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Study of the aluminium content in AGB winds using ALMA

Indications for the presence of gas-phase $(\text{Al}_2\text{O}_3)_n$ clusters

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ABSTRACT

Context. The condensation of inorganic dust grains in the winds of evolved stars is poorly understood. As of today, it is not yet known which molecular clusters form the first dust grains in oxygen-rich ($\text{C/O} < 1$) asymptotic giant branch (AGB) winds. Aluminium oxides and iron-free silicates are often put forward as promising candidates for the first dust seeds.

Aims. We aim to constrain the dust formation histories in the winds of oxygen-rich AGB stars.

Methods. We obtained Atacama Large Millimeter/sub-millimeter array (ALMA) observations with a spatial resolution of 120×150 mas tracing the dust formation region of the low mass-loss rate AGB star, R Dor, and the high mass-loss rate AGB star, IK Tau. We detected emission line profiles of AlO, AlOH, and AlCl in the ALMA data and used these line profiles to derive a lower limit of atomic aluminium incorporated in molecules. This constrains the aluminium budget that can condense into grains.

Results. Radiative transfer models constrain the fractional abundances of AlO, AlOH, and AlCl in IK Tau and R Dor. We show that the gas-phase aluminium chemistry is completely different in both stars with a remarkable difference in the AlO and AlOH abundance stratification. The amount of aluminium locked up in these three molecules is small, $\leq 1.1 \times 10^{-7}$ w.r.t. H_2 , for both stars, i.e. only $\leq 2\%$ of the total aluminium budget. An important result is that AlO and AlOH, which are the direct precursors of alumina (Al_2O_3) grains, are detected well beyond the onset of the dust condensation, which proves that the aluminium oxide condensation cycle is not fully efficient. The ALMA observations allow us to quantitatively assess the current generation of theoretical dynamical-chemical models for AGB winds. We discuss how the current proposed scenario of aluminium dust condensation for low mass-loss rate AGB stars within a few stellar radii from the star, in particular for R Dor and W Hya, poses a challenge if one wishes to explain both the dust spectral features in the spectral energy distribution (SED) in interferometric data and in the polarized light signal. In particular, the estimated grain temperature of Al_2O_3 is too high for the grains to retain their amorphous structure. We advocate that large gas-phase $(\text{Al}_2\text{O}_3)_n$ clusters ($n > 34$) can be the potential agents of the broad $11 \mu\text{m}$ feature in the SED and in the interferometric data and we propose potential formation mechanisms for these large clusters.

Key words. stars: AGB and post-AGB – stars: individual: IK Tau and R Dor – astrochemistry – instrumentation: interferometers – stars: mass-loss – circumstellar matter

1. Introduction

It is well known from observations that low and intermediate mass stars ($0.8 M_\odot < M < 8 M_\odot$) develop a low-velocity ($v \sim 5\text{--}15 \text{ km s}^{-1}$) stellar wind at the end of their life during the asymptotic giant branch (AGB) phase. At a mass-loss rate between 10^{-8} to $10^{-4} M_\odot/\text{yr}$, material is stripped away from the stellar surface. Convection-induced pulsations lift material to greater heights where the temperature is low enough for gas to condense into dust grains (Habing & Olofsson 2003; Freytag et al. 2017). Radiation pressure on these grains is thought to be the main trigger for the onset of the stellar wind. While theoretical simulations based on this scenario predict stellar wind

properties in broad agreement with observations of carbon-rich ($\text{C/O} > 1$) AGB stars, more fine-tuning of the models is required for oxygen-rich (O-rich) AGB stars. Indeed, in these O-rich environments, the first dust seeds thought to form close to the stellar surface are metal oxides and pure Fe-free silicates such as Al_2O_3 , SiO_2 , TiO_2 , Mg_2SiO_4 , and MgSiO_3 (Gail & Sedlmayr 1999). While their low near-infrared extinction efficiency prevents them from sublimating, it means that these glassy condensates make a negligible contribution to the radiative acceleration. The formation of both carbon and silicate grains (Höfner & Andersen 2007) or scattering of stellar radiation by micron-sized ($\sim 0.3 \mu\text{m}$) Fe-free silicates (Höfner 2008) have been proposed as possible alternatives to solve this acceleration

