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Modeling accretion shocks at the disk-envelope interface

Sulfur chemistry

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ABSTRACT

Context. As material from an infalling protostellar envelope hits the forming disk, an accretion shock may develop which could (partially) alter the envelope material entering the disk. Observations with the Atacama Large Millimeter/submillimeter Array indicate that emission originating from warm SO and SO₂ might be good tracers of such accretion shocks.

Aims. The goal of this work is to test under what shock conditions the abundances of gas-phase SO and SO₂ increase in an accretion shock at the disk-envelope interface.

Methods. Detailed shock models including gas dynamics are computed using the Paris-Durham shock code for non-magnetized *J*-type accretion shocks in typical inner envelope conditions. The effect of pre-shock density, shock velocity, and strength of the ultraviolet radiation field on the abundance of warm SO and SO₂ is explored. Compared with outflows, these shocks involve higher densities ($\sim 10^7 \text{ cm}^{-3}$), lower shock velocities ($\sim \text{few km s}^{-1}$), and large dust grains ($\sim 0.2 \mu\text{m}$) and thus probe a different parameter space.

Results. Warm gas-phase chemistry is efficient in forming SO under most *J*-type shock conditions considered. In lower-velocity ($\sim 3 \text{ km s}^{-1}$) shocks, the abundance of SO is increased through subsequent reactions starting from thermally desorbed CH₄ toward H₂CO and finally SO. In higher velocity ($\geq 4 \text{ km s}^{-1}$) shocks, both SO and SO₂ are formed through reactions of OH and atomic S. The strength of the ultraviolet radiation field is crucial for SO and in particular SO₂ formation through the photodissociation of H₂O. Thermal desorption of SO and SO₂ ice is only relevant in high-velocity ($\geq 5 \text{ km s}^{-1}$) shocks at high densities ($\geq 10^7 \text{ cm}^{-3}$). Both the composition in the gas phase, in particular the abundances of atomic S and O, and in ices such as H₂S, CH₄, SO, and SO₂ play a key role in the abundances of SO and SO₂ that are reached in the shock.

Conclusions. Warm emission from SO and SO₂ is a possible tracer of accretion shocks at the disk-envelope interface as long as a local ultraviolet field is present. Observations with the Atacama Large Millimeter/submillimeter Array at high-angular resolution could provide further constraints given that other key species for the gas-phase formation of SO and SO₂, H₂S and H₂CO, are also covered. Moreover, the *James Webb* Space Telescope will give access to other possible slow, dense shock tracers such as H₂, H₂O, and [S I] 25 μm .

Key words. Astrochemistry – shock waves – stars: formation – stars: protostars – stars: low-mass – ISM: abundances

1. Introduction

It is currently still unknown how much reprocessing of material occurs during the collapse from a prestellar core to a protostar and disk. Two scenarios can be considered: the inheritance scenario where the chemical composition is conserved from cloud to disk, or the (partial) reset scenario where the envelope material is modified during its trajectory from envelope to disk. Some inheritance is suggested by the rough similarity between cometary and interstellar ices (Mumma & Charnley 2011; Drozdovskaya et al. 2018, 2019). On the other hand, some modification of envelope material is expected due to the increase of temperature and stellar ultraviolet radiation during infall (e.g., Aikawa et al. 1999; Visser et al. 2009). Moreover, as the infalling envelope hits the disk, an accretion shock may develop which will alter the composition of the material flowing into the disk, and therefore (partially) reset the chemistry. In the most extreme case, reset implies complete vaporization of all molecules back to atoms with sub-

sequent reformation. Milder versions of reset include sputtering of ices, and gas and ice chemistry modifying abundances.

Already in the earliest Class 0 phase of the low-mass star-formation process, an accretion disk around the young protostar is formed (e.g., Tobin et al. 2012; Murillo et al. 2013). The impact of infalling envelope material onto the disk can cause a shock, raising temperatures of gas and dust at the disk-envelope interface to values much higher than from heating by stellar photons alone (Draine et al. 1983; Neufeld & Hollenbach 1994). These low-velocity ($\sim \text{few km s}^{-1}$) accretion shocks at high densities ($\sim 10^7 \text{ cm}^{-3}$) are widely found in models and simulations (e.g., Cassen & Moosman 1981; Li et al. 2013; Miura et al. 2017), and often invoked in Solar System formation to explain phenomena like noble gas trapping (Owen et al. 1992). They are most powerful in the earliest stages when the disk is still small ($\lesssim 10 \text{ AU}$). However, that infalling material may eventually end up in the protostar and not in the disk. As the embedded disk

reaches a size of (several) tens of AU, the envelope material entering the disk will remain there and eventually form the building blocks of planets (e.g., Harsono et al. 2018; Manara et al. 2018; Tychoniec et al. 2020).

These accretion shocks have not yet been undisputedly detected. Typical optical shock tracers such as [O I] 6300 Å and [S II] 6371 Å are more sensitive to high-velocity shocks ($\geq 20 \text{ km s}^{-1}$, e.g., Podio et al. 2011; Banzatti et al. 2019) and difficult to detect in protostellar environments due to the high extinction by the surrounding envelope. Observations of mid-infrared shock tracers such as H₂O, high-J CO, [S I] 25 μm , and [O I] 63 μm suffer from outflow contamination in low-spatial resolution space-based observations (e.g., Kristensen et al. 2012; Nisini et al. 2015; Rivière-Marichalar et al. 2016) and from the Earth's atmosphere for ground-based observations; here the launch of the *James Webb* Space Telescope will provide a solution. At sub-mm wavelengths, classical diagnostics of shocks include sputtering or grain-destruction products such as CH₃OH and SiO (e.g., Caselli et al. 1997; Schilke et al. 1997; Gusdorf et al. 2008a,b; Guillet et al. 2009; Suutarinen et al. 2014). However, usually these species are only observed in relatively high-velocity outflows and jets through broad emission lines (e.g., Tychoniec et al. 2019; Taquet et al. 2020; Codella et al. 2020).

Sulfur bearing species such as SO and SO₂ have been suggested as possible accretion shock tracers. An enhancement of warm SO emission at the centrifugal barrier, the interface between envelope and disk, has been observed in the L1527 Class 0/I system with the Atacama Large Millimeter/submillimeter Array at $\sim 100 \text{ AU}$ scales (Sakai et al. 2014, 2017). A similar rotating structure in SO has been seen toward the Class I system Elias 29 (Oya et al. 2019). Equivalently, warm SO₂ emission possibly related to an accretion shock was observed toward the B335 Class 0 protostar (Bjerkeli et al. 2019) and a few Class I sources (Oya et al. 2019; Artur de la Villarmois et al. 2019). However, both SO and SO₂ are also seen related to outflow activity, both in large-scale outflows (e.g., Codella et al. 2014; Taquet et al. 2020) as in disk winds on smaller scales (e.g., Tabone et al. 2017; Lee et al. 2018). Moreover, high-angular resolution observations ($\sim 30 \text{ AU}$) of the Class I source TMC1A show narrow SO emission lines coming from a ring-shaped morphology which may be linked to the warm inner envelope (Harsono et al. 2021). Emission from warm SO and SO₂ is thus not an unambiguous tracer of accretion shocks; comparison to shock models is necessary to make robust conclusions.

In shocks, the gas temperature readily rises to $> 100 \text{ K}$ (Draine et al. 1983; Flower & Pineau des Forêts 2003; Godard et al. 2019), enough to ignite gas-phase formation of SO and SO₂ (e.g., Prasad & Huntress 1980; Hartquist et al. 1980). A key species for this is the OH radical, which is efficiently produced in shocks through the endothermic reaction between H₂ and atomic oxygen, but also through photodissociation of H₂O. The OH radical reacts with atomic sulfur to form SO and subsequently to form SO₂. Moreover, SO can also be formed from the SH radical reaction with atomic oxygen (Hartquist et al. 1980). Since the gas-phase chemistry of SO and SO₂ in shocks is dependent on radicals such as OH and SH, the abundance of these species is not solely determined through the temperature in the shock but also through the strength of the local ultraviolet field.

Alternatively, thermal desorption of SO and SO₂ ice in shocks can enhance the gas-phase abundances of SO and SO₂. In interstellar ices, OCS is the only securely identified sulfur-bearing species (Geballe et al. 1985; Palumbo et al. 1995), with a tentative detection for SO₂ (Boogert et al. 1997; Zasowski et al.

2009). H₂S is suggested as the main sulfur carrier in the ices (e.g., Vidal et al. 2017), but so far only upper limits could be derived (Jiménez-Escobar & Muñoz Caro 2011). To date, SO ice has not yet been detected. All aforementioned species are observed in cometary ices such as 67P/Churyumov-Gerasimenko (Calmonte et al. 2016; Rubin et al. 2019; Altwegg et al. 2019). Miura et al. (2017) modeled low-velocity (2 km s^{-1}) accretion shocks taking into account the dust dynamics but without any gas-phase chemistry included and could only explain the observations of Sakai et al. (2014) if SO was thermally desorbed while adopting dust grains that were smaller than $0.1 \mu\text{m}$. Sputtering seems to be irrelevant as this becomes only efficient at larger shock velocities ($> 10 \text{ km s}^{-1}$; e.g., Aota et al. 2015). However, to thermally desorb sulfur-bearing ices such as SO and SO₂ from the grains, the dust temperature has to reach values on the order of 40–70 K.

In this paper, we present a grid of irradiated low-velocity ($< 10 \text{ km s}^{-1}$) *J*-type accretion shock models at typical inner envelope densities ($10^5 - 10^8 \text{ cm}^{-3}$). This is a different parameter space than usually explored in, for example, higher-velocity ($> 10 \text{ km s}^{-1}$) outflows at lower densities ($\sim 10^4 \text{ cm}^{-3}$). The goal is to test under what shock conditions the abundance of warm SO and SO₂ is significantly increased. In Sect. 2, the model and input parameters are introduced. We present the results of our analysis in Sect. 3, after which we discuss these results in Sect. 4. Our conclusions are summarized in Sect. 5.

2. Accretion shock model

2.1. Shock model

The accretion shock at the disk-envelope interface is modeled using a slightly modified version of the Paris-Durham shock code (Flower & Pineau des Forêts 2003; Lesaffre et al. 2013; Godard et al. 2019). This publicly available numerical tool¹ computes the dynamical, thermal, and chemical structure of stationary plane parallel shock waves. The models therefore represent a continuous inflow of envelope material onto the disk and thus a continuous sequence of shocks. The location of the accretion shock is expected to move outward as the disk grows and evolves (Visser & Dullemond 2010).

In this paper, only stationary non-magnetized *J*-type accretion shocks are presented. The presence of a magnetic field could result in *C* or *CJ*-type shocks (e.g., Draine 1980; Flower & Pineau des Forêts 2003), see Appendix E for details. However, the length of *C*-type accretion shocks is strongly dependent on the initial conditions and the size of dust grains and may reach envelope scales of $\sim 1000 \text{ AU}$, which is not consistent with the emission of SO and SO₂ seen on $< 100 \text{ AU}$ scales (e.g., Sakai et al. 2014, 2017; Artur de la Villarmois et al. 2019; Oya et al. 2019). *CJ*-type shocks could be relevant in intermediate magnetized inner envelopes ($\sim 0.1 \text{ mG}$; Hull et al. 2017, Appendix E), but to compute *CJ*-type shocks across our specific parameter space (low velocities, high densities), the Paris-Durham code needs to be tuned, which is beyond the scope of this work. Nevertheless, the results of *CJ* and *C* type shocks are not expected to be different from *J*-type shocks in terms of chemistry.

In the following sections, the main physical and chemical quantities and properties that are included in the model are introduced. In particular, the most relevant physical and chemical processes and updates on the version of the code presented by Godard et al. (2019) are highlighted.

¹ <https://ism.obspm.fr>

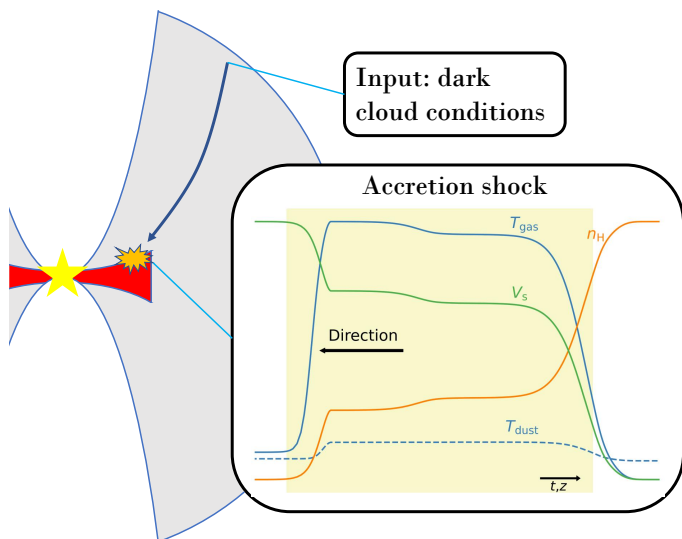


Fig. 1. Physical structure of a protostellar envelope. Starting with dark cloud conditions, an accretion shock model is calculated. The shock profile in the bottom panel is shown on a logarithmic scale, with the yellow region indicating the shocked region.

