First results of the Herschel key program "Dust, Ice and Gas In Time" (DIGIT): Dust and gas spectroscopy of HD100546


DOI: 10.1051/0004-6361/201014674

Publication date
2010

Document Version
Final published version

Published in
Astronomy & Astrophysics

Citation for published version (APA):
First results of the Herschel key program “Dust, Ice and Gas In Time” (DIGIT): Dust and gas spectroscopy of HD 100546*


(Affiliations are available in the online edition)

Received 31 March 2010 / Accepted 28 April 2010

ABSTRACT

Context. We present far-infrared spectroscopic observations, taken with the Photodetector Array Camera and Spectrometer (PACS) on the Herschel Space Observatory, of the protoplanetary disk around the pre-main-sequence star HD 100546. These observations are the first within the DIGIT Herschel key program, which aims to follow the evolution of dust, ice, and gas from young stellar objects still embedded in their parental molecular cloud core, through the final pre-main-sequence phases when the circumstellar disks are dissipated.

Aims. Our aim is to improve the constraints on temperature and chemical composition of the crystalline olivines in the disk of HD 100546 and to give an inventory of the gas lines present in its far-infrared spectrum.

Methods. The 69 μm feature is analyzed in terms of position and shape to derive the dust temperature and composition. Furthermore, we detected 32 emission lines from five gaseous species and measured their line fluxes.

Results. The 69 μm emission comes either from dust grains with ~70 K at radii larger than 50 AU, as suggested by blackbody fitting, or it arises from ~200 K dust at ~13 AU, close to the midplane, as supported by radiative transfer models. We also conclude that the forsterite crystals have few defects and contain at most a few percent iron by mass. Forbidden line emission from [C II] at 157 μm and [O I] at 63 and 145 μm, most likely due to photodissociation by stellar photons, is detected. Furthermore, five H2O and several OH lines are detected. We also found high-J rotational transition lines of CO, with rotational temperatures of ~300 K for the transitions up to J = 22–21 and T ~ 800 K for higher transitions.

Key words. stars: individual: HD 100546 – infrared: general – stars: pre-main sequence – techniques: spectroscopic

1. Introduction

Circumstellar disks around young stars are the birthplaces of planetary systems. To understand planet formation, it is vital to study the processes that govern the evolution of gas and dust in these disks. PACS provides unique information in this field through observations of far-infrared (IR) solid-state features which are particularly sensitive to temperature and elemental composition. Moreover, PACS is well suited to study the warm gas of a few 100 K, complementary to the hot gas probed in near-IR and the cold gas observed at millimeter wavelengths.

An intensively studied pre-main-sequence star is the Herbig B9.5Vn star HD 100546. While signs of ongoing accretion indicate the star’s infancy (Deleuil et al. 2004), the estimated age of 10 Myr makes it unusually old for a star with a disk (van den Ancker et al. 1997). HD 100546 is nearby (103 pc) and optical/near-IR scattered light imaging has revealed a wealth of structures in the disk (e.g., Grady et al. 2001; Augereau et al. 2001; Ardila et al. 2007). Based on the mid- to near-IR excess ratios, Bouwman et al. (2003) suggested an inner cavity in the disk. Later observations confirmed this gap (e.g., Grady et al. 2005; Benisty et al. 2010). This remarkable feature and the longevity of the HD 100546 disk may point to a young planet in the inner 10 AU, making it a prime target for detailed studies.

The system of HD 100546 has a rich 2.4–180 μm spectrum as observed with ISO (Malfait et al. 1998), showing a striking similarity with that of the comet Hale-Bopp (Crovisier et al. 1997). Juhasz et al. (2010) investigated the Spitzer spectrum of the source. Strong forsterite emission is observed in both ISO and Spitzer data. A continuum fit indicated crystalline dust at 210 K and 50 K (Malfait et al. 1998).