deficit. After its postulation in 2008 by Höfner et al., the latter scenario got more and more support from observations; the first such observations were reported by Norris et al. (2012) using VLT/NACO data of three low mass-loss rate AGB stars (R Dor, W Hya, and R Leo). The mid-infrared interferometric observations of the semi-regular AGB star RT Vir with the MIDI instrument at VLTI also lent support to the presence of iron-free silicates between 2 and 3 stellar radii (Sacuto et al. 2013). Recent polarized light data obtained with the VLT/SPHERE instrument offer us another way to look at the inner wind chemical and morphological structure. Again, these data prove the presence of grains close to the star, albeit it is not evident to put stringent constraints on the grain size (Khoury et al. 2016; Ohnaka et al. 2016). However, neither the NACO or the SPHERE data can pinpoint the exact chemical constitution of these micron-sized transparent grains.

Considering the abundance of molecular species and the temperatures at which they can exist in solid phase, thermodynamic equilibrium studies show that possible candidates for the first nucleation seeds are SiO, SiO₂, MgO, Fe, Al₂O₃, TiO, and TiO₂ (Jeong et al. 2003). However, taking into account the thermodynamic properties, MgO nucleation is thought to be completely negligible in O-rich AGB stars. Although Al₂O₃ dust is stable at very high temperatures, gaseous Al₂O₃ has a low abundance, calling into question its role as the first candidate. For a long time, TiO and TiO₂ were considered to be the best candidates as primary condensates, since the nucleation rate of the more abundant SiO was thought to peak around 600 K (as based on the thermodynamic model of Schick 1960, using a single laboratory determination of the SiO vapour pressure), which is significantly lower than the typical temperatures at the inner edge of the dust shells. However, more recent laboratory experiments by Nuth & Ferguson (2006) and Gail et al. (2013) have proved that the onset of SiO nucleation starts at much higher temperatures than was previously found, although the calculated SiO condensation temperatures are still ~100 K lower than the observed temperatures of around 880 K. Recent ALMA observations of TiO and TiO₂ transitions in Mira (*o* Ceti) show these two molecules are abundant in the gas phase, implying titanium oxides may not be important first condensates as previously thought (Kamiński et al. 2017).

The A-X infrared bands of AlO have been detected on a fair fraction of AGB stars (Banerjee et al. 2012). Rotational transitions of AlO and AlOH were reported for the oxygen-rich supergiant VY CMa by Tenenbaum & Ziurys (2009, 2010). Kamiński et al. (2016) have reported on the detection of several AlO lines in *o* Ceti with ALMA. The remarkable integrated-intensity map of the $N = 6-5$ AlO transition at 230 GHz indicates the molecule is present within $4R_*$ from the stellar surface and shows a patchy distribution. The molecule AlOH was also potentially detected in *Herschel* and APEX data, while AlH and atomic Al I lines were detected in optical spectra. Takigawa et al. (2016) have reported on ALMA long-baseline observations of several molecules, including AlO, in W Hya. Finally, De Beck et al. (2017) has studied eight oxygen-rich AGB stars; AlO could clearly be identified in two of these stars and for four other targets tentative detections were shown.

Another quest focusses on the potential chemical differentiation between grains formed in low mass-loss rate versus high mass-loss rate O-rich AGB stars. Observations indicate that low mass-loss rate stars form primarily dust that preserves the spectral properties of Al₂O₃, and stars with higher mass-loss rates form dust with properties of warm

silicate oxides (Karovicova et al. 2013)¹. It is absolutely unknown if the silicates form via a heteromolecular homogeneous nucleation process consuming Mg, SiO, and H₂O molecules (Goumans & Bromley 2012) or if the silicates form on top of the alumina grains (heterogeneous growth), hence gradually accelerating the wind.