2.1.1. Geometry

The shock is calculated following a stationary plane parallel structure, see Fig. 1. A shock front propagates with a velocity V_s in the direction of negative distance z . The entire structure is irradiated with an isotropic radiation field equal to the isotropic standard interstellar radiation field scaled with a factor G_0 (Mathis et al. 1983), as was implemented by Godard et al. (2019). In protostellar systems, the ultraviolet radiation can originate both from the accretion onto the protostar itself or from shocks in the bipolar jets (e.g., van Kempen et al. 2009; Yıldız et al. 2012; Benz et al. 2016). Moreover, the extinction A_V between the source of ultraviolet irradiation and the shock at the disk-envelope interface can be as high as ~ 10 mag (e.g., Drozdovskaya et al. 2015) and is assumed to be taken into account with the choice of G_0 . Therefore, the exact source of ultraviolet radiation is irrelevant here as long as a reasonable range in G_0 is covered and since the radiation field is assumed to be isotropic. A diluted black body of 100 K representing far-infrared dust emission from the accretion disk is added to the radiation field. Assuming an emitting region of 4.5 AU at a distance of 100 AU, the black body is diluted with a factor $W = 5 \times 10^{-4}$ (for details, see Appendix A of Tabone et al. 2020), and is constant for all models. Additionally, secondary ultraviolet photons created through the collisional excitation of H_2 by electrons produced by cosmic ray ionization are included.

2.1.2. Radiative transfer and photon processes

Photodissociation and photoionization processes of both the gas and dust are taken into account. The cross sections are taken from the Leiden database (Heays et al. 2017). To save computation time, the photoreaction cross sections were only used for the photodissociation of CH, CH_3 , CH_4 , NH, CN, O_2 , OH, H_2O , H_2CO , SH, H_2S , CS, OCS, SO, and SO_2 , and the photoionization of C, S, CH, CH_3 , CH_4 , O_2 , OH, H_2O , H_2CO , SH, H_2S , and SO. Photodissociation and photoionization of all other species are also included in the chemistry, but using the analytical expression of Heays et al. (2017). Due to the self-shielding properties of H_2 , CO, and N_2 , photodissociation of these species is as-

sumed to be negligible. The destruction of atoms and molecules by far-ultraviolet photons between 911 Å and 2400 Å are calculated including both continuum and discrete processes.

2.1.3. Grains and PAHs

The grains are assumed to have grown in protostellar envelopes to larger sizes than the typical interstellar medium distribution (e.g., Guillet et al. 2007; Miotello et al. 2014; Harsono et al. 2018; Galametz et al. 2019). Here, we assume that all grains have an initial core size of 0.2 μm (Guillet et al. 2020), and are either neutral, or single negatively or positively charged. Moreover, the dust grains are assumed to be dynamically coupled to the gas. Grain-grain interactions are not included since vaporization and shattering of dust grains are only relevant for *J*-type shocks moving at high velocities (> 20 km s^{-1} ; Guillet et al. 2009). Dust coagulation could potentially be relevant in changing the grain size distribution (Guillet et al. 2011, 2020), but is not included here.

Other large species are polycyclic aromatic hydrocarbons (PAHs), which are modeled here as single particles with a size of 6.4 Å (Flower & Pineau des Forêts 2003). In most studies of interstellar shocks, the abundance of polycyclic aromatic hydrocarbons is set to the typical value of the interstellar medium ($\sim 10^{-6}$ with respect to the proton density $n_H = 2n(H_2) + n(H) + n(H^+)$; Draine & Li 2007). However, given that polycyclic aromatic hydrocarbons freeze out in dense clouds, the abundance of polycyclic aromatic hydrocarbons in the gas is set to $\sim 10^{-8}$ with respect to n_H (Geers et al. 2009).

2.1.4. Chemical network

The chemical network adopted in this work consists of 143 species which can take place in about a thousand reactions, both in the gas phase and in the solid state. The network is largely similar to that used by Flower & Pineau des Forêts (2015) and Godard et al. (2019), with the addition of important sulfur bearing ices such as SO and SO_2 and their desorption and adsorption reactions. Moreover, the binding energies of all ice species were updated (e.g., 1800 K, 3010 K, and 2290 K for SO, SO_2 , and H_2S , respectively; Penteado et al. 2017). For a density of 10^8 cm^{-3} , this results in a sublimation temperature of 37 K, 62 K, and 47 K for SO, SO_2 , and H_2S , respectively. For simplicity, grain surface reactions have been disabled since the effect of these reactions on timescales of the shock is negligible. Sputtering of ices is not relevant here since in *J*-type shocks all material is coupled in one fluid.

2.1.5. Heating and cooling

The thermal balance is calculated consistently throughout the shock. The main sources of gas heating in non-magnetized *J*-type shocks are through viscous stresses and compression of the medium at the shock front where T_{gas} increases adiabatically over a few mean free paths. A second form of heating occurs through irradiation coming from the incident local ultraviolet radiation field and through gas-dust thermal coupling.

The gas is cooled through atomic lines of C, N, O, S, and Si, ionic lines of C^+ , N^+ , O^+ , S^+ , and Si^+ , and rotational and rovibrational lines of H_2 , OH, H_2O , NH_3 , ^{12}CO , and ^{13}CO (Flower & Pineau des Forêts 2003; Lesaffre et al. 2013; Godard et al. 2019). All cooling by atomic and ionic lines, as well as rovibrational lines of H_2 are calculated in the optically thin limit. The cool-

Table 1. Fiducial model parameters and explored parameter range

	Units	Fiducial	Range
n_{H}	cm^{-3}	10^7	$10^5 - 10^8$
ζ_{H_2}	s^{-1}	10^{-17}	–
G_0	Mathis	1	$10^{-3} - 10^2$
V_s	km s^{-1}	3	1 – 10

ing of OH is calculated using an analytical cooling function (Le Bourlot et al. 2002), which holds up to densities of 10^{10} cm^{-3} . However, for NH_3 , local thermodynamic equilibrium effects become relevant already for densities $\gtrsim 10^8 \text{ cm}^{-3}$. The cooling through pure rotational lines of NH_3 has therefore been recalculated numerically using the RADEX software (van der Tak et al. 2007) in the optically thin limit, see Appendix A. Cooling by rovibrational lines of H_2O , ^{12}CO , and ^{13}CO is computed using the tabulated values of Neufeld & Kaufman (1993) which take into account opacity effects. Additionally, the gas can be cooled through collisions with dust grains which subsequently radiate away the heat.

The temperature of the dust is calculated from the equilibrium between heating through absorption of photons, thermal collisions with the gas which can both heat and cool the dust, and cooling through infrared emission (Godard et al. 2019). The far-infrared radiation field of the 100 K black body is of particular importance for the heating for dust grains in the pre and post-shock regimes as it sets a minimum dust temperature of $\sim 22 \text{ K}$. A dust temperature of $< 20 \text{ K}$ would imply that even the most volatile species like CO and N_2 are frozen out onto dust grains. However, observations of embedded protostellar systems suggest that CO and N_2 are not frozen out in the inner regions of such systems (e.g., van ’t Hoff et al. 2018). A higher dust temperature would result in more ice species being desorbed prior to the shock, but the effect on the shock structure and chemistry is negligible.

2.2. Input parameters

The range of physical parameters explored in this paper is presented in Table 1. Important physical parameters are the initial proton density n_{H} , the cosmic ray ionization rate of H_2 ζ_{H_2} , the strength of the local incident radiation field, and the shock velocity V_s . Depending on the mass of the source and size of the disk, the density of infalling envelope material impacting the disk can be $\sim 10^5 - 10^8 \text{ cm}^{-3}$ (e.g., Harsono et al. 2015). Therefore, the investigated range of n_{H} covers $\sim 10^5 - 10^8 \text{ cm}^{-3}$, with the fiducial value at 10^7 cm^{-3} . The radiation field is parameterized as a combination of the isotropic standard interstellar radiation field multiplied with a factor G_0 (Mathis et al. 1983) and infrared emission from the disk (which is constant for all models). The strength of the local ultraviolet radiation field at the disk-envelope interface is highly uncertain and (among other things) dependent on the distance to the source of ultraviolet radiation (see Sect 2.1.1). Without extinction, the local G_0 value could be as high as 10^4 at $\sim 100 \text{ AU}$ (Visser et al. 2012). However, taking into account the extinction the local strength will likely not be much higher than a G_0 of ~ 100 . Therefore, a large range in G_0 is considered: $10^{-3} - 10^2$, with a fiducial value of $G_0 = 1$. The fiducial V_s is set to 3 km s^{-1} , which is roughly equal to the velocity of the infalling envelope at 100 AU for a $0.5 M_{\odot}$ star. Shock velocities of $1 - 10 \text{ km s}^{-1}$ are considered, which correspond to an infalling envelope at $10 - 1000 \text{ AU}$. The initial gas temperature is set to 25 K .

Input abundances of smaller molecules and atoms are set to match typical low-mass dark cloud conditions (van der Tak et al. 2003; Boogert et al. 2015; Navarro-Almaida et al. 2020; Tafalla et al. 2021; Goicoechea & Cuadrado 2021). The initial gas-phase abundance of atomic S and O is set to 10^{-6} with respect to n_{H} , with most of the oxygen and carbon reservoirs locked up in refractory grain cores or ices such as H_2O and CH_4 (10^{-4} and 10^{-6} , respectively; Boogert et al. 2015). H_2S ice is assumed to be the dominant sulfur carrier with an abundance of $\sim 2 \times 10^{-5}$ (e.g., Vidal et al. 2017; Navarro-Almaida et al. 2020). The abundance of both SO and SO_2 ice was set to 10^{-7} (e.g., Boogert et al. 2015; Rubin et al. 2019; Altwegg et al. 2019), with the initial gas-phase abundance at 10^{-9} (van der Tak et al. 2003). The full list of input abundances is presented in Appendix B.

3. Results

3.1. Temperature and density

The gas and dust temperature, T_{gas} and T_{dust} , respectively, and density structure of the fiducial shock model are shown in the left of Fig. 2. At the start of the shock, T_{gas} increases to $\sim 500 \text{ K}$, which is in agreement with the analytic expression for a weakly magnetized and fully molecular shock derived by Lesaffre et al. (2013) from the Rankine-Hugoniot relations,

$$T_{\text{gas,max}} \approx 53V_s^2, \quad (1)$$

where V_s is the initial shock velocity in units of km s^{-1} . Following the jump in temperature, the gas is cooled down through radiation of rotational and rovibrational molecular lines and atomic lines. The most dominant coolant right after the shock front is H_2 , while cooling by (optically thick) pure rotational and rovibrational lines of CO and H_2O take over as the temperature drops below $\lesssim 400 \text{ K}$. Cooling of the gas through gas-dust thermal coupling is negligible. Cooling through NH_3 in the optically thin limit is significant for those shocks with $V_s \gtrsim 6 \text{ km s}^{-1}$ since in those shocks the abundance of NH_3 is significantly increased through high-temperature gas-phase chemistry. The end of the shock is defined as the distance where T_{gas} drops below 50 K , which corresponds to $\sim 3 \text{ AU}$ for the fiducial model. In the post-shock, the thermal balance between gas and dust is determined by heating of the dust through the far-infrared radiation field, gas-grain collisions that transfer the heat from dust to gas, and cooling of the gas through predominantly CO rotational transitions.

The dust temperature is very relevant for these shocks since it increases to values where thermal desorption of ices occurs. In the shock, the material compresses, increasing the density by up to two orders of magnitude (i.e., to about 10^9 cm^{-3} in Fig. 2). Due to the thermal coupling between the gas and dust, T_{dust} increases to $\sim 30 \text{ K}$ for the fiducial model, enough for thermal sublimation of volatile species such as CH_4 , but not for more strongly bound ices such as H_2S , H_2O , and CH_3OH . However, as shown in Fig. 3, T_{dust} can become as large as $> 70 \text{ K}$ in the highest-velocity ($\sim 10 \text{ km s}^{-1}$) shocks in the densest (10^8 cm^{-3}) media, resulting in thermal sublimation of several ices including H_2S , SO, and SO_2 . Dust heating through ultraviolet radiation is only significant for strongest irradiated environments with $G_0 \gtrsim 50$ and only up to $T_{\text{dust}} \lesssim 25 \text{ K}$. Since photodesorption of ices is negligible on timescales of the shock (see Eq. (C.2)), thermal desorption is the dominant mechanism in releasing ices to the gas phase.

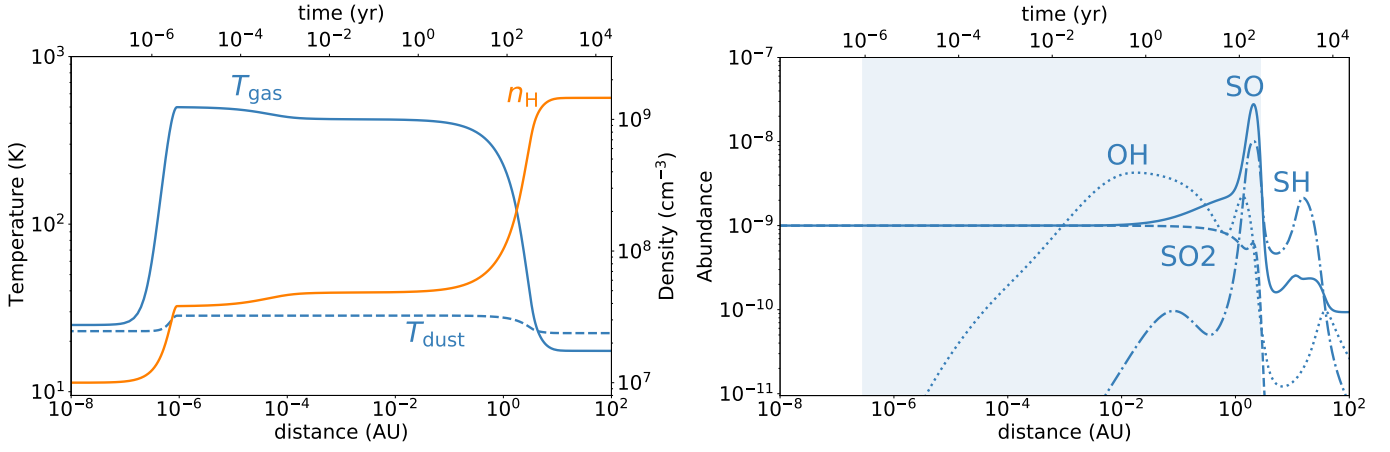


Fig. 2. *Left:* Temperature and density structure in the fiducial *J*-type shock. The shock starts at 0 AU and ends at about 5 AU. The dust temperature does not increase enough for thermal sublimation of SO and SO₂. *Right:* Abundance profiles of SO (solid) and SO₂ (dashed) and the key species for gas-phase formation of these species: SH (dot-dashed) and OH (dotted). The region where $T_{\text{gas}} > 50$ K is indicated with the shaded blue region.

