

# Gas-phase SO<sub>2</sub> in absorption towards massive protostars<sup>\*</sup>

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**Abstract.** We present the first detection of the  $\nu_3$  ro-vibrational band of gas-phase SO<sub>2</sub> in absorption in the mid-infrared spectral region around 7.3  $\mu\text{m}$  of a sample of deeply embedded massive protostars. Comparison with model spectra shows that the derived excitation temperatures correlate with previous C<sub>2</sub>H<sub>2</sub> and HCN studies, indicating that the same warm gas component is probed. The SO<sub>2</sub> column densities are similar along all lines of sight suggesting that the SO<sub>2</sub> formation has saturated, but not destroyed, and the absolute abundances of SO<sub>2</sub> are high ( $\sim 10^{-7}$ ). Both the high temperature and the high abundance of the detected SO<sub>2</sub> are not easily explained by standard hot core chemistry models. Likewise, indicators of shock induced chemistry are lacking.

**Key words.** star-formation: gas-phase molecules – ISM: abundances – ISM: molecules

## 1. Introduction

Observations with the Infrared Space Observatory (*ISO*) have dramatically increased our knowledge of the active chemistry occurring within star-forming regions. Extensive studies have revealed a vast richness of solid-state molecules embedded in icy grain mantles (Ehrenfreund & Schutte 2000) which highlight the crucial role of grain surface chemistry in molecule formation. Paralleling this, the infrared and submillimeter (Boonman et al. 2000; Lahuis & van Dishoeck 2000; van der Tak et al. 2000a) observations of gas-phase molecules directly probe the chemical and physical conditions of the star-formation process. By combining the solid-state and gas-phase observations, a detailed picture of the evolving chemistry emerges which can serve as a stringent test of proposed chemical models of star-forming regions.

Sulphur-bearing species are particularly interesting to study as they were originally proposed as tracers of shocks since the increased availability of OH radicals will lead to enhanced abundances of specific molecular species (Hartquist et al. 1980). The chemistry of sulphur-bearing molecules in warm gas is essentially governed by neutral-neutral reactions involving H<sub>2</sub>S formed on grain surfaces and subsequently evaporated into the gas-phase

(cf. Charnley 1997). The destruction of H<sub>2</sub>S frees atomic sulphur, which can then readily react with OH and O<sub>2</sub> to produce SO. SO<sub>2</sub> is easily formed through the conversion of SO by OH. Above  $\sim 200$ –300 K, the OH radicals are driven into H<sub>2</sub>O and the formation of SO<sub>2</sub> is halted. Except for the detection of the  $\nu_3$  band of gas-phase SO<sub>2</sub> in emission towards Orion (van Dishoeck et al. 1998), only purely rotational lines of SO<sub>2</sub> in the submillimeter have been detected toward massive star-forming regions. Abundances of  $\sim 10^{-9}$  (Schreyer et al. 1997) are typically derived which are much lower than model predictions (Charnley 1997). SO<sub>2</sub> abundances up to  $10^{-7}$  are only found in Orion-KL in the so-called plateau gas associated with the low-velocity outflow (e.g. Blake et al. 1987; Sutton et al. 1995). This gas is known to contain high abundances of OH (Melnick et al. 1987), and hence, the formation of SO<sub>2</sub> is intimately connected with the availability of reactive OH.