A promising way to unravel the coupling between nucleation and wind generation is by studying at high spatial resolution the molecules contributing to the dust formation. This possibility is offered by ALMA (the Atacama Large Millimeter/submillimeter array). We were granted ALMA Cycle 2 observing time (project 2013.1.00166.S) to investigate the molecular content in the inner wind region of the low mass-loss rate semi-regular AGB star R Dor ($\dot{M} \sim 9 \times 10^{-8} M_{\odot}/\text{yr}$) and the high mass-loss rate Mira star IK Tau ($\dot{M} \sim 4.5 \times 10^{-6} M_{\odot}/\text{yr}$). We obtained the first ALMA spectral scan for O-rich AGB stars in band 7 at a spatial resolution of $\sim 0''.12 \times 0''.15$. The full data scan will be presented in Decin et al. (in prep.). In this paper, we wish to advance our knowledge of aluminium-bearing molecules, clusters, and grains in AGB winds by considering the two stars above and another low mass-loss rate AGB star, W Hya. Some properties of the three targets are introduced in Sect. 2.1. The ALMA observations of R Dor and IK Tau are presented in Sect. 2.2. Since the aim is to correlate the gaseous aluminium content with the amount of aluminium locked up into grains, we briefly summarize the most important dust properties of the three targets in Sect. 2.3. The results of the ALMA data analysis are presented in Sect. 3. The molecular abundance fractions of AlO, AlOH, and AlCl are derived, and the aluminium fraction incorporated in gas-phase and solid-state species is presented. We show that observations indicate the potential presence of large gas-phase (Al₂O₃)_{*n*} clusters in Sect. 4 and discuss how these large clusters can be formed. In addition, we calculate the emission of very small (Al₂O₃)_{*n*} ($n \leq 4$) clusters. The conclusions are summarized in Sect. 5.

2. Observations

2.1. Target selection

To study the intricate coupling between gas and dust chemistry and the impact of the formation of some type of dust species on the wind dynamics, for our ALMA observations we selected the two O-rich AGB stars that best represent the classes of low mass-loss rate and high mass-loss rate O-rich AGB stars, R Dor and IK Tau. The reasons for their selection are their proximity (60 pc and 260 pc, respectively), their rich infrared and submillimeter spectra with signatures of many dust species and molecules, and the fact that both targets are very well studied with various instruments. Each target has a very different dust emission spectrum (see Sect. 2.3); the spectrum of IK Tau is dominated by amorphous olivine dust (magnesium-iron silicates (Mg_{2-*x*}Fe_{*x*})SiO₄, with $0 \leq x \leq 2$), while the infrared spectrum of R Dor does not show the presence of olivine dust but species such as corundum (an aluminium oxide, Al₂O₃), the aluminium-calcium bearing silicate gehlenite, the magnesium-aluminium member spinel, and CO₂ gas have been detected (Heras & Hony 2005; Khouri 2014). This supports the hypothesis that the dust condensation sequence in the low mass-loss rate object R Dor has experienced a process called “freeze-out” (Tielens 1990),

¹ This differentiation does not mean that no silicates are present in low mass-loss rate stars. Their presence might stay unnoticed in the spectral energy distribution (SED) owing to, for instance, their glassy character.

where the sequence is not completed and stops at an intermediate product when the wind density drops below the density required for the next step; in this particular case the freeze-out occurs after the formation of aluminium oxide with only a small amount of silicon condensing into dust grains.

In IK Tau the formation of silicon-bearing dust species seems to be much more efficient. This can happen through either a one-step process, i.e. the growth of micron-sized Fe-free silicates close the central star, or a two-step process might take place, i.e. the growth of aluminium oxides around a few stellar radii (R_\star), after which the grains were coated with silicates around ~ 10 stellar radii.

R Dor is the closest AGB star to our Sun with a gas mass-loss rate value of $\sim 0.9\text{--}2 \times 10^{-7} M_\odot/\text{yr}$ and an expansion velocity of only $\sim 5.7 \text{ km s}^{-1}$ (Schöier et al. 2004; Khouri 2014; Maercker et al. 2016; Van de Sande et al. 2017). This star is a semi-regular variable with two pulsation modes that have periods of 332 days and about 175 days; the latter mode has a much smaller amplitude (Bedding et al. 1998). The angular diameter of this star is ~ 60 mas (Bedding et al. 1997; Norris et al. 2012). From an analysis of *Herschel* and sub-millimeter data, Khouri (2014) and Van de Sande et al. (2017) have derived that some 90% of atomic silicon is locked up in gaseous SiO and the remainder in silicates. Data from VLTI/NACO and SPHERE/ZIMPOL support the presence of a close halo of large transparent grains ($\sim 0.3 \mu\text{m}$) at $\sim 1.5\text{--}2 R_\star$ (Norris et al. 2012; Khouri et al. 2016).