The ISO spectrum shows a strong line of [O I] 63 μm and a weaker [C II] 158 μm line. In the near-IR, Brittain et al. (2009) and van der Plas et al. (2009) observed CO ro-vibrational emission lines with a rotational temperature of ~1000 K and found no CO gas in the inner dust cavity. Observations of optical [O I] and Balmer line emission, on the other hand, demonstrate that the inner disk is not completely devoid of gas (Acke & van den Ancker 2006). At submillimetre wavelengths, Panić et al. (2010) have...
detected pure rotational lines of CO up to $J = 7$–$6$ probing a warm (60–70 K) layer in the outer disk (~100 AU). Information on the intermediate temperatures is still lacking.

Here we present an analysis of narrow features (about 1 μm) in the far-IR spectrum of HD 100546 over the full PACS range that was obtained within the Herschel key program “Dust, Ice and Gas In Time” (DIGIT).

2. Observation and data reduction

We used the spectral energy distribution (SED) mode, covering the wavelength range from 55 to 210 μm (see Poglitsch 2010, for details of the instrument and calibration). Four up/down scans of the grating for two nod positions were performed with a chopper throw of 1′. Our data were processed in HIPE 3.0, provided by the Herschel science center, up to level 1. We did not apply the integration ramp linearity, cross-talk and response drift corrections as these were not yet scientifically validated.

After level 1 processing, we extracted the spectra for the two nod positions separately from the central spaxel (spatial element). The wavelength–dependent point spread function (PSF) correction was taken from the PACS calibration document PICC-KL-TN-041. We first uniformly rebinned the spectra, using an over-sampling factor of 1 relative to the spectral resolution of the instrument and an up-sampling factor of 2, before combining the spectra of the two nod positions. The current flux calibration and relative spectral response function (RSRF) were derived from ground-based measurements. The correction factors, determined from the first in-space calibrations were applied (for details see the PACS calibration document PICC-KL-TN-041). We estimate that the absolute flux calibration is accurate within 30%. The RSRF is valid for spectral features not much wider than about 1 μm. For broader features, the RSRF is not yet well calibrated. Therefore, we will focus on spectral features narrower than about 1 μm. The wavelength calibration is accurate down to the spectral resolution as can be seen from the gas-phase lines, most of which are found within 0.02 μm of their laboratory wavelength.

3. Results and discussion

The PACS data confirm the tentative ISO detection of the 69 μm forsterite band first reported by Malfait et al. (1998) and deliver a band profile of very high quality (see Fig. 1). The feature is characterized by a signal-to-noise ratio of about 10. In addition, we find a wealth of narrow molecular gas lines in the spectrum of HD 100546, coming from CO, H₂O, and OH, and [C II] and [O I] fine-structure lines (see Table 1).

In this section we present an analysis of the 69 μm forsterite peak in terms of dust temperature and composition. In addition, we provide a compilation of the detected gas lines in the far-IR spectrum of HD 100546 and derive the CO-rotational temperature. A full spectral decomposition, including a search for shallow and wide features (e.g. ice and hydrated silicate features), will be presented in a follow-up paper.

3.1. Forsterite

The “69 μm” band is a characteristic signature of forsterite – the Mg-rich end member of olivines (Henning 2010). In order

$$F_{68-71 \mu m}(\lambda) = F_{cont}(\lambda) + \sum_{i=1}^{5} w_i \cdot \kappa(T_{dust}^i, \lambda) \cdot B_\lambda(\lambda, T_{dust}^i).$$

Here $F_{cont}$ is the local continuum, and the $\kappa$ values are the mass absorption coefficients, computed from the optical constants of the Suto et al. (2006) forsterite sample, at the different dust temperatures $T_{dust}^i$. Finally, the $w_i$ values are the relative weights of the flux at the different temperatures. In addition, we had to

Fig. 1. The continuum-subtracted forsterite emission at 69 μm (histogram) overlaid with the best-fit model (red line) which is given by Eq. (1). The contributions of the different temperatures are indicated. The dotted vertical line locates the peak position of 70 K pure forsterite and the dashed one the position in the PACS spectrum, fitted with a Lorentz profile, which is not shown here.