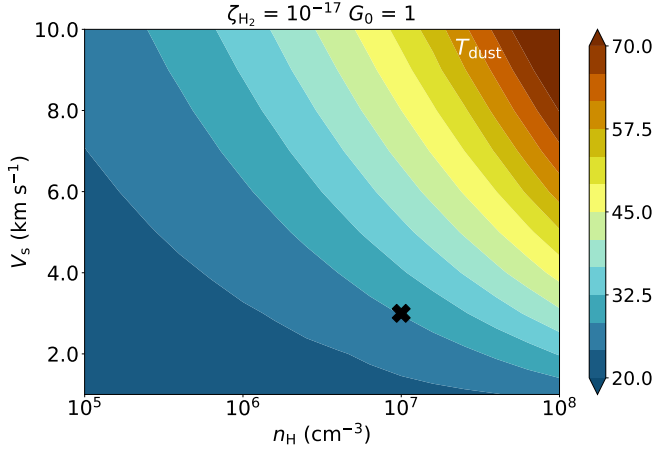


Fig. 3. Maximum dust temperature reached (in color) in shock models as function of initial n_{H} and V_{s} . All other physical parameters are kept constant to the fiducial values and listed on top of the figure. The black cross indicates the position of the fiducial model.

3.2. Chemistry of SO and SO₂

The abundance profiles of SO and SO₂ in the fiducial model are presented in the right part of Fig. 2. Toward the end of the shock, as the gas and dust are cooling down, a significant increase in the abundances of SO is visible up to the $\sim 5 \times 10^{-8}$ level. The abundance of SO₂ does not increase above the dark cloud abundance of 10^{-9} (van der Tak et al. 2003). The maximum abundance reached in the shock for SO and SO₂ as function of different initial n_{H} and V_{s} is presented in Fig. 4. In the following subsections different parts of the parameter space are highlighted using the relevant chemical formation pathways in Fig. 5.

3.2.1. Low-velocity shocks: ~ 3 km s⁻¹

In low-velocity (~ 3 km s⁻¹) shocks, thermal desorption of SO and SO₂ ice does not occur since T_{dust} only rises to $\lesssim 35$ K. Any increase in abundance thus originates from gas-phase chemistry. Here, the relevant chemical reaction leading to the formation of SO is,



The main formation pathway here of SH is via the green route of Fig. 5, that is, through the reaction of H₂CO with S⁺. Since H₂CO only desorbs directly from the ice when $T_{\text{dust}} \gtrsim 65$ K, H₂CO is formed through the gas-phase reaction of the CH₃ radical with atomic O. To get a significant amount of CH₃ in the gas, CH₄ ice needs to desorb (at $T_{\text{dust}} \gtrsim 25$ K; Penteado et al. 2017) and subsequently photodissociate into CH₃ and atomic H. The S⁺ ion originates from the photo-ionization of atomic S. The strongest increase in abundance through this route at intermediate densities ($n_{\text{H}} \sim 10^7$ cm⁻³). It is important to note that H₂CO is a representative here of any hydrocarbon reacting with S⁺ to form SH.

Some SO₂ is also formed in low-velocity shocks through a reaction of SO with OH (see Reaction (4)). However, the abundance of SO₂ does not increase above the initial dark cloud abundance of 10^{-9} because no significant amount of OH is available at these shock velocities.

3.2.2. High-velocity shocks: > 4 km s⁻¹

For shocks propagating at higher velocities ($\gtrsim 4$ km s⁻¹), both SO and SO₂ are efficiently formed through gas-phase chemistry, see Fig. 4. After the start of the shock, H₂O is readily formed through reactions of OH with H₂ (Flower & Pineau des Forêts 2010, see also Fig. D.1). As the shock cools down to below $\lesssim 300$ K, the production of H₂O stops, but OH is still being formed through photodissociation of H₂O. This increases the OH abundance in the gas which stimulates the formation of SO and SO₂ through the red route of Fig. 5,



The highest abundances are achieved in less dense environments ($\lesssim 10^6$ cm⁻³), with the maximum abundance of both SO and SO₂ dropping when moving toward intermediate densities ($\sim 10^6 - 10^7$ cm⁻³). In the latter conditions, NH₃ becomes a dominant coolant in the tail of the shock. Therefore, the shock cools down more quickly and hence the region of favorable conditions for SO and SO₂ formation ($100 \lesssim T_{\text{gas}} \lesssim 300$ K) is reduced and their maximum abundances are lower.

At the highest densities ($n_{\text{H}} > 10^7$ cm⁻³), thermal desorption of sulfur-bearing ices becomes relevant for the chemistry (i.e., the blue route in Fig. 5). SO ice thermally sublimates as the dust

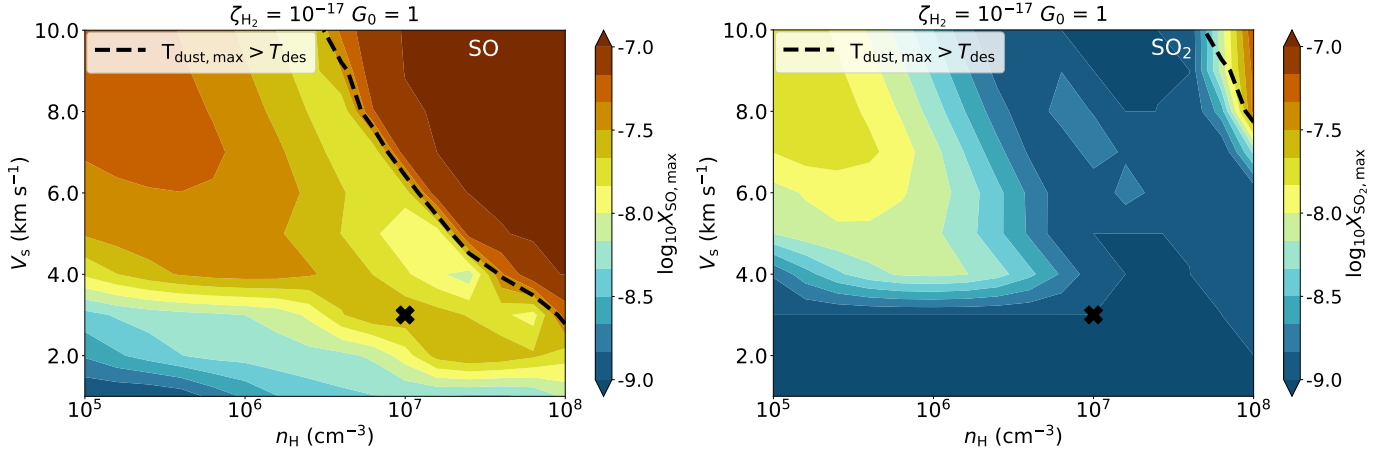


Fig. 4. Maximum abundance reached (in color) of SO (left) and SO₂ (right) in shock models as function of initial n_H and V_s . All other physical parameters are kept constant to the fiducial values and listed on top of the figure. The black cross indicates the position of the fiducial model. The dashed black line shows the ice line, i.e., where 50% of the ice is thermally desorbed into the gas in the shock.

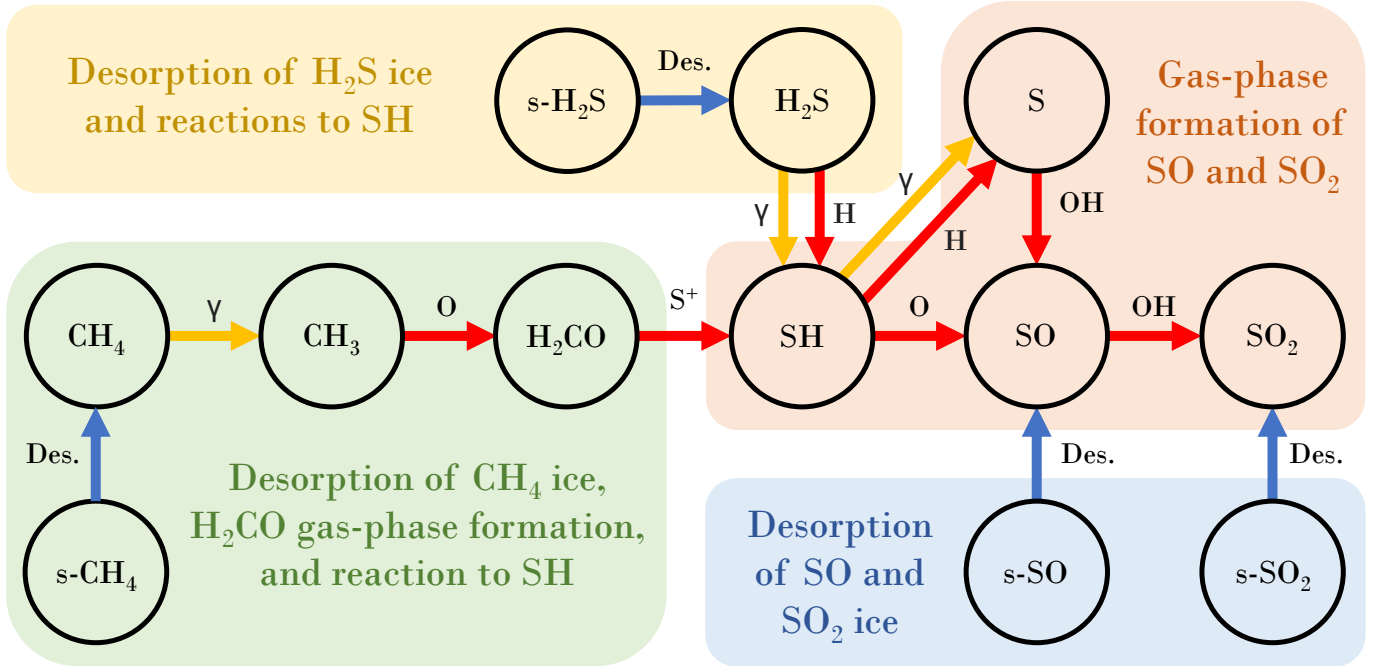


Fig. 5. Chemical reactions leading to the formation of gas-phase SO and SO₂. Species denoted as s-X are located in the ice mantles of dust grains. Reverse reactions are not shown for clarity. Red arrows denote chemical reactions with the annotated species, yellow arrows photodissociation reactions, and blue arrows ice desorption via either thermal or non-thermal mechanisms. Relevant chemical pathways are highlighted with a colored background.

temperature reaches ~ 37 K, leading to a strong increase in its gas-phase abundance (see Fig. 4). Thermal desorption of SO₂ only occurs for the highest velocity shocks in the densest media. However, desorption of H₂S ice (at $T_{\text{dust}} \sim 47$ K) is also relevant for the gas-phase chemistry as subsequent photodissociation and reactions with atomic H lead to an increased abundance of SH and atomic S (i.e., the yellow route of Fig. 5).

3.2.3. Dependence on ultraviolet radiation field

As shown in Reactions (2)-(4) and Fig. 5, gas-phase formation of SO and SO₂ is dependent on radicals such as OH and SH. These radicals can be created in various ways, including high-temperature chemical reactions (Prasad & Huntress 1980;

Hartquist et al. 1980). However, a more dominant physical process for enhancing the abundance of OH and SH is the photodissociation of H₂O, H₂S, and interestingly also CH₄. This process is dependent on the strength of the local ultraviolet radiation field, which is parametrized here with G_0 .

In Fig. 6, the maximum abundance of SO as function of V_s and n_H is presented for $G_0 = 10^{-2}$. Similar figures of both SO and SO₂ for various G_0 are presented in Appendix D.2. Decreasing the strength of the ultraviolet field decreases the amount of SO and SO₂ produced in high-velocity ($\gtrsim 4$ km s⁻¹) shocks due to less OH being produced through H₂O photodissociation. On the other end, increasing G_0 shifts the spot for efficient SO and SO₂ formation in the shock to higher densities to create the favorable conditions for their chemistry. Moreover,

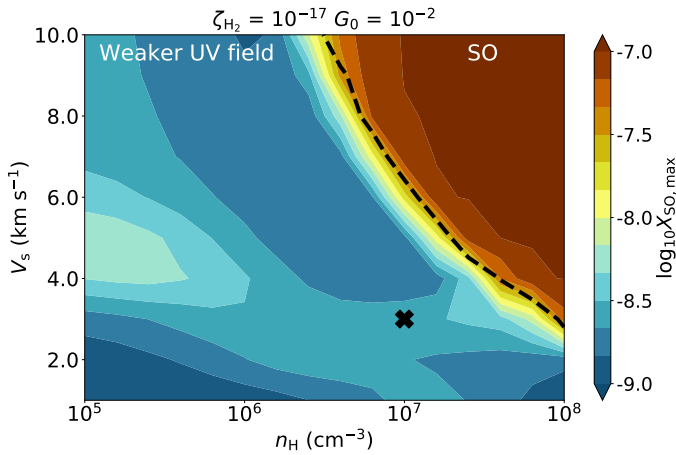


Fig. 6. Similar figure as Fig. 4, but now for SO in shocks with a weaker ultraviolet field ($G_0 = 10^{-2}$).