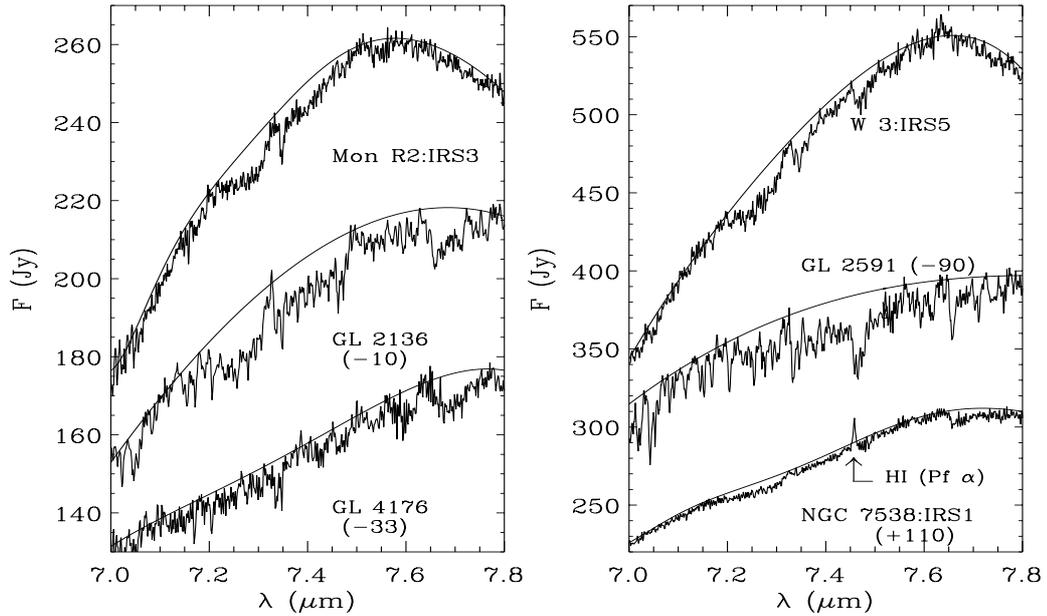
Here we present the first detection of infrared gas-phase SO<sub>2</sub> in absorption in the  $\nu_3$  ro-vibrational band towards a sample of embedded massive protostars.

## 2. Observations and reduction

High resolution AOT 6 ( $\lambda/\Delta\lambda \geq 1600$ ) grating mode observations of the massive protostars presented in this article were made with the Short Wavelength Spectrometer (SWS) on-board *ISO*. The  $\nu_3$  ro-vibrational mode of SO<sub>2</sub> lies within Band 2C which suffers from instrumental fringing of varying severity. In order to extract unimpeded SO<sub>2</sub>  $\nu_3$  absorption profiles, the fringes were corrected for by

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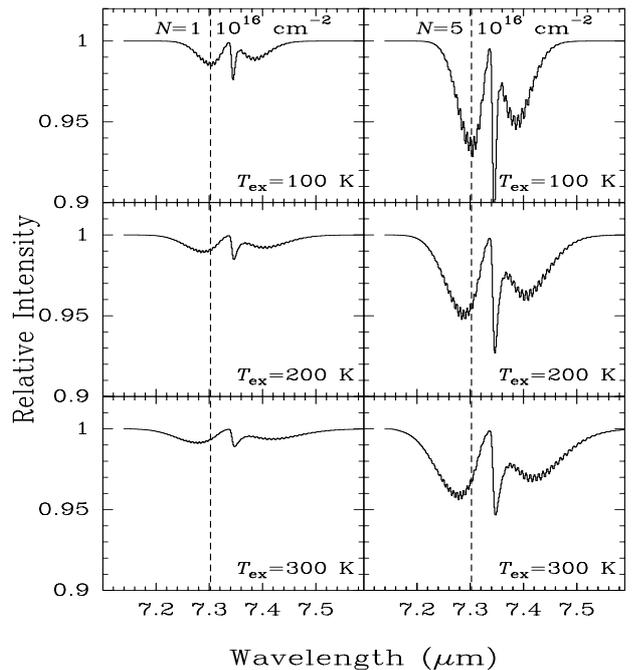


**Fig. 1.** ISO-SWS AOT 6 spectra towards six massive protostars. The thin solid lines indicate the locally defined 4th order polynomials adopted as the continua. Some of the sources were offset for clarity by a constant factor indicated in the brackets.

dividing the observed fluxes by cosine functions fitted to the data in wavenumber space (Lahuis & van Dishoeck 2000). The data were flat-fielded to the average level and then rebinned to the wavelength grid with a constant bin-size of  $0.003 \mu\text{m}$  which corresponds to  $\lambda/\Delta\lambda \sim 2500$ . The fully reduced 7–8  $\mu\text{m}$  spectra are shown in Fig. 1, where the noise level is approximately 4–5% for  $3\sigma$  significance.