Based on the peak position of the band, we can immediately conclude that the olivines cannot contain more than 3–4% iron as this would cause the peak to appear at significantly longer wavelengths than observed. The PACS observations provide much better constraints on the iron content than the shorter-wavelength Spitzer data (Juhász et al. 2010) because silicate bands in that spectral range are less sensitive to a change in Mg/Fe ratio. Furthermore, the small $FWHM$ of the band profile indicates crystals with few defects (Koike et al. 2010). The small $FWHM$ also shows that the dust emission comes from a narrow range of temperatures.

Determining the temperature of the dust that produces the 69 μm feature from intrinsic properties of this band is more difficult because even a small amount of iron could balance a lower temperature. Assuming pure forsterite results in a dust temperature of about 200 K (Fig. 2). To constrain the amount of colder dust, we fitted the observed profile with a weighted sum of laboratory data (Suto et al. 2006) at different temperatures, ranging from 50–300 K (Fig. 1). The measured flux $F_{68-71 \mu m}$ is given by

Table 1. Peak wavelengths and line fluxes for the identified gas lines between 60 and 190 \( \mu m \).

<table>
<thead>
<tr>
<th>Species</th>
<th>Transition</th>
<th>Peak ( \mu m )</th>
<th>Linerlux ( 10^{-17} \text{[W/m}^2\text{]} )</th>
</tr>
</thead>
<tbody>
<tr>
<td>[O i]</td>
<td>( ^{3}P_{1}−^{3}P_{2} )</td>
<td>63.19</td>
<td>554.37 ± 5.0</td>
</tr>
<tr>
<td>[O i]</td>
<td>( ^{3}P_{2}−^{3}P_{1} )</td>
<td>145.54</td>
<td>55.70 ± 1.3</td>
</tr>
<tr>
<td>[C ii]</td>
<td>( ^{2}P_{2}/^{2}P_{1} )</td>
<td>157.75</td>
<td>31.87 ± 1.0</td>
</tr>
<tr>
<td>CO</td>
<td>( J = 14−13 )</td>
<td>186.01</td>
<td>7.42 ± 0.9</td>
</tr>
<tr>
<td>CO</td>
<td>( J = 15−14 )</td>
<td>173.64</td>
<td>11.50 ± 0.8</td>
</tr>
<tr>
<td>CO</td>
<td>( J = 16−15 )</td>
<td>162.82</td>
<td>8.25 ± 0.9</td>
</tr>
<tr>
<td>CO</td>
<td>( J = 17−16 )</td>
<td>153.28</td>
<td>10.49 ± 0.8</td>
</tr>
<tr>
<td>CO</td>
<td>( J = 18−17 )</td>
<td>144.79</td>
<td>9.85 ± 1.0</td>
</tr>
<tr>
<td>CO</td>
<td>( J = 19−18 )</td>
<td>137.21</td>
<td>8.86 ± 0.7</td>
</tr>
<tr>
<td>CO</td>
<td>( J = 20−19 )</td>
<td>130.38</td>
<td>7.10 ± 0.7</td>
</tr>
<tr>
<td>CO</td>
<td>( J = 21−20 )</td>
<td>124.18</td>
<td>5.94 ± 0.6</td>
</tr>
<tr>
<td>CO</td>
<td>( J = 22−21 )</td>
<td>118.59</td>
<td>5.92 ± 0.9</td>
</tr>
<tr>
<td>CO(^{i})</td>
<td>( J = 23−22 )</td>
<td>113.47</td>
<td>4.76 ± 1.2</td>
</tr>
<tr>
<td>CO(^{i})</td>
<td>( J = 24−23 )</td>
<td>108.77</td>
<td>10.56 ± 1.3</td>
</tr>
<tr>
<td>CO(^{i})</td>
<td>( J = 25−24 )</td>
<td>100.47</td>
<td>12.71 ± 2.4</td>
</tr>
<tr>
<td>CO(^{i})</td>
<td>( J = 27−26 )</td>
<td>96.78</td>
<td>10.20 ± 1.4</td>
</tr>
<tr>
<td>CO(^{i})</td>
<td>( J = 28−27 )</td>
<td>93.35</td>
<td>10.00 ± 1.0</td>
</tr>
<tr>
<td>CO(^{i})</td>
<td>( J = 29−28 )</td>
<td>90.17</td>
<td>11.83 ± 1.6</td>
</tr>
<tr>
<td>CO(^{i})</td>
<td>( J = 30−29 )</td>
<td>87.19</td>
<td>8.46 ± 1.0</td>
</tr>
<tr>
<td>CO(^{i})</td>
<td>( J = 31−30 )</td>
<td>84.40</td>
<td>4.90 ± 1.5</td>
</tr>
<tr>
<td>OH</td>
<td>( 1/2, 7/2−5/2 )</td>
<td>71.17</td>
<td>9.17 ± 1.4</td>
</tr>
<tr>
<td>OH(^{i})</td>
<td>( 1/2, 7/2−5/2 )</td>
<td>71.22</td>
<td>10.10 ± 1.1</td>
</tr>
<tr>
<td>OH(^{i})</td>
<td>( 3/2, 7/2−5/2 )</td>
<td>84.42</td>
<td>14.96 ± 1.7</td>
</tr>
<tr>
<td>OH(^{i})</td>
<td>( 3/2, 7/2−5/2 )</td>
<td>84.59</td>
<td>14.96 ± 1.5</td>
</tr>
<tr>
<td>OH(^{i})</td>
<td>( 1/2, 5/2−3/2 )</td>
<td>98.73</td>
<td>8.74 ± 1.5</td>
</tr>
<tr>
<td>OH(^{i})</td>
<td>( 1/2, 3/2−1/2 )</td>
<td>163.41</td>
<td>2.60 ± 0.6</td>
</tr>
<tr>
<td>p-H(2)O</td>
<td>( 3_{2}−2_{1} )</td>
<td>90.00</td>
<td>14.32 ± 1.2</td>
</tr>
<tr>
<td>o-H(2)O(^{i})</td>
<td>( 4_{1}−3_{2} )</td>
<td>113.53</td>
<td>3.64 ± 0.9</td>
</tr>
<tr>
<td>o-H(2)O</td>
<td>( 3_{2}−2_{1} )</td>
<td>136.48</td>
<td>4.52 ± 0.7</td>
</tr>
<tr>
<td>p-H(2)O</td>
<td>( 3_{1}−2_{0} )</td>
<td>138.54</td>
<td>3.17 ± 0.7</td>
</tr>
<tr>
<td>o-H(2)O</td>
<td>( 2_{1}−1_{0} )</td>
<td>179.54</td>
<td>5.3 ± 0.8</td>
</tr>
</tbody>
</table>