Table 2. Input abundances of key Sulfur-bearing species

Case	S	SO	SO ₂	s-SO	s-SO ₂	s-H ₂ S
Fiducial	1(-6)	1(-9)	1(-9)	1(-7)	1(-7)	2(-5)
Low-S	1(-8)	1(-9)	1(-9)	1(-7)	1(-7)	2(-5)
High-S	2(-5)	1(-9)	1(-9)	1(-7)	1(-7)	1(-7)

Notes. a(b) represents a $\times 10^b$. Species denoted as s-X are located in the ice mantles of dust grains. All other abundances are the same as those presented in Appendix B.

for $G_0 \gtrsim 10$ photodissociation of thermally desorbed H₂S (i.e., when $n_H \gtrsim 10^7 \text{ cm}^{-3}$ and $V_s \gtrsim 4 \text{ km s}^{-1}$) significantly increases the abundance of atomic S (i.e., yellow route of Fig. 5) which leads to an increase in abundances of SO and in particular SO₂ (see Fig. D.3).

At lower shock velocities ($V_s \lesssim 4 \text{ km s}^{-1}$), water is not formed in the shock and hence photodissociation of H₂O does not increase the abundance of SO and SO₂. The increase of SO abundance is here due to the green route of Fig. 5, which is initialized by thermal desorption and subsequent photodissociation of CH₄. This formation route is most relevant for $G_0 = 1$; for lower G_0 , the ultraviolet radiation field is not strong enough to photodissociate significant amounts of CH₄. For higher G_0 , higher densities are required for the efficient production of SO through the green route.

3.2.4. Dependence on initial conditions

The abundance of all species, including SO and SO₂, is dependent on the initial abundances. The interstellar ice abundances of SO and SO₂ are not well known, with only a tentative detection for SO₂ (Boogert et al. 1997; Zasowski et al. 2009). Here, the ice abundances of SO and SO₂ are estimated at the 10^{-7} level based on what is found in cometary ices (Calmonte et al. 2016; Rubin et al. 2019; Altwegg et al. 2019). However, they are only relevant in the shocks where thermal desorption of these ices occurs (i.e., the blue route of Fig. 5 and to the right of the dashed line in Fig. 7). For those shocks, increasing or decreasing the ice abundances with an order of magnitude will also result in an increase or decrease of the gas-phase abundances with an order of magnitude, given that the ice abundance is dominant over what is achieved through gas-phase chemistry alone.

For shocks where the abundance of SO and SO₂ is mainly increased through gas-phase chemistry (i.e., to the left of the dashed line in Fig. 7), the maximum abundance reached is mostly directly dependent on the initial abundance of atomic sulfur. To test the effect of initial atomic S abundance on the maximum abundances of SO and SO₂, shock models are calculated for a lower gas-phase atomic sulfur abundance (Low-S: $X_S = 10^{-8}$) and assuming that almost all sulfur is atomic and in the gas phase (High-S: $X_S = 2 \times 10^{-5}$). The initial abundances for these cases are presented in Table 2. The resulting maximum SO and SO₂ abundances for these models are shown in Fig. 7.

In the Low-S case, the maximum abundance of SO drops for both low and high velocity shocks (excluding higher densities where the abundance is increased through thermal ice sublimation). This is straightforward to interpret since both S and S⁺ are main formation precursors of SO in both the red and green formation routes of Fig. 5, respectively. At lower densities ($\lesssim 10^6 \text{ cm}^{-3}$), the maximum SO₂ abundance drops in a similar way since less SO is formed. However, at intermediate densities ($\sim 10^7 \text{ cm}^{-3}$), slightly more SO₂ is formed in high velocity shocks compared to Fig. 4. This is because atomic S and SO are ‘competing’ to react with OH to form respectively SO and SO₂. Dropping the initial atomic S abundance results in more SO₂ being formed, also because the majority of the gas-phase SO here originates from thermal ice sublimation.

In the High-S case, not only the chemistry but also the thermal structure of the shock is slightly altered since cooling through atomic S becomes significant. Furthermore, more SO is formed through both the red and green routes of Fig. 5. Interestingly, the maximum abundance of SO₂ attained decreases compared to Fig. 4 since with more atomic S in the gas phase to react with OH, less OH is available to react with SO toward SO₂.

The initial abundances of other species are also important for the chemistry of SO and SO₂. Increasing or decreasing the amount of atomic oxygen results in respectively more or less OH and H₂O formed in the shock and thus directly affects the abundances of SO and SO₂. Moreover, reactions with atomic carbon are a main destruction pathway for SO (forming CS; Hartquist et al. 1980), hence an increased initial atomic carbon abundance results in lower abundances of both SO and SO₂ in the shock.

In reality, the initial dark cloud abundances may also be altered during the infall from envelope to disk since both the temperature and the strength of the ultraviolet radiation field increase (e.g., Aikawa et al. 1999; Visser et al. 2009; Drozdovskaya et al. 2015). To take this into account, a pre-shock model should be calculated. In Appendix C, the effect of calculating such a pre-shock model on the abundance of SO attained in the shock is presented. Any alterations on the pre-shock conditions are only important for the chemistry of SO and SO₂ if the timescale over which the pre-shock is calculated is longer than about 10% of the photodesorption timescale. For typical infalling envelopes, this is not expected to be relevant.

3.3. Effect of grain size and PAHs

The size of the dust grains has an effect on both the physical structure of the shock and on the chemistry (e.g., Guillet et al. 2007; Miura et al. 2017). In Fig. 8, the abundance profiles of SO and SO₂ are shown for the fiducial model and for the same model while adopting a typical interstellar medium grain size distribution (computed using a single size of $\sim 0.03 \mu\text{m}$; Mathis et al. 1977; Godard et al. 2019). The decrease in grain size leads to stronger coupling between gas and dust and thus to a slightly higher dust temperature ($\sim 35 \text{ K}$ against $\sim 30 \text{ K}$ for $0.2 \mu\text{m}$

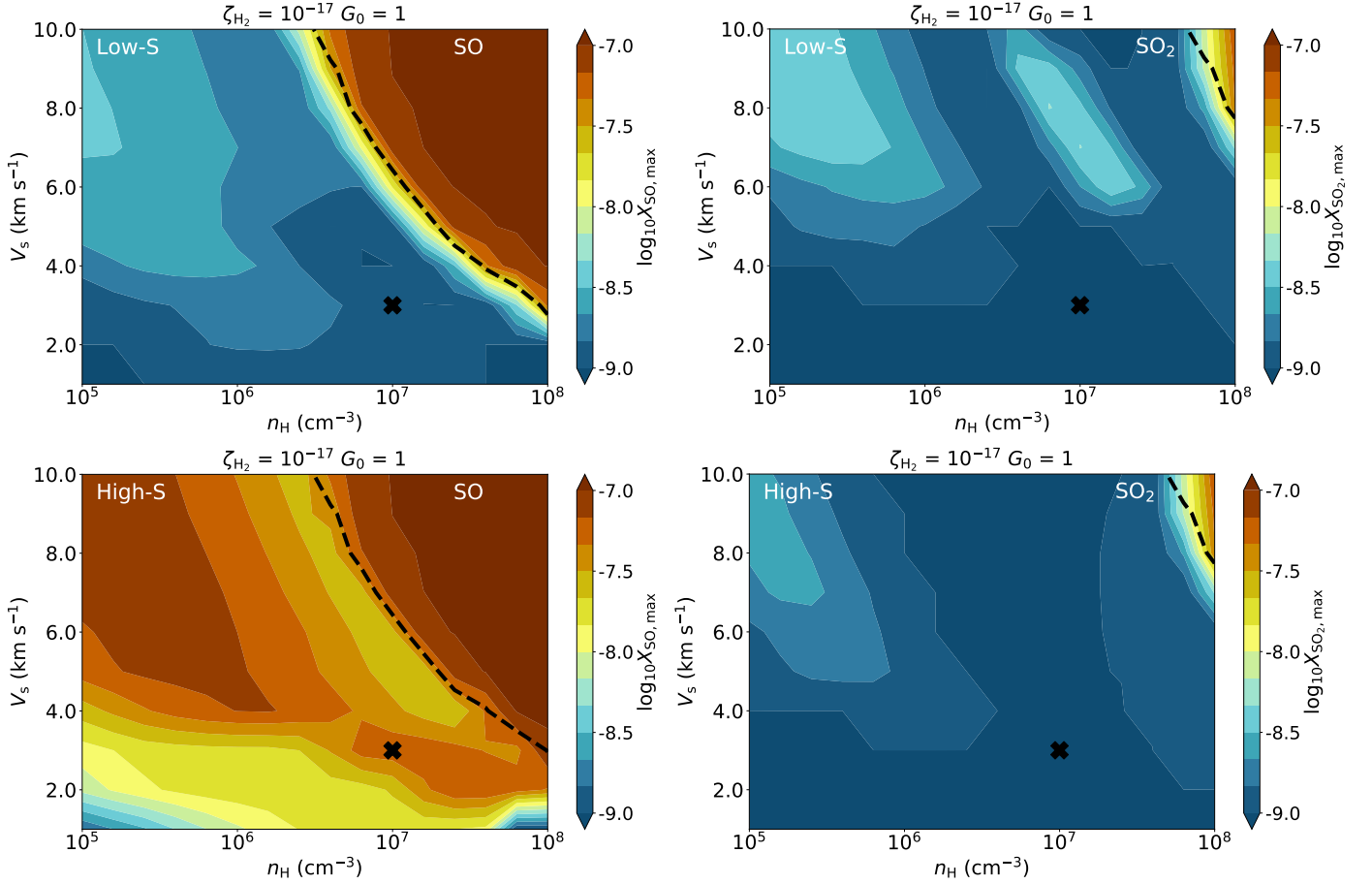


Fig. 7. Maximum abundance reached (in color) of SO (left) and SO₂ (right) in shock models as function of initial n_{H} and V_s for an initial gas-phase atomic S abundance of 10^{-8} (top row) and 10^{-5} (bottom row). All other physical parameters are kept constant to the fiducial values and listed on top of the figure. The black cross indicates the position of the fiducial model. The dashed black line shows the ice line, i.e., where 50% of the ice is thermally desorbed into the gas in the shock.

grains). Moreover, the increased gas-grain coupling results in the dust being a dominant coolant in the tail of the shock reducing the length of the shock (~ 1 AU against ~ 3 AU for the fiducial model). Nevertheless, the abundance of SO is increased in the shock, but the maximum abundance reached is about a factor of 2 – 3 lower compared to the fiducial model. No significant increase is seen for SO₂. Miura et al. (2017) found that the dust temperature easily reaches ~ 50 K in low-velocity shocks taking into account dust aerodynamics for ISM-size dust grains. The full treatment of dust physics, including aerodynamic heating and grain-grain interactions (Guillet et al. 2007, 2009, 2011, 2020; Miura et al. 2017), will likely move the ice line of Fig. 4 toward lower n_{H} and V_s , but is beyond the scope of this work.

Contrary to dust grains, the effect of polycyclic aromatic hydrocarbons on the physical structure of J -type shocks is negligible since polycyclic aromatic hydrocarbons do not contribute to the cooling and only little to the heating if a strong ultraviolet field is present. However, polycyclic aromatic hydrocarbons do play a key role in the ionization balance and are therefore highly relevant for the chemistry (Flower & Pineau des Forêts 2003). In Fig. 8, the abundances of SO and SO₂ are also shown for the fiducial model with a polycyclic aromatic hydrocarbon abundance of 10^{-6} (equal to the value derived in the local diffuse interstellar medium; Draine & Li 2007). The maximum abundance of SO is a factor 3 – 4 lower when the abundance of polycyclic aromatic hydrocarbons is at the interstellar medium level because

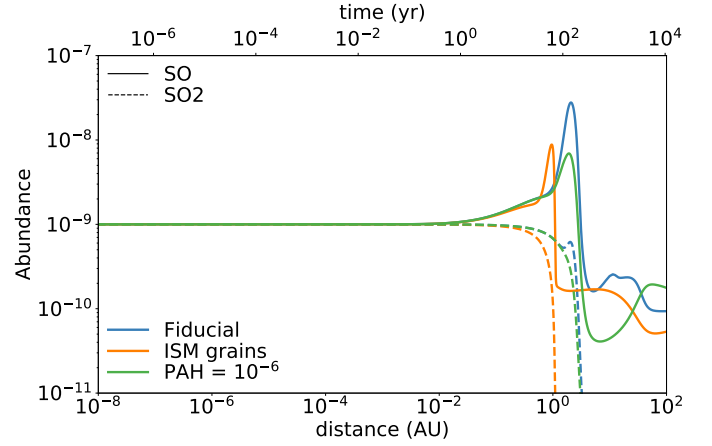


Fig. 8. Abundance profile of SO (solid) and SO₂ (dashed) for the fiducial model (blue) with a polycyclic aromatic hydrocarbon abundance of 10^{-8} using a grain size of $0.2 \mu\text{m}$. Overplotted are the same model using a typical interstellar medium grain size of $0.03 \mu\text{m}$ (orange) and a model with a polycyclic aromatic hydrocarbon abundance of 10^{-6} (green). All other parameters are kept at their fiducial value (see Table 1).

the polycyclic aromatic hydrocarbons are dominating the ionization balance. Consequently, the abundance of other ions, include S^+ , drops by more than an order of magnitude. Given that the

green route of Fig. 5 is the dominant SO formation route here and dependent on S^+ , less SO and subsequently SO_2 are formed. For high-velocity shocks, this effect is not apparent since Reactions (3) and (4) dominate the formation of SO and SO_2 and are independent of ions.

