{CN}} = 0.07$. 

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Table 2. Summary of derived physical parameters.

| Molecule   | $\tau$ | $T_{\text{rot}}$ [K] | $N_{\text{rot}}$ [cm$^{-2}$] | Molecular abundance |
|------------|--------|----------------------|-----------------------------|---------------------|
| C$_3$H     | 0.39   | 3.50E+15             | 3.5 x 10$^{-8}$             |                     |
| H$^3$CN    | 0.07   | 5.80E+13             | 5.7 x 10$^{-10}$            |                     |
| CS         | 5.1    | 6.63E+15             | 6.6 x 10$^{-8}$             |                     |
| SO         | 0.9    | 26 ± 4               | 9.76E+14                    | 9.7 x 10$^{-9}$     |
| CH$_3$OH   | <1     | 10 ± 1               | 7.85E+15                    | 7.8 x 10$^{-8}$     |

3.5. Optical depth using isotopic abundance

We detected two isotopologues of two sulphur-bearing species: C$^{34}$S and CS, and $^{33}$SO and SO. We estimated the optical depths of CS and SO using the isotopologue line intensity ratios, as was done in Eq. (1), where $a$ is the intrinsic intensity ratio of $^{32}$S/$^{34}$S = 22.5 (Kahane et al. 1988). Our measured intensity ratio of $T_A$(CS)/$T_A$(C$^{34}$S) = 4.9 gives $\tau_{\text{CS}} = 5.1$, and our intensity ratio of $T_A$(SO)/$T_A$(SO) = 15.2 gives $\tau_{\text{SO}} = 0.9$.

3.6. Column densities and abundances

To estimate column densities from individual molecular species, we used the prescription outlined in Goldsmith & Langer (1999) for species with measurable optical depths (see Sects. 3.4 and 3.5) except for the column density of CH$_3$OH, which was obtained from the population diagram. For other molecules the statistical weights, upper energy levels, partition functions, and Einstein coefficients for individual transitions were sourced from the Leiden Atomic and Molecular Database (LAMDBA; Schöier et al. 2005) and CDMS catalogues with an assumed excitation temperature of 18.75 K with the exception of SO, for which we used the measured rotation temperature (see Sect. 3.2).

Saito et al. (2007) estimated the molecular hydrogen column density of the molecular core in S235AB to be $1.0 \times 10^{22}$ cm$^{-2}$ based on their C$^{18}$O observations at comparable angular resolution. By comparing their results with the column densities measured in this work we estimated molecular abundances. These, alongside the aforementioned physical gas parameters, are summarised in Table 2.

4. Discussion and conclusions

Observations of S235AB by Sun et al. (2012) detected SiO emission at a peak brightness consistent with our observations; however, velocity wings were not detectable at the 0.017 K rms sensitivity of their observations. Our SiO spectrum (Fig. 1) exhibits wide velocity wings reaching up to 75 km s$^{-1}$ from the core velocity. Our deeper observations confirm the presence of a fast molecular jet in S235AB. Discovery of a high-velocity jet in this source is of particular significance in light of the recent water maser results of Burns et al. (2015). Those authors find both blueshifted and redshifted high-velocity ($\sim$50 km s$^{-1}$ from the core velocity) maser features in a bipolar configuration matching the alignment of the slow (<5 km s$^{-1}$ from the core velocity) NNW–SSE molecular outflow seen in HCO$^+$ and C$^{34}$S, reported by Felli et al. (2004).

The velocity of the SiO gas (~75 km s$^{-1}$) is much faster than that of the molecular outflow in S235AB making it likely that the SiO emission traces the primary molecular jet. Furthermore, as can be seen in Fig. 1, the SiO jet gas and water masers have similar terminal velocities, which suggests likely association, although this should be confirmed with mapping observation.

Molecular outflows are thought to be produced as linear momentum from a protostellar jet is transferred into ambient gas around to the protostar. This gas becomes entrained at a slower velocity and at larger radii from the inner jet. The fast SiO gas in S235AB-MIR therefore indicates the presence of an entraining jet, which is the driving source for the water masers and molecular outflows seen in this region.

Burns et al. (2015) report evidence of possible jet rotation in their maser observations. Our confirmation of a molecular jet in S235AB-MIR introduces the possibility of investigating the spatial orientation of the jet via mapping observations, which may unearth further evidence of jet rotation.

Our survey investigated the physical properties of numerous molecular species in S235AB, which revealed a cold, dense gas environment with high opacity in several tracers typical of the young cores in which MYSOs are embedded. Our survey results provide information on strengths and line widths of various gas tracers, which will be useful for further follow-up observations.

We detected maser emission in the CH$_3$OH (5–4) E transition at 84.521188 GHz emanating from an ensemble of blended features. With regard to thermal CH$_3$OH emission at 97 GHz, we measured a low rotation temperature for the methanol gas (~10 K) derived from the population diagram (Fig. 3). In this respect S235AB conforms to the trend reported by Minier & Booth (2002) who find low rotation temperatures to be a systematic feature among their sample of massive star forming regions. They postulate that the 97 GHz methanol gas may trace cooler gas at the outer parts of the methanol cloud.

With regard to molecular abundances, we compare our results to Gerner et al. (2014) who have investigated a sample of 59 massive star forming regions categorised into different stages of evolution which, in order of ascending evolutionary stage, are infrared dark clouds, high-mass protostellar objects, hot molecular cores, and ultra compact HII regions. A comparison with their results was possible for C$_2$H, SO, CS, and CH$_3$OH, which are common to both our work and theirs. For the cases of C$_2$H; SO, and CH$_3$OH our estimated abundances in S235AB are consistent with the hot molecular core category of Gerner et al. (2014; see their Fig. 3), while CS matches somewhere between the hot molecular core and high-mass protostellar object categories (see their Fig. 5). The abundance of H$^3$CN in S235AB is similar to those seen in another five well-studied massive star forming regions, the survey of which was reported in Zinchenko et al. (2009).

To conclude, the molecular abundances in S235AB are typical of a young massive star forming region. Its evolutionary stage close to that of a hot molecular core (this work and Felli et al. 2004) and preceding the formation of a HII region, as is indicated by the lack of centimetre emission (Ishii et al. 1995; Felli et al. 2006). Nonetheless, it is a very active region as can be seen by the presence of slow molecular outflows (Felli et al. 2004), a maser jet (Burns et al. 2015), and a fast molecular jet.

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