Notes. Only the formal errors are given, not including systematic errors due to the PACS spectral response function. (\(^{i}\) Debledoned lines, CO not used in rotational diagram.

specify a dust shape distribution in order to calculate the mass absorption values. We used the DHS model (distribution of hollow spheres, Min et al. 2003) with a grain size of 0.1 \( \mu m \), which has been found to be a good representation of observed silicate profiles (e.g., Juhász et al. 2010).

Under the assumption of pure forsterite, no more than a few percent of the dust emission can be coming from the 50 K component. This could be up to 9%, if grains of \( \sim 5 \mu m \) radius are assumed. Such large grains can only exist in the deeper layers of the disk, where they are invisible to Spitzer, as Juhász et al. (2010) rule out grains \( \geq 1 \mu m \). The bulk of the emission, under the assumption of pure forsterite, comes from \( \sim 150−200 \) K.

However, a temperature of 150 to 200 K for the 69 \( \mu m \) band would lead to tremendously higher fluxes in the 20 to 40 \( \mu m \) region than observed by ISO and Spitzer. Additionally, the predicted relative strength between the 24- and 33 \( \mu m \) forsterite bands would be inverted compared to the observations. In fact, temperatures of only 70 K (Bouwman et al. 2003) or even 50 K (Malfait et al. 1998) were derived from the ISO data.