3.4. Other molecular shock tracers: SiO, H_2O , H_2S , CH_3OH , and H_2

Other classical molecular shock tracers include SiO, H_2O , H_2S , CH_3OH , and H_2 . Abundance maps similar to Fig. 4 of SiO, H_2O , H_2S , and CH_3OH are presented in Appendix D.1. SiO is a tracer of shocks with significant grain destruction since then the silicon which is normally locked in dust grains can react with the OH radical to form SiO (e.g., Caselli et al. 1997; Schilke et al. 1997; Gusdorf et al. 2008a,b; Guillet et al. 2009). It is often observed in the high velocity bullets of jets originating from young Class 0 protostars (e.g., Guilloteau et al. 1992; Tychoniec et al. 2019; Taquet et al. 2020). However, the abundance of SiO is not significantly increased in our shock models because most of the Si remains locked up in silicates in the dust grains.

Water is the most abundant ice in protostellar envelopes (Boogert et al. 2015; van Dishoeck et al. 2021), but also frequently observed in outflow shocks (e.g., Flower & Pineau des Forêts 2010; Herczeg et al. 2012; Kristensen et al. 2012; Nisini et al. 2013; Karska et al. 2018). Here, H_2O is efficiently produced in shocks with $V_s \gtrsim 4$ km s $^{-1}$ (see Fig. D.1) and very important for the gas-phase chemistry of SO and SO_2 . Thermal desorption of H_2O is negligible.

H_2S is suggested as a main carrier of sulfur in ices (e.g., Vidal et al. 2017), despite being undetected thus far (Jiménez-Escobar & Muñoz Caro 2011). In the gas phase it is generally observed close to Class 0 protostars where the emission may originate from thermal desorption (Tychoniec et al. 2021), or in outflow shocks (e.g., Holdship et al. 2016). In our shock models, gas-phase formation of H_2S becomes efficient at slightly higher velocities than for water ($V_s \gtrsim 5$ km s $^{-1}$, see Fig. D.1). Moreover, for high-velocity shocks in dense media, H_2S ice is thermally sublimated.

Methanol is a tracer of lower-velocity (< 10 km s $^{-1}$) outflow shocks where ices are sputtered off the dust grains (e.g., Suutarinen et al. 2014). However, since sputtering is not relevant in our single-fluid J -type accretion shock models, no significant increase of CH_3OH abundance is evident except at the highest densities ($\gtrsim 10^8$ cm $^{-3}$) and velocities ($\gtrsim 10$ km s $^{-1}$) where CH_3OH ice thermally sublimates.

Emission from warm H_2 is generally observed toward shocks in protostellar jets (e.g., McCaughrean et al. 1994). Contrary to the molecules discussed above, the abundance of H_2 remains constant along the shock and only thermal excitation occurs. Hence, H_2 can be a powerful diagnostic of accretion shocks in inferring the temperature from a rotational diagram. The population of the lowest 50 rotational and rovibrational levels of H_2 are computed during the shock (Flower & Pineau des Forêts 2003). The rotational diagram of H_2 for the fiducial model is presented in Fig. 9. Since the temperature in this shock reaches about ~ 500 K, levels up to ~ 6000 K are collisionally populated. The population of all higher- E_{up} levels is set during the formation of H_2 on the dust grains following a Boltzmann distribution at 17248 K (1/3 of the dissociation energy of H_2 ; Black & van Dishoeck 1987). Using the rotational diagram of Fig. 9, a rotational temperature ~ 390 K can be derived for this shock.

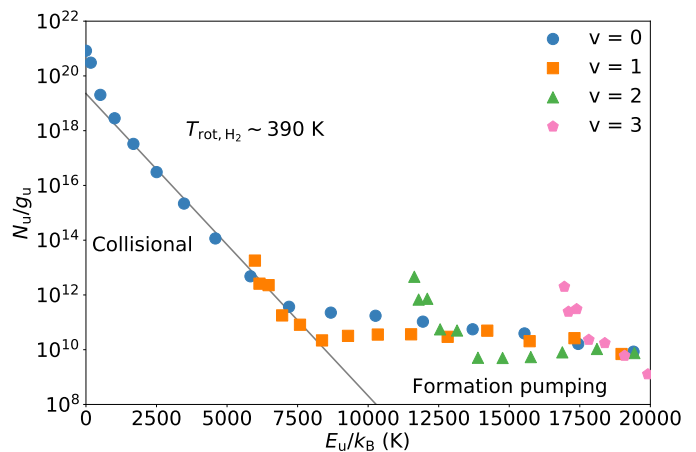


Fig. 9. Rotational diagram of H_2 created from the fiducial shock model. The levels of different vibrational states are indicated with different symbols and colors. Note that only the pure rotational levels with $E_u/k_B \lesssim 6000$ K are collisionally populated. The fit to the pure rotational levels is shown in gray line with the derived rotational temperature annotated.

4. Discussion

4.1. Comparison to SO and SO_2 with ALMA

Emission of warm ($T_{ex} > 50$ K) SO and SO_2 around the disk-envelope interface has been suggested as a possible tracer of accretion shocks (Sakai et al. 2014, 2017; Bjerkeli et al. 2019; Oya et al. 2019; Artur de la Villarmois et al. 2019). Here, we have shown that the abundance of both SO and SO_2 can increase in shocks in typical inner envelope conditions. The abundance of SO is estimated to be $\sim 10^{-8}$ in the suggested accretion shock around the L1527 Class 0/I protostar (Sakai et al. 2014), but this is uncertain since the density of H_2 was guessed. Furthermore, by itself this does not provide strong constraints on the physical conditions of the shock (see Fig. 4), other than that a weak ultraviolet field should be present if SO ice is not thermally sublimated ($G_0 > 10^{-2}$; Fig. D.2). However, it remains difficult to derive abundances with respect to n_H from observations.

Recent the Atacama Large Millimeter/submillimeter Array observations show that the amount of warm SO_2 seems to increase with the bolometric luminosity (i.e., stronger ultraviolet field; Artur de la Villarmois et al. 2019). This is in line with our results since the abundance of SO_2 only increases when a significant ultraviolet field is present (i.e., $G_0 \gtrsim 1$, see Fig. D.3). Especially for higher velocity shocks (> 4 km s $^{-1}$) at higher densities ($> 10^7$ cm $^{-3}$) the maximum abundance of SO_2 increases with 2 orders of magnitude between $G_0 = 1$ and $G_0 = 100$. However, besides a significant UV field, a shock velocity of $V_s \gtrsim 4$ km s $^{-1}$ is necessary to increase the abundance of SO_2 and the maximum abundance reached in the shock is also dependent on the initial density n_H .

Comparing the ratio of observed column densities to those derived from our shock models can more accurately constrain the physical conditions in a shock. In Fig. 10, the column density ratio of SO_2/SO in the shock is presented as function of V_s and n_H for $G_0 = 1$ and $G_0 = 100$. The only protostellar source where both SO and SO_2 have been detected related to a possible accretion shock is Elias 29 (Oya et al. 2019), where a column density ratio of $0.5^{+0.4}_{-0.2}$ is derived. When $G_0 = 1$, this would imply a low-density shock ($< 10^6$ cm $^{-3}$) propagating at either low (< 2 km s $^{-1}$) or high (> 7 km s $^{-1}$) velocities, see upper panel in

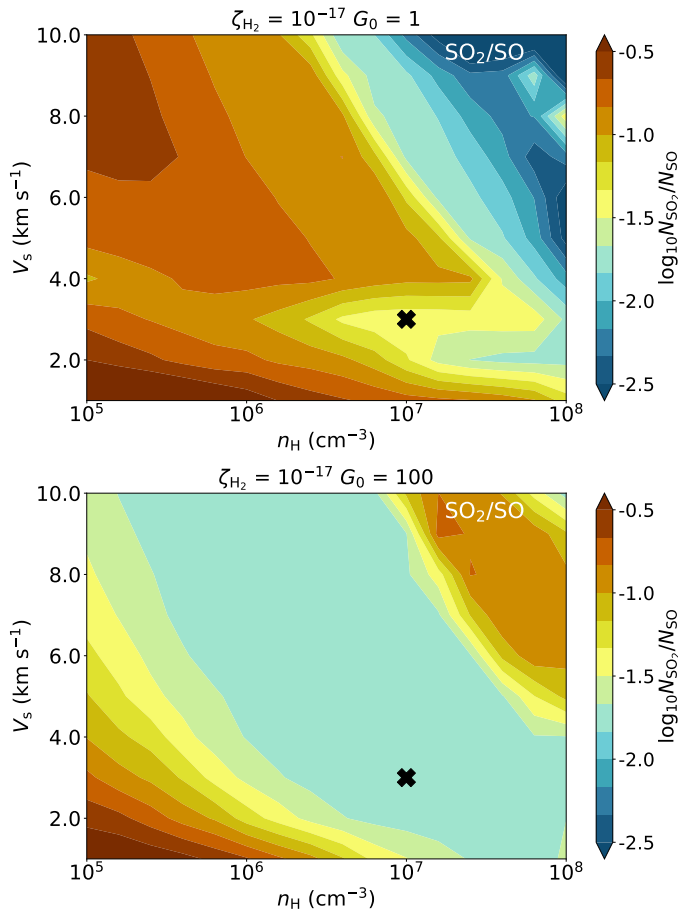


Fig. 10. Ratio of column densities of SO₂ and SO as function of n_{H} and V_s for $G_0 = 1$ (top) and $G_0 = 100$ (bottom). All other physical parameters are kept constant to the fiducial values and listed on top of the figure. The black cross indicates the position of the fiducial model.

Fig. 10. For $G_0 = 100$, shocks propagating at higher velocities (> 4 km s⁻¹) at high densities ($> 10^7$ cm⁻³) produce SO₂/SO column density ratios similar to that of Elias 29 (Oya et al. 2019, see lower panel in Fig. 10).

Since both SO and SO₂ emission is also linked to outflow activity and passive heating in the inner envelope (Codella et al. 2014; Tabone et al. 2017; Lee et al. 2018; Taquet et al. 2020; Harsono et al. 2021), high spatial resolution observations with the Atacama Large Millimeter/submillimeter Array are necessary to both spatially and spectrally disentangle the different protostellar components and determine their emitting area. Moreover, to robustly test if the SO and SO₂ emission traces the accretion shock, key species for their gas-phase formation such as H₂S and H₂CO should be observed on similar scales. This also allows for more accurately constraining the physical conditions of a shock.

4.2. Predicting H₂, H₂O, and [S I] with JWST

With the launch of the *James Webb* Space Telescope, near-infrared and mid-infrared shock tracers such as H₂, H₂O, and [S I] can be used to search for accretion shocks. Although the abundance of H₂ does not increase in accretion shocks, it is still thermally excited and therefore a good shock tracer at near-infrared and mid-infrared wavelengths (e.g., McCaughrean et al. 1994). Observing the rotational sequence of H₂ in the near-infrared and mid-infrared could thus put constraints on the

physical properties of the shock. For low-velocity shocks ($\lesssim 5$ km s⁻¹), most of the strong H₂ transitions (i.e., 0-0 S(3) and 0-0 S(5)) are located in the the *James Webb* Space Telescope/MIRI range (5 – 28 μ m). For higher velocity shocks ($\gtrsim 5$ km s⁻¹), also vibrational levels of H₂ get populated and the strongest H₂ transitions shift toward the NIRSpec range of the *James Webb* Space Telescope (1 – 5 μ m).

Warm water is also a well-known and important tracer of shocks (e.g., Herczeg et al. 2012; Kristensen et al. 2012; Nisini et al. 2013; van Dishoeck et al. 2021). Observing H₂O toward an accretion shock is a clear indication of a high-velocity shock ($\gtrsim 4$ km s⁻¹, see Fig. D.1). Warm water should be easily detectable with the *James Webb* Space Telescope as both the vibrational band around 6 μ m and several pure rotational transitions at > 10 μ m fall within the MIRI range (5 – 28 μ m). Moreover, the detection of mid-infrared lines of the OH radical can be an indication of H₂O photodissociation (Tabone et al. 2021), and therefore be spatially co-located with emission of warm SO and SO₂ in the Atacama Large Millimeter/submillimeter Array observations.

An alternative shock tracer in the the *James Webb* Space Telescope/MIRI band is the [S I] 25 μ m transition. It is often observed in protostellar outflows (e.g., Neufeld et al. 2009; Lahuis et al. 2010; Goicoechea et al. 2012), where the intensity of the line can range from a few % of the H₂ 0-0 S(3) and 0-0 S(5) lines to a factor of a few higher. It is therefore uncertain whether [S I] 25 μ m will be detectable toward accretion shocks, also given that the sensitivity of MIRI drops at the highest wavelengths. Moreover, at 25 μ m, the angular resolution of the *James Webb* Space Telescope is $\sim 1''$ (i.e., 300 AU at a distance of 300 pc) which complicates the spatial disentanglement of an accretion shock with outflow emission.