### 3. Absorption features

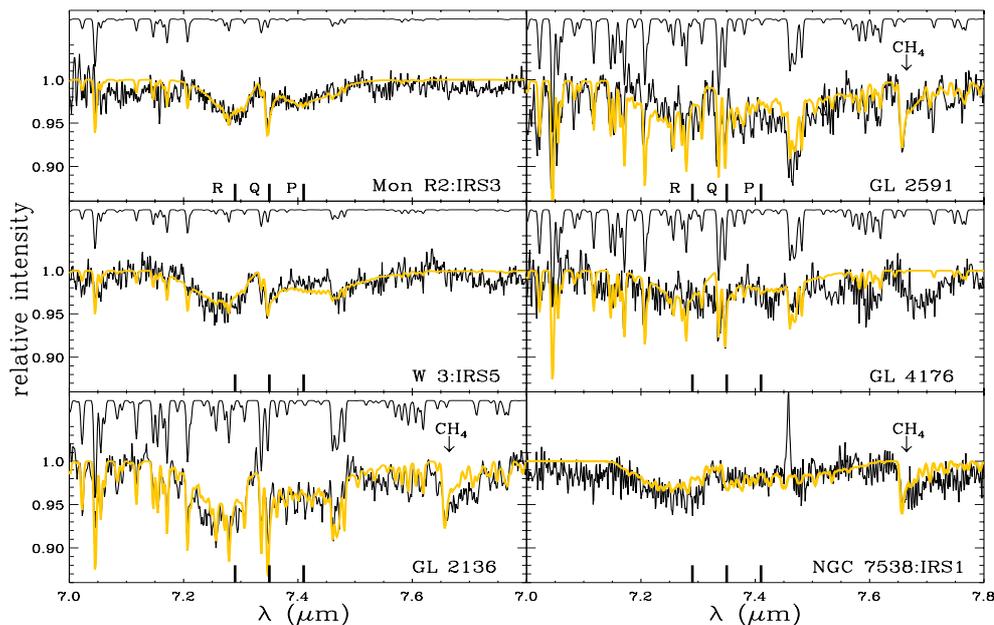
The 7–8  $\mu\text{m}$  spectra (Fig. 1), towards all lines of sight, display a richness of broad and narrow absorption features attributable to solid-state and gas-phase molecular species. The region is dominated by the red wing of the 6.85  $\mu\text{m}$  feature and the blue wing of the 10  $\mu\text{m}$  silicate feature (Keane et al. 2001). The feature near 7.6  $\mu\text{m}$  is well fitted by gas-phase and/or solid CH<sub>4</sub> (Boogert et al. 1997; Dartois et al. 1998). The spectra in Fig. 1 show evidence for weak features between 7.2  $\mu\text{m}$  and 7.4  $\mu\text{m}$ . The spectrum of Mon R2:IRS3 is particularly revealing in that it shows a narrow absorption feature at 7.342  $\mu\text{m}$  flanked by broader red- and blue-shifted bands. This structure is reminiscent of the *P*, *Q*, and *R* branch structure of gaseous molecules. The other sources show similar structure albeit less pronounced due to the presence of gas-phase H<sub>2</sub>O absorption lines (Boonman et al. 2000). A weak broad feature has been seen toward W 33A centered at 7.25  $\mu\text{m}$  and has been attributed to solid HCOOH (Keane et al. in prep.). However, this feature is easily distinguished from the spectral structure observed here as it is shifted to the blue and cannot explain the observed *Q*- and *P*-branch structure. We attribute the spectral structure between 7.2  $\mu\text{m}$  and 7.4  $\mu\text{m}$  to gas-phase SO<sub>2</sub>.



**Fig. 2.** Synthetic gas-phase SO<sub>2</sub> spectra calculated at various temperatures and column densities for a spectral resolution  $R \sim 2000$ . For all panels the Doppler parameter is  $3 \text{ km s}^{-1}$ . The dashed line shows the shift in the *R*-branch as a function of increasing excitation temperature.

### 4. Gas-phase SO<sub>2</sub>

The modeling of the spectra has been performed using synthetic spectra from Helmich (1996) combined with the molecular line data from the HITRAN 2000 database (<http://www.hitran.com>). Following the same analysis as in Lahuis & van Dishoeck (2000) and Boonman et al. (2000) a homogeneous source has been assumed with a



**Fig. 3.** The continuum divided spectra upon which the best fitting models (grey) are superimposed. Also shown are the H<sub>2</sub>O model spectra (offset) used for the modeling except in the case of NGC 7538:IRS1. The position of the gas-phase CH<sub>4</sub> band is indicated for the sources where it was included in the modeling along with the SO<sub>2</sub> Q, R, and P branches (thick solid lines).