These results can be reconciled in two ways. First, the dust producing the 69 \( \mu m \) feature is at 70–50 K. The shift from the band position in the 70–50 K laboratory data to the one observed by PACS could be caused by an admixture of iron. Given the information in Fig. 3, we need at least 2 but no more than 3% for such a shift, considering that the data were taken at \( \sim 300 \) K.

An alternative explanation is based on the optical depth of the disk which decreases with increasing wavelength. The \( \tau = 1 \) surface is deep into the disk at 69 \( \mu m \), even reaching the midplane (Mulders, in prep.). The disk of HD 100546 is a special case in this respect: because of its large inner hole, the midplane temperature at 13 AU is much higher (\( \sim 150–200 \) K) than in disks without a hole so that the far-infrared features can be produced in heated layers close to the midplane from which the radiation at 70 \( \mu m \) can escape. In the most extreme case the emission could arise from pure forsterite of 200 K temperature, located in the midplane at 13 AU.

In this scenario the location of the forsterite near the inner edge of the disk supports the crystallization by collisional cascades or shocks induced by the suggested giant planet.

3.2. Gas lines

We identified 32 gas lines (Table 1) from the ISO line lists.\(^{2}\) CO lines up to the \( J = 31−30 \) transition were detected as well as H\(2\) and OH lines. Selected molecular lines are shown in Fig. 4. Furthermore, we confirm the ISO detections of [O I] and [C II] fine-structure lines (Lorenzetti et al. 2002). The integrated line fluxes, obtained from fitted Lorentz profiles, are in Table 1. The [O I] line fluxes agree within 30% with those seen by ISO (Lorenzetti et al. 2002), while the [C II] line flux is about a factor of 3 weaker. Significant flux outside the pointsource PSF is detected for the [C II] line, which is not seen for any other line, suggesting that extended emission dominates the larger ISO beam. Through comparison of the flux ratios [O I]\(_{3}\)/[O I]\(_{1}\) and [O I]\(_{3}/[C II]_{18}\) with the models from Kamp et al. (2010) a gas mass of \( \sim 10^{-3} M_\odot \) is derived.

\(^{2}\) http://www.mpe.mpg.de/ISO/linelists/index.html
According to Brittain et al. (2009) the 4.7 μm CO fundamental emission shows both a UV fluorescent and a thermal component. The latter was fitted with a column density of at least \(2.3 \times 10^{15} \text{ cm}^{-2}\) (for discussion, see Brittain et al. 2009). The submillimeter data give a typical CO column density of \(10^{15} \text{ cm}^{-2}\) (Panić et al. 2010). From Fig. 5, assuming that the emission originates from the area between 13 and 100 AU, we derive column densities of \(10^{16}\) to \(10^{17} \text{ cm}^{-2}\) in both models. These values indicate that PACS probes the part of the disk between the near-IR and submillimeter data.

4. Conclusions

The PACS data provide the strongest constraints yet on the composition of olivines in a protoplanetary disk. We found the olivines to be extremely iron-poor (less than 3–4% iron). Our observations can be modeled with dust at \(-70^\circ\text{K}\), consistent with ISO results, but requiring at least 2% iron. On the other hand the determination of the temperature of the forsterite based on intrinsic features of the 69 μm band instead of using a continuum analysis offers a second option: the emission may emerge from pure forsterite at 200 K and 13 AU close to the midplane. This region is optically thick for shorter wavelengths.

The far-infrared spectrum of HD 100546 contains a wealth of molecular gas lines including CO, H$_2$O, and OH. We found CO transitions up to \(J = 31-30\) which arise from gas in the temperature range of 300 to 800 K and appear to sample gas in the surface layers of the disk at temperatures and surface densities intermediate to those probed by submillimeter and near-infrared CO measurements.

Acknowledgements. Support for this work, part of the Herschel open time key program, was provided by NASA through an award issued by the Jet Propulsion Laboratory, California Institute of Technology. G. Meeus is partly supported by Spanish grant AYA 2008-01727. The authors thank the referee C. Grady for helpful comments that improved the paper.

References

Henning, Th. 2010, ARA&A, in press

Page 5 is available in the electronic edition of the journal at http://www.aanda.org