5. Summary

We have modeled low-velocity (≤ 10 km s⁻¹) non-magnetized *J*-type shocks at typical inner envelope densities (10^5 – 10^8 cm⁻³) using the Paris-Durham shock code to test SO and SO₂ as possible molecular tracers of accretion shocks. The main conclusions of this work are as follows:

- In low-velocity (~ 3 km s⁻¹) shocks, SO can be efficiently formed at intermediate densities ($\sim 10^7$ cm⁻³) through the SH radical reacting with atomic O. Here, SH is formed through S⁺ reacting primarily with H₂CO; though the latter species is a representative of any hydrocarbon.
- Both SO and SO₂ can be efficiently formed through reactions of atomic S with OH in high-velocity ($\gtrsim 4$ km s⁻¹) shocks in low-intermediate dense environments ($\lesssim 10^7$ cm⁻³). The formation of SO and SO₂ occurs in the end of the shock, where the abundance of OH is enhanced through photodissociation of earlier formed H₂O.
- Thermal desorption of SO ice can occur in high-velocity ($\gtrsim 4$ km s⁻¹) shocks at high densities ($\gtrsim 10^7$ cm⁻³), while thermal desorption of SO₂ ice is only relevant at the highest shock velocities ($\gtrsim 10$ km s⁻¹) and highest densities ($\gtrsim 10^8$ cm⁻³).
- The chemistry of both SO and SO₂ through all formation routes is strongly linked to the strength of the local ultraviolet radiation field since the formation of both the OH and SH radicals is linked to photon processes such as photodissociation and photoionization. In particular, SO₂ is only formed in the shock through gas-phase chemistry if the strength of the local ultraviolet radiation field is greater or equal to the isotropic standard interstellar radiation field.

- The composition of infalling material, in both gas-phase species like atomic O and S and in the ices such as H₂S, CH₄, SO, and SO₂, is highly relevant for the chemistry of SO and SO₂ in accretion shocks.
- Observations of warm SO and SO₂ should be complemented with key species for their formation such as H₂S and H₂CO. Moreover, the launch of the *James Webb* Space Telescope will add additional near-infrared and mid-infrared accretion shock tracers such as H₂, H₂O, and [S I] 25 μ m.

Our results highlight the key interplay between both physics and chemistry in accretion shocks. The abundance of gas-phase molecules such as SO and SO₂ is not solely determined through high-temperature gas-phase chemistry, but also thermal sublimation of ices and photodissociation through ultraviolet radiation play an important role. Additional high-angular resolution observations with the Atacama Large Millimeter/submillimeter Array are necessary to spatially disentangle the disk from the envelope in embedded systems in order to assess the physical properties of accretion shocks at the disk-envelope interface. Moreover, future facilities such as the *James Webb* Space Telescope will provide additional near-infrared and mid-infrared shock tracers such as H₂, OH, and H₂O. It is crucial to attain the physical structure of accretion shocks since it determines the chemical composition of material that enters the disk and is eventually incorporated in planets.

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Appendix A: Updated cooling of NH₃

Typically, cooling by NH₃ in the optically thin limit is calculated using the empirical relation derived by [Le Bourlot et al. \(2002\)](#). However, this relation does not take into account that the populations get thermalized if the density gets above the critical density. Therefore, the cooling by NH₃ is recalculated using the RADEX software ([van der Tak et al. 2007](#)). Assuming all emission is optically thin (i.e., low column densities) and emitted through rotational lines, a grid of gas temperatures and H₂ densities is set. The full range explored is 5–5000 K and 10²–10¹² cm⁻³, respectively. The linewidth is set to 2 km s⁻¹. The cooling (in erg s⁻¹) for each grid point is calculated by summing over the emission of each spectral line and dividing that by the assumed column density.

Using the cooling grid computed with RADEX, a cooling function, Λ , with T_{gas} and n_{H_2} as free parameters can be derived. Minimum χ^2 fitting of the grid computed with RADEX was used to calculate Λ , which resulted in,

$$\Lambda_{\text{NH}_3} = \Lambda_{\infty} \left(1 - \exp \left[\frac{-n_{\text{H}_2}}{n_0} \right] \right), \quad (\text{A.1})$$

with Λ_{∞} and n_0 numerically derived as,

$$\Lambda_{\infty} = 2.85 \times 10^{-15} \left(1 - \exp \left[\frac{-T_{\text{gas}}}{81.7} \right] \right)^4, \quad (\text{A.2})$$

$$n_0 = 8.33 \times 10^8 \left(1 - \exp \left[\frac{-T_{\text{gas}}}{125} \right] \right). \quad (\text{A.3})$$

Λ_{∞} is the cooling of NH₃ in the local thermodynamic equilibrium limit, whereas n_0 is a measure of a critical density above which local thermodynamic equilibrium effects become relevant. Both parameters are dependent on T_{gas} .

The function in Eq. (A.1) is presented in Fig. A.1 for a few values of T_{gas} . Overplotted are the RADEX calculations and the empirical relation of [Le Bourlot et al. \(2002\)](#). Indeed, at high densities ($\gtrsim 10^9$ cm⁻³) the latter relation starts overproducing the cooling computed with RADEX by orders of magnitude. However, the updated cooling function represent the cooling by RADEX within a factor of 3 over the entire temperature and density range. Since densities of $\gtrsim 10^9$ cm⁻³ are frequently reached in our shock models, the updated cooling function was adopted in the updated version of the shock code.

In high-temperature shocks, significant amounts of NH₃ might be produced through gas-phase chemistry, enhancing the density and thus the optical depth of the spectral lines. Moreover, the density of NH₃ can be increased through ice sublimation in high-density shocks. If the lines of NH₃ become optically thick, Eq. A.1 is no longer valid and cooling tables similar to those of [Neufeld & Kaufman \(1993\)](#) should be calculated for NH₃ taking into account opacity effects. This is beyond the scope of this work.

Appendix B: Input abundances

The input abundances used in the shock models are presented in Table B.1. The total elemental abundances match those of [Flower & Pineau des Forêts \(2003\)](#). Most of the oxygen, carbon, and sulfur budget is locked up in dust grains and in the ice mantles. The initial abundances of oxygen and nitrogen-bearing ices are equivalent to those presented by [Boogert et al. \(2015\)](#), while the sulfur-bearing ice abundances are matched to those found in both protostellar envelopes and comets ([Calmonte et al.](#)

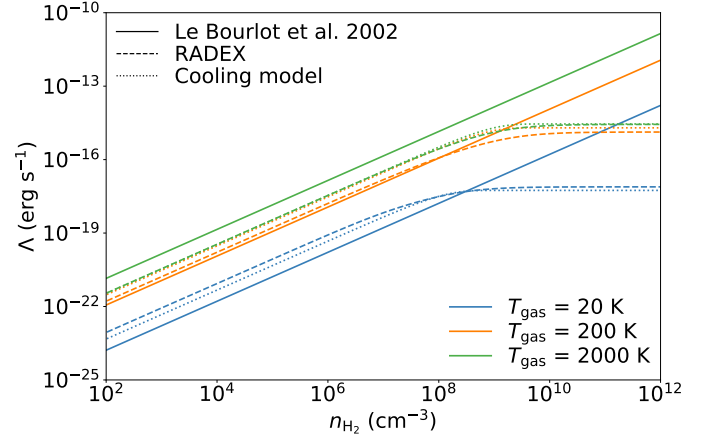


Fig. A.1. Cooling by NH₃ as function of n_{H_2} for various T_{gas} (colors). The empirical relation by [Le Bourlot et al. \(2002\)](#) is shown as the solid lines, the RADEX computations in the dashed line, and the derived cooling function of Eq. (A) in the dotted line.

[2016](#); [Rubin et al. 2019](#); [Altwegg et al. 2019](#); [Navarro-Almaida et al. 2020](#)). The gas-phase abundances of the key sulfur-bearing molecules, SO, SO₂, CS, OCS, and H₂S, are taken from [van der Tak et al. \(2003\)](#). All other molecular gas-phase abundances are set equal to those derived on cloud scales by [Tafalla et al. \(2021\)](#). The gas-phase abundance of atomic S in protostellar envelopes is highly uncertain. It is set here at a 10⁻⁶ level, which is consistent with the models of [Goicoechea & Cuadrado \(2021\)](#). The effect of initial atomic S abundance on the maximum abundances of SO and SO₂ attained in the shock is presented in Sect. 3.2.4.

Table B.1. Dark cloud input abundances

H	5.0×10^{-5}	Grains	4.6×10^{-11}
H ₂	5.0×10^{-1}	s-H ₂ O	1.0×10^{-4}
He	1.0×10^{-1}	s-O ₂	1.0×10^{-8}
C	1.0×10^{-8}	s-CO	8.3×10^{-6}
O	1.0×10^{-6}	s-CO ₂	1.3×10^{-5}
CO	8.5×10^{-5}	s-CH ₄	1.5×10^{-6}
C ₂ H	2.0×10^{-9}	s-N ₂	3.9×10^{-6}
C ₃ H ₂	2.0×10^{-10}	s-NH ₃	1.0×10^{-6}
CH ₃ OH	1.5×10^{-9}	s-CH ₃ OH	1.9×10^{-5}
N	1.0×10^{-6}	s-H ₂ CO	6.2×10^{-6}
CN	1.5×10^{-9}	s-HCO ₂ H	7.2×10^{-6}
HCN	3.0×10^{-9}	s-OCS	2.1×10^{-7}
HNC	1.0×10^{-9}	s-H ₂ S	1.8×10^{-5}
N ₂	3.5×10^{-5}	s-SO	1.0×10^{-7}
S	1.0×10^{-6}	s-SO ₂	1.0×10^{-7}
H ₂ S	1.0×10^{-9}	c-O	1.4×10^{-4}
CS	3.0×10^{-9}	c-Si	3.4×10^{-5}
SO	1.0×10^{-9}	c-Mg	3.7×10^{-5}
SO ₂	1.0×10^{-9}	c-Fe	3.2×10^{-5}
OCS	2.0×10^{-9}	c-C	1.6×10^{-4}
Si	1.0×10^{-10}	HCO ⁺	1.5×10^{-9}
Fe	1.5×10^{-8}	N ₂ H ⁺	1.5×10^{-10}
PAH	1.0×10^{-8}		

Notes. Abundances are with respect to n_{H} . Species in ice mantles on dust grains are indicated as s-X. Species in composing the cores of dust grains are denoted as c-X.

Appendix C: Changing the pre-shock conditions

To test the effect of alteration of infalling dark cloud material in the inner envelope on an accretion shock, various pre-shock models are calculated. Since the increase of the temperature and protostellar radiation field are negligible in the outer parts of the envelope (e.g., Drozdovskaya et al. 2015), changes in temperature, composition, and ionization balance are most efficient in the inner ~ 1000 AU of the envelope when the material is already approaching the disk and protostar. Given that the infall velocity goes as $v(r) = \sqrt{2GM/r}$, it takes about ~ 100 years to move from 1000 AU to disk scales (~ 100 AU) for a $0.5 M_{\odot}$ star. However, depending on the mass of the protostar and the size of the disk, this pre-shock timescale $t_{\text{pre-shock}}$ might range from, for example, $\sim 10 - 10^4$ years. Therefore, the effect of infall time (i.e., $t_{\text{pre-shock}}$) on the accretion shock is explored here. The thermal balance is included in the pre-shock model, but for simplicity the density is assumed to remain constant.

Given that the infalling material is ionized and photodissociated on a timescale of (Heays et al. 2017),

$$\tau_{\text{ionize/disso}} \sim \frac{10 - 100}{G_0} \text{ yrs}, \quad (\text{C.1})$$

the ionization balance is fully altered for all $t_{\text{pre-shock}} \geq 10$ years if the strength of the local ultraviolet field is $G_0 \geq 1$. For models with weaker ultraviolet fields ($G_0 = 10^{-3} - 10^{-1}$), the ionization balance is dependent on $t_{\text{pre-shock}}$. Moreover, the ice composition may be altered on the photodesorption and adsorption timescales for $0.2 \mu\text{m}$ grains of (Hollenbach et al. 2009; Godard et al. 2019),

$$\tau_{\text{photodes}} \approx \frac{2 \times 10^5}{G_0} \text{ yrs}, \quad (\text{C.2})$$

$$\tau_{\text{adsorb}} \approx \frac{10^9 - 10^{10}}{n_{\text{H}}} \text{ yrs}. \quad (\text{C.3})$$

It is important to note here that if $\tau_{\text{ionize/disso}} < \tau_{\text{adsorb}}$, any photodesorbed species are dissociated before they can adsorb back on the grain. This increases the abundance of atomic and ionic species.

The abundances of S^+ , S , SO , SO_2 , H_2S , and $\text{s-H}_2\text{S}$ during a pre-shock calculation are presented in Fig. C.1 for the fiducial model conditions (see Table 1). Following Eq. (C.1), the ionization balance of S^+/S is set after ~ 10 years. However, at longer timescales of $\sim 10^4 - 10^5$ years, the abundances of both S^+ and S increase above the initial abundance of atomic S . On these timescales, H_2S ice is photodesorbed (see Eq. (C.2)) and subsequently photodissociated, therefore enhancing the abundance of both atomic S and S^+ . It is interesting to note that since the abundance of H_2S is much higher than that of atomic S in the gas phase, already on timescales of $0.1\tau_{\text{photodes}}$ the abundance of atomic S is increased with a factor of a few. Meanwhile, the abundance of SO is increased through gas-phase chemistry via Reaction (3), and SO_2 is photodissociated into SO and atomic O .