single temperature  $T_{\text{ex}}$  and column density  $N$ . Since the SO<sub>2</sub> models are not sensitive to the linewidth, a Doppler  $b$  parameter of  $3 \text{ km s}^{-1}$  is adopted here, corresponding to the mean value of the submillimeter SO<sub>2</sub> lines. Figure 2 illustrates the expected spectral structure of the  $\nu_3$  ro-vibrational band of gas-phase SO<sub>2</sub> for different column densities and excitation temperatures. A global comparison with the observations shows that the observed features imply typically a column of a few times  $10^{16} \text{ cm}^{-2}$  of warm ( $\gtrsim 200 \text{ K}$ ) SO<sub>2</sub>.

Using these models, we have made detailed fits to the observed absorption features. Half of the sources show the presence of strong gas-phase H<sub>2</sub>O absorption in the  $\nu_2$  ro-vibrational band extending well into the 7.2–7.5  $\mu\text{m}$  region. Therefore the H<sub>2</sub>O model fits of Boonman et al. (2000) have been included in the modeling of gas-phase SO<sub>2</sub>. In the sources GL 2591, NGC 7538:IRS1, and GL 2136 gas-phase CH<sub>4</sub> is also present and this has been included in the models, although it affects the SO<sub>2</sub> band only moderately ( $\lesssim 2\%$ ). The best fitting models have been determined using the reduced  $\chi^2_{\nu}$ -method and are shown in Fig. 2. The corresponding excitation temperatures and column densities are listed in Table 1.

## 5. Discussion

Molecular abundances can serve as a direct means of probing the chemical history of star-formation. The derived SO<sub>2</sub> excitation temperatures range from 200–700 K and are in good agreement with those inferred for HCN and C<sub>2</sub>H<sub>2</sub> (Table 1), which are good tracers of warm gas (Lahuis & van Dishoeck 2000). The SO<sub>2</sub> column densities show little variation from source to source with typical

abundances of  $4\text{--}8 \times 10^{-7}$  relative to the total H<sub>2</sub>. The infrared SO<sub>2</sub> abundances are roughly consistent with the SO<sub>2</sub> abundances of  $\sim 10^{-7}$  observed in the submillimeter towards the Plateau, the Compact Ridge, and the Hot Core in Orion (Sutton et al. 1995). The relative constancy of the Orion SO<sub>2</sub> abundances is striking given the physical differences that exist between the afore mentioned regions in Orion. On the other hand, the derived SO<sub>2</sub> abundances in hot cores are much higher than SO<sub>2</sub> abundances in dark clouds (Irvine et al. 1983). More recently, Hatchell et al. (1998) have found gas-phase SO<sub>2</sub> abundances of  $5 \times 10^{-10}$  to  $2 \times 10^{-8}$  in hot core regions, which are a factor of  $\geq 10$  less than what is derived here. Some of this difference may well reflect the beam dilution suffered by the submillimeter observations. Thus, given the Orion template, there are two possible origins for the high abundance of gaseous SO<sub>2</sub>: hot core chemistry or shock induced chemistry.

In hot core chemistry, the SO<sub>2</sub> originates from oxidation of sulphur bearing species by OH (Charnley et al. 1997). This limits the SO<sub>2</sub> to gas with temperatures in the range  $\sim 100$  to  $\sim 200 \text{ K}$ . The observed SO<sub>2</sub> temperature is well above this, though it is possible that this high temperature reflects radiative pumping by the dust. Moreover, HCN and C<sub>2</sub>H<sub>2</sub> appear in gas with similar temperatures to that of the SO<sub>2</sub>. This is difficult to reconcile since the formation of SO<sub>2</sub> becomes very inefficient for temperature above  $\sim 230 \text{ K}$ , whereas the route to HCN is greatly enhanced (Charnley 1997; Boonman et al. 2001; Rodgers & Charnley 2001). Thus, SO<sub>2</sub> cannot be abundant in gas which has become enriched in HCN through the removal of OH. The SO<sub>2</sub> and HCN abundances must therefore peak at different radii from the protostar in order for standard hot core models to be compatible. Alternatively, the SO<sub>2</sub>