IK Tau is a high mass-loss rate Mira-type AGB star with a gas mass-loss rate of $\sim 4.5 \times 10^{-6} M_\odot/\text{yr}$, an expansion velocity of 17.7 km s^{-1} (Decin et al. 2010a; Maercker et al. 2016), and a pulsation period of ~ 470 days (Wing & Lockwood 1973). The angular diameter is estimated around 20 mas (Decin et al. 2010a). Currently, a dozen different molecules and some of their isotopologues have been discovered in IK Tau, including CO, HCN, SiO, SiS, SO, SO_2 , NaCl, NS, NO, HCO^+ , H_2CO , PO, PN, and H_2S (Milam et al. 2007; Kim et al. 2010; Decin et al. 2010a,b; De Beck et al. 2013, 2015; Velilla Prieto et al. 2017). Some molecules including SiO and SiS are depleted in the circumstellar envelope, possibly owing to condensation onto dust grains. Other molecules such as HCN demonstrate the impact of pulsation-driven shocks in the inner wind zone which forces the chemistry into a non-equilibrium state (Duari et al. 1999; Cherchneff 2006).

W Hya resembles in many aspects the low mass-loss rate AGB star R Dor. Thanks to its proximity (~ 78 pc; Knapp et al. 2003) and its brightness, R Dor is also well studied with various observational techniques from the visible to the radio. This semi-regular variable has a light curve showing a clear periodicity with period of 389 days (Uttenthaler et al. 2011). The molecular abundance stratification in its circumstellar envelope was studied in detail by Khouri et al. (2014a,b). Based on an extensive grid of CO, SiO, H_2O , and HCN lines (and some of their isotopologues) the thermodynamic structure of the wind was derived and they deduced that about one-third of the silicon atoms is locked up in dust particles. The SED of W Hya is characterized by the same dust emission peaks as R Dor (see Sect. 2.3) and also for this star large scattering grains in a halo close the star are detected via polarized light signal (Norris et al. 2012; Ohnaka et al. 2016). W Hya was also observed by ALMA in band 7 (project 2015.1.01466.S, P.I. A. Takigawa) with a spatial resolution of ~ 40 mas at three selected frequency bands (330.40–331.34 GHz, 342.40–343.34 GHz, and 344.03–345.91 GHz) encompassing the AIO $N = 9\text{--}8$ and

Table 1. ALMA observation parameters for the Al-compounds in IK Tau and R Dor.

Species	Chan ^a (MHz)	Vel ^b (km s^{-1})	Beam (mas \times mas, PA) ^c	σ_{rms} (mJy)
IK Tau				
Continuum	15.2×10^3		180×160 , 27°	0.047
AlCl	1.95	1.7	160×130 , 43°	1.6
AlCl	5.9	5	200×160 , 42°	1.4
AlOH	1.95	1.7	150×140 , -22°	5.0
AlOH	5.9	5	190×170 , -1°	2.4
AIO	1.95	1.7	160×130 , -15°	5.0
AIO	5.9	5	190×170 , -20°	2.6
R Dor				
Continuum	11.6×10^3		150×140 , -5°	0.047
AlCl	0.97	0.9	150×130 , 20°	2.7
AlCl	2.5	2.2	180×180 , 10°	1.4
AlOH	0.97	0.9	170×120 , -48°	3.8
AlOH	2.5	2.2	200×160 , -48°	2.4
AIO	0.97	0.9	160×130 , -26°	4.0
AIO	2.5	2.2	190×160 , -30°	1.8

Notes. ^(a) “Chan” is the channel width for the line cubes, or the total line-free bandwidth (spread between 335–362 GHz) for the continuum. ^(b) “Vel” is the spectral resolution of the image cube. ^(c) PA is the position angle measured from north to east.

AlOH $J = 11\text{--}10$ transition. AIO is detected in their data (Takigawa et al. 2016).