The effect of the length of the pre-shock model on the maximum abundances of SO in the subsequent accretion shock is presented in Fig. C.2. Up to $t_{\text{pre-shock}} \lesssim 1000$ years, the maximum abundance reached in the shock is independent of the time over which the pre-shock model is calculated. In the $t_{\text{pre-shock}} = 10000$ years case, photodesorption and subsequent photodissociation of H_2S increases the initial atomic S abundance and hence the maximum abundance of SO reached in the shock. For different strengths of the ultraviolet radiation field (i.e., different G_0), the timescale for photodesorption changes (i.e., Eq. (C.2)). $t_{\text{pre-shock}}$ does thus not affect the final abundances of SO (and

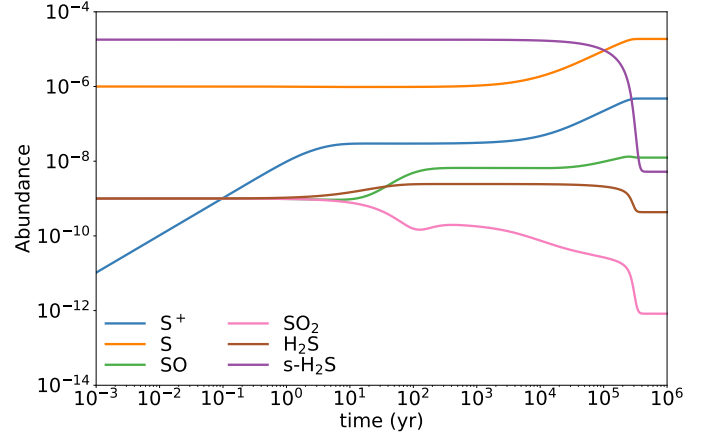


Fig. C.1. Abundances of S^+ , S , SO , SO_2 , H_2S , and $\text{s-H}_2\text{S}$ during a pre-shock calculation. All physical parameters are kept constant to the fiducial values.

SO_2) in the shock if less than roughly $\sim 10\%$ of the photodesorption timescale τ_{photodes} , given that $\tau_{\text{adsorb}} > \tau_{\text{ionize/disso}}$.

Appendix D: Abundance grids

Appendix D.1: H_2S , H_2O , SiO , and CH_3OH for $G_0 = 1$

In Fig. D.1, the maximum abundance for H_2S , H_2O , SiO , and CH_3OH are presented as function of initial n_{H} and V_s . All other parameters are set to their fiducial value (e.g., $G_0 = 1$).

Appendix D.2: SO and SO_2 for various G_0

In Figs. D.2 and D.3, the maximum abundance for respectively SO and SO_2 are presented as function of initial n_{H} and V_s for various strengths of the ultraviolet radiation field (i.e., $G_0 = 10^{-3} - 10^2$).

Appendix E: Higher magnetized environments

The focus of this paper is non-magnetized J -type shocks. However, in reality, magnetic fields are present in protostellar envelopes where observations hint at intermediate or strong magnetic field strengths (i.e., $\sim 0.1 - 1$ mG; Hull et al. 2017). The structure of shocks depends strongly on the strength of the magnetic field. In weakly magnetized regions, the emerging shock will remain a magnetized J -type shock with a similar structure to non-magnetized J -type shocks. However, in higher magnetized environments, the ions and electrons can decouple from the neutral species through the Lorentz force. While slowing down through the Lorentz force, the ions exert a drag force on the neutral fluid. Both fluids slow down and compress, resulting in an increase of the temperature and the emergence of a C -type (strong magnetic field strength) and CJ -type (intermediate magnetic field strength) shocks (Draine 1980; Flower & Pineau des Forêts 2003; Godard et al. 2019). Below, the results of C and CJ -type accretion shocks are shortly discussed, where in both cases a pre-shock model was calculated over a timescale $t_{\text{pre-shock}} = 100$ years (see Appendix C for details).

Appendix E.1: C -type shocks

In interstellar C -shocks, the mass of the ionized fluid is dominated by small dust grains (which couple well to ions) and ion-

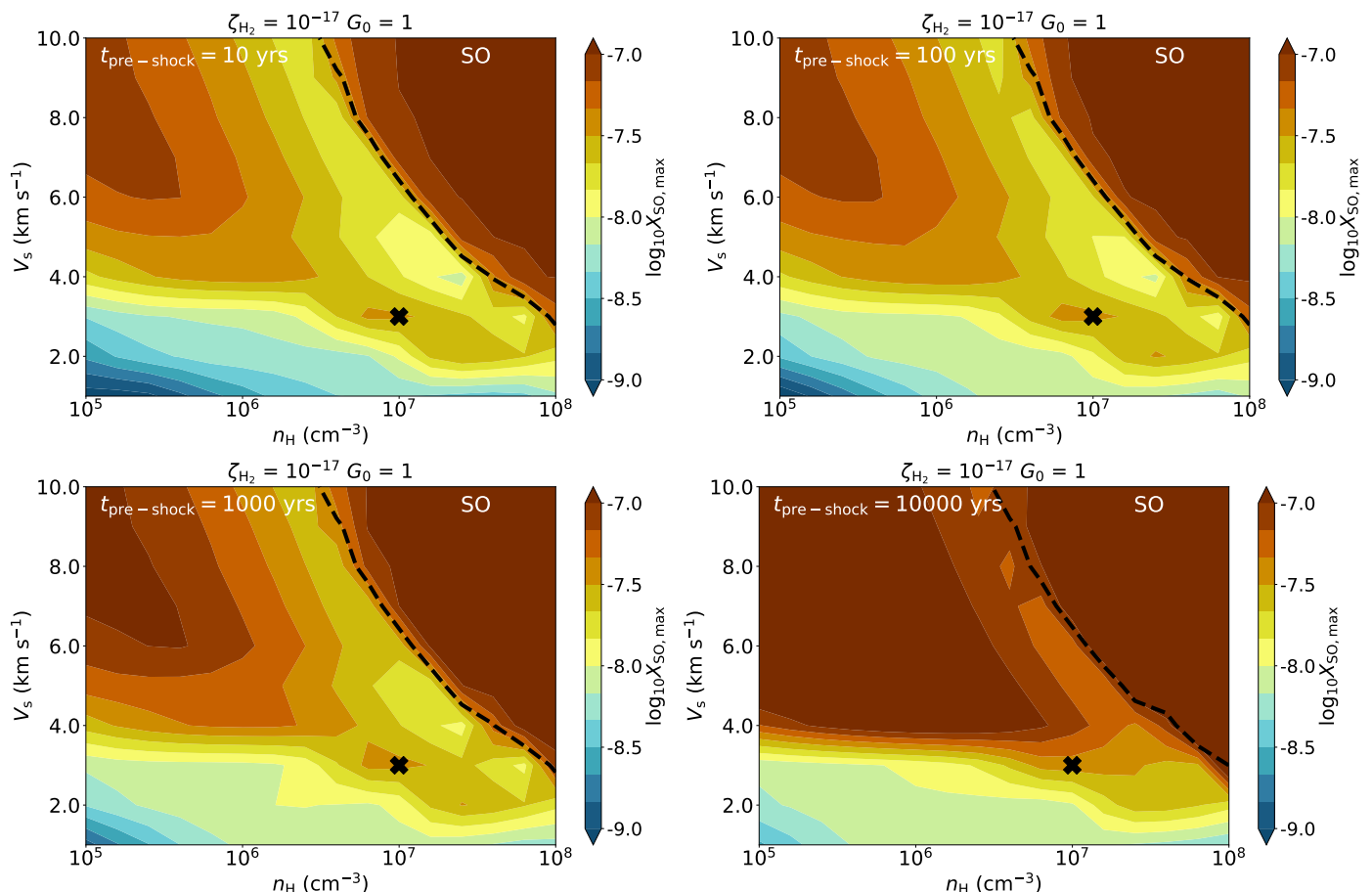


Fig. C.2. Maximum abundance reached (in color) of SO in shock models as function of initial n_{H} and V_s after calculating a pre-shock model for 10 (top left), 100 (top right), 1000 (bottom left), and 10000 (bottom right) years. All other physical parameters are kept constant to the fiducial values and listed on top of the figure. The black cross indicates the position of the fiducial model. The dashed black line shows the ice line, i.e., where 50% of the ice is thermally desorbed into the gas in the shock.

ized polycyclic aromatic hydrocarbons (e.g., [Godard et al. 2019](#)). However, since dust grows to larger grains (here assumed at $0.2 \mu\text{m}$) in protostellar envelopes, it will likely be better coupled to the neutral fluid ([Guillet et al. 2007, 2020](#)). Moreover, polycyclic aromatic hydrocarbons are lacking in the gas, likely frozen out onto dust grains ([Geers et al. 2009](#)). Hence, the mass of the ion fluid exerting a drag force on the neutrals is determined solely by small ions such as S^+ , C^+ , HCO^+ , and N_2H^+ . Having more small ions (e.g. due to a stronger ultraviolet radiation field) therefore strongly affects the structure of the *C*-type shock.

Since the decoupling between ions and neutrals is very slow, the length of *C*-type shocks is on envelope scales (i.e., $\gtrsim 1000 \text{ AU}$) for most of the parameter space considered (see Table 1). Only for high velocity ($> 8 \text{ km s}^{-1}$) *C*-shocks at high densities ($> 10^7 \text{ cm}^{-3}$) the total length of the shock is on disk scales of $< 100 \text{ AU}$. For such shocks, the abundance of SO is slightly increased through gas-phase chemistry following the green route of Fig. 5, while the abundance of SO_2 is not increased in these *C*-type shocks.

Contrary to *J*-type shocks, an increased polycyclic aromatic hydrocarbons abundance also affects the physical structure of a *C*-type shock as ionized polycyclic aromatic hydrocarbons carry most of the mass of the ion fluid if dust grains are assumed to be coupled to the neutrals ([Flower & Pineau des Forêts 2003](#)). A higher abundance of polycyclic aromatic hydrocarbons results in a stronger decoupling between ions and neutrals and conse-

quently a decrease in the shock length. Hence, in a larger part of the parameter space, *C*-type shocks on disk scales ($< 100 \text{ AU}$) are produced. However, given the lack of polycyclic aromatic hydrocarbon emission from Class 0 and I sources, a high polycyclic aromatic hydrocarbon abundance is not realistic.

In reality, any small grains that are present in protostellar envelopes could be coupled to the ions ([Guillet et al. 2007; Godard et al. 2019](#)). In turn, this also results in a larger fraction of the parameter space having a shock length of $< 100 \text{ AU}$. However, the full treatment of grain dynamics in *C*-type shocks would have to include a separate fluid for each grain size ([Ciolek & Roberge 2002; Guillet et al. 2007](#)) with grain-grain interactions such as vaporization, shattering, and coagulation ([Guillet et al. 2009, 2011](#)). This is beyond the scope of this work.

Appendix E.2: *CJ*-type shocks

With decreasing magnetic field strength, the decoupling between ions and neutrals becomes weaker. The motion of the neutral velocity is no longer completely dominated by the ion-neutral drag, but also by the viscous stresses and thermal pressure gradient ([Godard et al. 2019](#)). The resulting *CJ*-shock structure is presented in left part of Fig. E.1 for $B \sim 160 \mu\text{G}$. All other parameters are kept at their fiducial value (see Table 1). Initially, the drag of the ions slows down the neutrals, increasing the gas temperature to $\sim 50 \text{ K}$. However, here the viscous stresses and

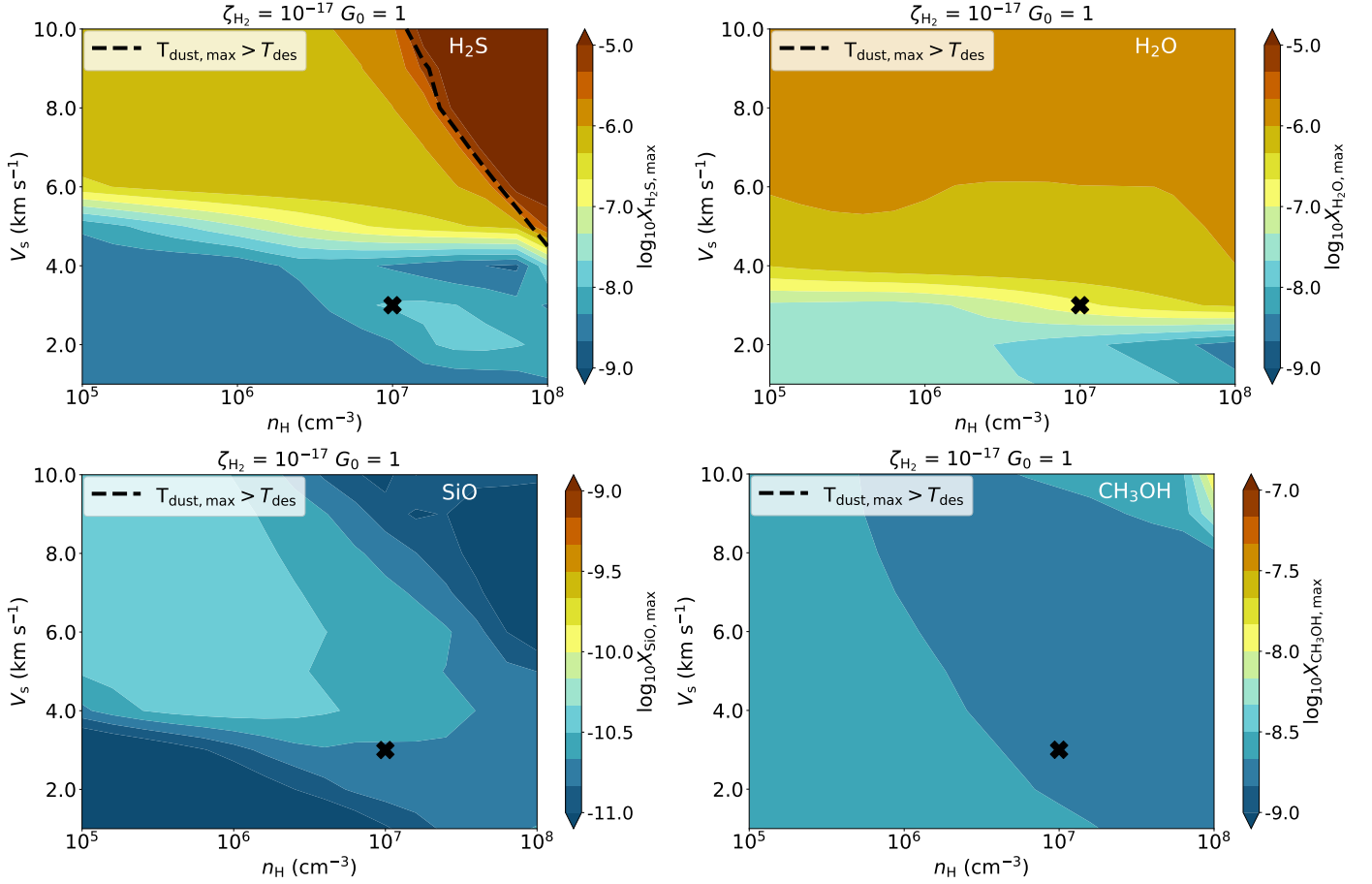


Fig. D.1. Similar figure as Fig. 4, but now for H_2S (top left), H_2O (top right), SiO (bottom left), and CH_3OH (bottom right). All other physical parameters are kept constant to the fiducial values and listed on top of the figure. The black cross indicates the position of the fiducial model. The dashed black line shows the ice line, i.e., where 50% of the ice is thermally desorbed into the gas in the shock.