**Table 1.** SO<sub>2</sub> excitation temperatures and column densities in comparison with other gas-phase molecules.

Source	$T_{\text{ex}}$ (K)					$N$ ( $10^{16}$ cm <sup>-2</sup> )						$\frac{N(\text{SO}_2)}{N(\text{H}_2)^c}$ $10^{-7}$
	SO <sub>2</sub>	H <sub>2</sub> O <sup>a</sup>	CO <sub>hot</sub> <sup>b</sup>	HCN <sup>c</sup>	C <sub>2</sub> H <sub>2</sub> <sup>c</sup>	SO <sub>2</sub>	H <sub>2</sub> O <sup>a</sup>	CO <sub>total</sub> <sup>b</sup>	CO <sub>hot</sub> <sup>b</sup>	HCN <sup>c</sup>	C <sub>2</sub> H <sub>2</sub> <sup>c</sup>	
Mon R2:IRS3	225 <sup>+50</sup> <sub>-70</sub>	300	310 <sup>d</sup>	—	—	4 ± 0.8	60	980 <sup>d</sup>	444 <sup>d</sup>	—	—	8.2
W 3:IRS5	450 <sup>+100</sup> <sub>-100</sub>	400	577	400	500	5 ± 0.8	40	2580	1260	0.5	0.3	3.8
GL 2136	350 <sup>+50</sup> <sub>-100</sub>	500	580	600	800	6 ± 0.8	150	2200	1500	3.5	1.5	5.5
GL 2591	750 <sup>+70</sup> <sub>-100</sub>	450	1010	600	900	6 ± 0.4	350	1280	558	4	2	6.3
GL 4176	350 <sup>+175</sup> <sub>-75</sub>	400	≥500 <sup>c</sup>	500	700	4 ± 1.0	150	1600 <sup>c</sup>	800 <sup>c</sup>	2	1	5.0
NGC 7538:IRS1	700 <sup>+300</sup> <sub>-400</sub>	176 <sup>b</sup>	176	600	800	4 ± 1.0	<20	1740	840	1	0.8	4.7

<sup>a</sup> Boonman et al. (2000) unless otherwise noted; <sup>b</sup> Mitchell et al. (1990) unless otherwise noted, using <sup>13</sup>CO and assuming <sup>12</sup>CO/<sup>13</sup>CO = 60; <sup>c</sup> Lahuis & van Dishoeck (2000); <sup>d</sup> Giannakopoulou et al. (1997).

may be formed on grains surfaces and then be released into the gas by evaporation. Grain surface chemistry models predict an abundance of  $\sim 3 \times 10^{-3}$  relative to H<sub>2</sub>O on the ice (Tielens & Hagen 1982; Tielens private communication), which is a factor  $\gtrsim 10$  less than what is derived here. Another aspect is that the SO<sub>2</sub> column density does not vary between the sources, whereas H<sub>2</sub>O shows large variation. The constancy of the SO<sub>2</sub> column density can be explained by the fact that the SO<sub>2</sub> is only abundant in a narrow zone between  $\sim 90$  K (the ice evaporation temperature) and  $\sim 230$ – $300$  K (the OH  $\rightarrow$  H<sub>2</sub>O transition), whose mass does not vary much in spite of the different total masses of the sources (Doty et al., in prep.).

The presence of SO and SO<sub>2</sub> has often been quoted as evidence for shocks (Hartquist et al. 1980). The degree to which the SO<sub>2</sub> abundance is enhanced depends on whether most of the sulphur is initially in atomic form ( $\sim 10^{-7}$ , Pineau des Forêts et al. 1993) or locked up in stable molecules ( $\sim 10^{-8}$ , Leen & Graff 1988). A good aspect of the shock induced chemistry hypothesis is that the SO<sub>2</sub> and HCN may be colocated in gas which contains freshly (i.e.,  $\lesssim 3.4 \times 10^4$  yr) released grain mantle molecules. However, the lack of increased SO<sub>2</sub> line widths at submillimeter wavelengths would seem to indicate the decay of shock activity within the region sampled by the submillimeter observations. In addition, the presence of fragile molecules sensitive to destruction by shocks (e.g. H<sub>2</sub>CO; van der Tak et al. 2000b) makes shock-induced chemistry less likely as the source of SO<sub>2</sub> for these sources.

In general, the gaseous SO<sub>2</sub>/H<sub>2</sub>S ratio serves as a sensitive chemical clock for the star formation process and searches for these molecules at high spectral resolution are needed to help resolve the issue of the origin of the gas-phase SO<sub>2</sub>.

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