thermal pressure take over, resulting in a jump in temperature and density as the neutral fluid jumps across the sonic point (Gordard et al. 2019). Afterwards, the ions keep exerting a small drag force on the neutrals until the fluids recouple at about ~ 80 AU. Since the heating induced by this drag is lower than the cooling of the gas, the shock gradually cools down. The abundances of SO and SO_2 do show an increase in the CJ -shock, see right panel of Fig. E.1. Here, SO is predominantly formed through Reaction (2) with SH coming from the green route of Fig. 5.

Given that the strength of the magnetic field on inner envelope scales might be high enough to create CJ -type shocks (Hull et al. 2017), these type of shocks are very relevant. However, in order to compute CJ -type shocks over the entire parameter space considered in this paper, the Paris-Durham shock code has to be tweaked to our specific parameter space. This is beyond the scope of this work. Nevertheless, as shown in Fig. E.1, the chemical formation routes of Fig. 5 are still valid for CJ -type shocks since similar physical conditions are attained.

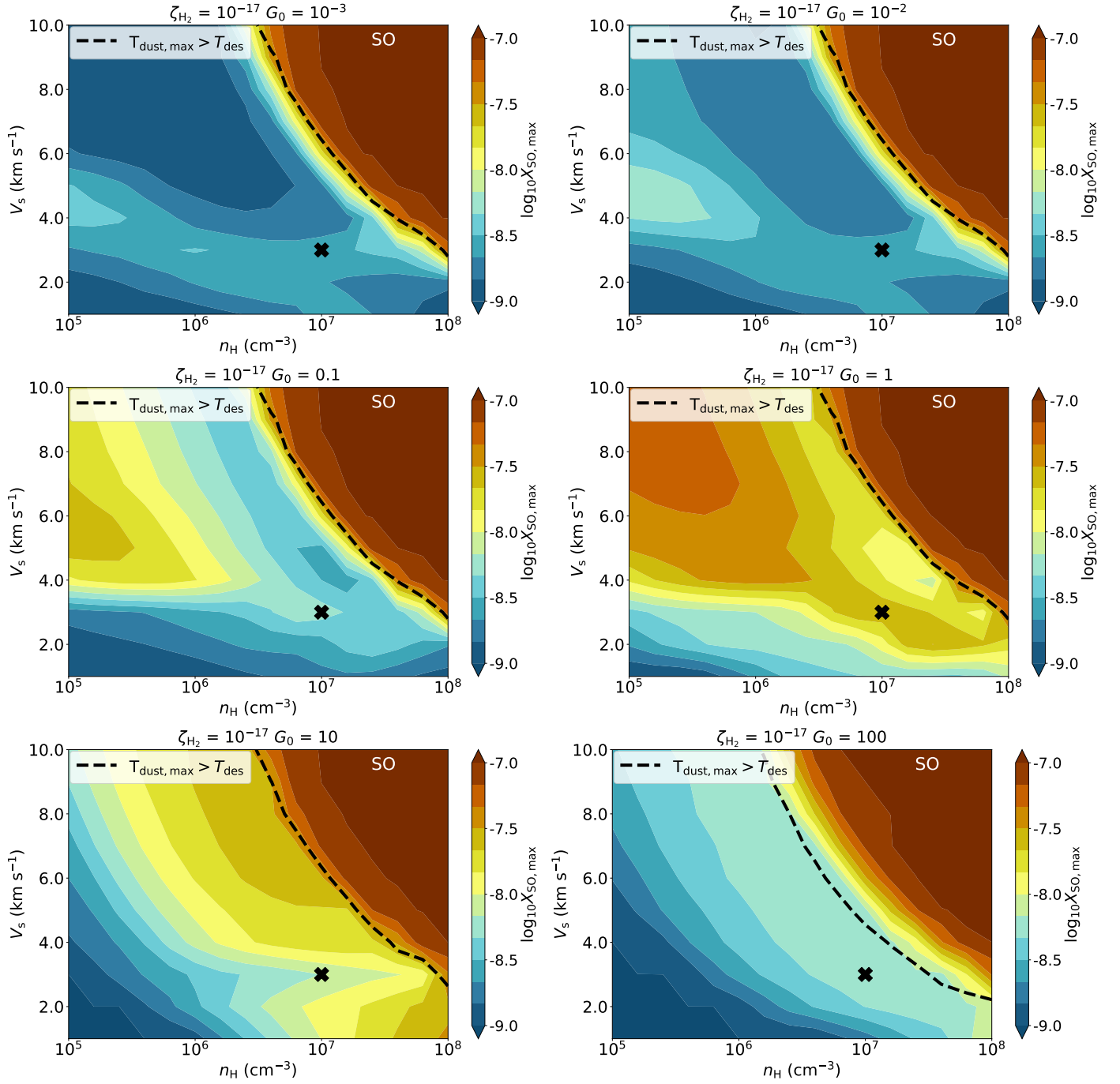


Fig. D.2. Maximum abundance reached (in color) of SO in shock models as function of initial n_H and V_s for $G_0 = 10^{-3}$ (top left), $G_0 = 10^{-2}$ (top right), $G_0 = 0.1$ (middle left), $G_0 = 1$ (middle right), $G_0 = 10$ (bottom left), and $G_0 = 100$ (bottom right). The black cross indicates the position of the fiducial model. The dashed black line shows the ice line, i.e., where 50% of the ice is thermally desorbed into the gas in the shock.

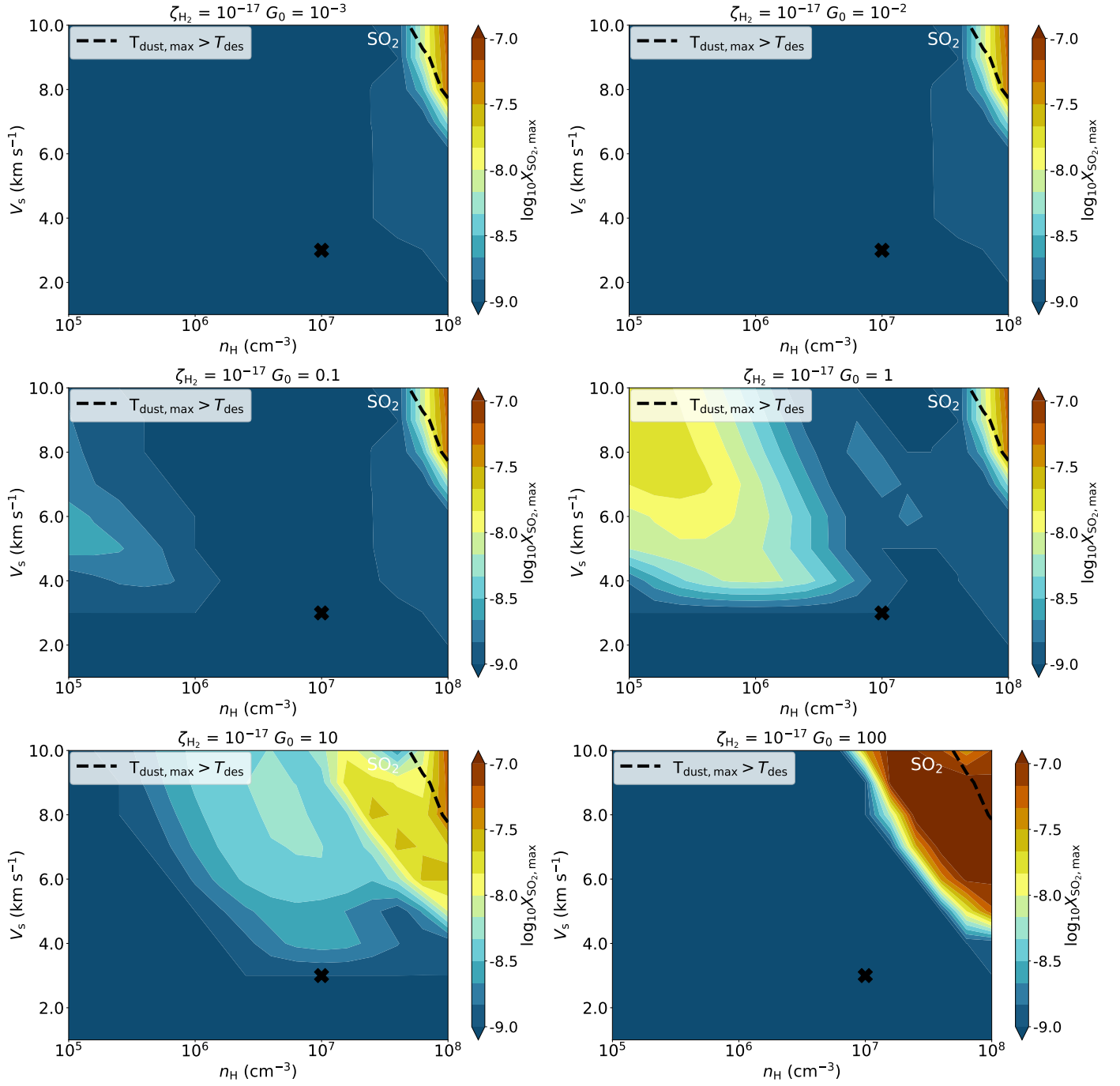


Fig. D.3. Maximum abundance reached (in color) of SO_2 in shock models as function of initial n_{H} and V_s for $G_0 = 10^{-3}$ (top left), $G_0 = 10^{-2}$ (top right), $G_0 = 0.1$ (middle left), $G_0 = 1$ (middle right), $G_0 = 10$ (bottom left), and $G_0 = 100$ (bottom right). The black cross indicates the position of the fiducial model. The dashed black line shows the ice line, i.e., where 50% of the ice is thermally desorbed into the gas in the shock.

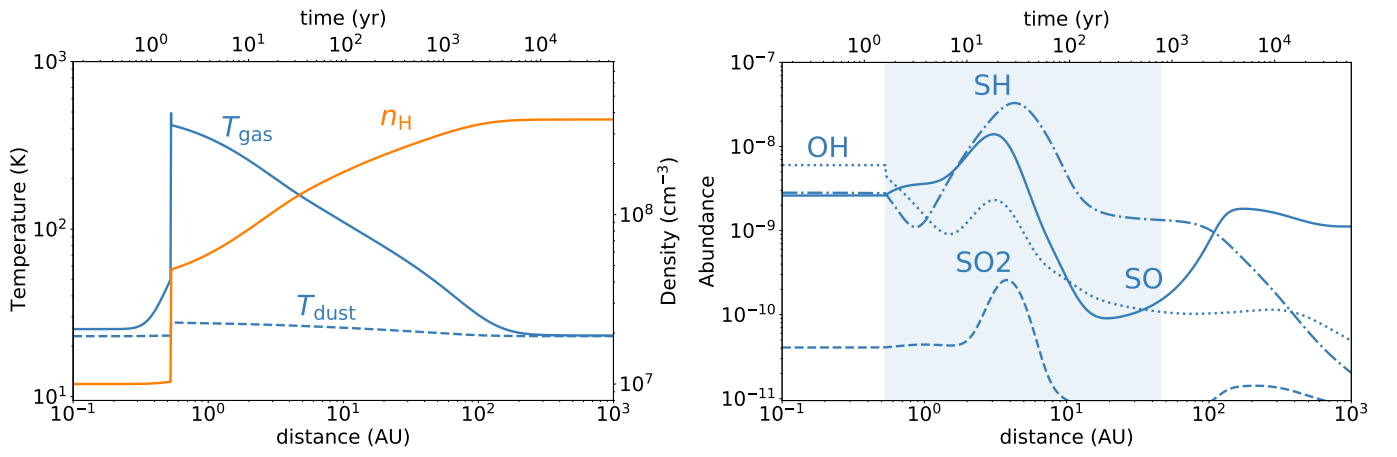


Fig. E.1. Same as Fig. 2 but for a *CJ*-type shock with $B \sim 160 \mu\text{G}$. All other parameters are kept at their fiducial value (see Table 1). In the right panel, the region where $T_{\text{gas}} > 50$ K is indicated with the shaded blue region.