2.2. ALMA observations

2.2.1. ALMA observing strategy

Spectral scans of IK Tau and R Dor were obtained covering the frequencies between 335–362 GHz, using four separate observations each during August–September 2015. The observations and data reduction used the ALMA pipeline for calibration; we followed this by identifying and imaging the line-free continuum, which was then used for self-calibration. The solutions were applied to all data and spectral cubes were made adjusted to constant v_{LSR} . The process is described in more detail in Decin et al. (in prep.); here, we concentrate on the parameters affecting the Al compound cubes. The astrometric accuracy is ~ 17 mas and the flux scale is accurate to $\sim 7\%$. The relative accuracy (e.g. aligning different lines in the same source) is limited only by the signal-to-noise (S/N) and would be < 1 mas for S/N of 200, or 33 mas (about 1 pixel) for a faint 3σ detection of a compact source. The rms noise (σ_{rms}) varies across the band depending on intrinsic atmospheric transmission and the weather at the time of observations. The final image parameters are summarized in Table 1. Owing to the different elevations as seen from ALMA, we achieved a higher resolution and used a smaller pixel size (30 mas versus 40 mas) for R Dor versus IK Tau. For each line, we made cubes at higher and lower spectral resolution (see info in Table 1) using different weighting and spectral averaging. In addition, for AIO, we further smoothed the wide-channel data set with a 500 mas Gaussian kernel. The other lines detected and the sub-mm continuum properties will be discussed in future papers.

2.2.2. Al-bearing species detected with ALMA

Chemical models show that the most likely carriers of aluminium in the wind of oxygen-rich evolved stars are AlH, AlO, AlOH, AlO₂, and AlCl (Gobrecht et al. 2016). AlO₂ is also predicted to be abundant along with some other small aluminium clusters very close to the star. The main isomers of AlO₂ is linear and its rotational lines are not observable. AlO, AlOH, AlH, and Al were detected in α Ceti (Kamiński et al. 2016). AlH does not have a rotational line transition in the targeted ALMA frequency range. The following other three aluminium-bearing molecules are detected in IK Tau and R Dor with ALMA: AlO $N = 9-8$ (rest frequency, ν_{rest} , at 345.457 GHz, lower state energy $E_{\text{low}} = 66$ K), AlOH $J = 11-10$ ($\nu_{\text{rest}} = 346.156$ GHz, $E_{\text{low}} = 83$ K), and AlCl $J = 24-23$ ($\nu_{\text{rest}} = 349.444$ GHz, $E_{\text{low}} = 193$ K).

The transitions of these three molecules are split up by hyperfine components due to the nuclear spin, $I = 5/2$, of aluminium. AlOH and AlCl are closed-shell species with a $^1\Sigma^+$ ground state. AlO is a radical with a $^2\Sigma$ ground state. The rotational levels of this molecule are therefore split by both fine and hyperfine interactions. The net result is that each rotational transition consists of 10–12 favourable, closely spaced, hyperfine components (Tenenbaum & Ziurys 2009, 2010).

Figure 1 shows the flux densities extracted for an aperture size of 320 mas (300 mas) and 800 mas (1000 mas) for IK Tau (R Dor). The channel maps for AlOH in IK Tau and AlO in R Dor are shown in Figs. 2 and 3, respectively. The other channel maps are presented in Appendix A. Table 2 gives the measured properties of each molecular transition, including the velocity width, the shift of the centroid w.r.t. the LSR velocity of the star, the angular width, and the peak flux and integrated flux within the specified velocity interval. The angular width was obtained from the zeroth moment (total intensity) by measuring the mean flux in annuli of 60 (80) mas, centred on the position of R Dor (IK Tau), assumed to be the continuum peak. The angular size was taken as the distance out to $2\sigma_{\text{rms}}$. The measurements were also used to compute the azimuthally averaged flux densities to compare with 1D models solving the non-LTE radiative transfer equation for the specific molecule and with models solving the chemical network by Gobrecht et al. (2016) in Sect. 3. The emission of some molecular transitions shows some irregular morphologies (as described in the next subsections) but there is no obvious preferred direction, so this is a reasonable approximation although not allowing for clumpiness.

IK Tau

- AlO: looking at Fig. A.1, AlO remains at first sight barely detected in IK Tau. This is reflected by the line profile for the 320 mas extraction aperture (Fig. 1). However, enlarging the extraction aperture to 800 mas, a clear signature of AlO emission is seen with a peak flux of 0.19 Jy and integrated line flux of 3.2 Jy km s^{-1} . The SMA observations by De Beck et al. (2017) also show a weak AlO emission feature with an integrated line flux of $2.33 \text{ Jy km s}^{-1}$ for an aperture of $1 \times 1''$. This points towards faint extended AlO emission that is probably genuinely clumpy. To verify this idea, we averaged the AlO emission over five channels and smoothed the emission with a $0''.5$ Gaussian kernel; see Fig. A.2. Albeit the S/N is still low and we caution that the smoothing is on a scale comparable to the maximum recoverable scale (MRS), this figure indeed hints at extended clumpy emission.

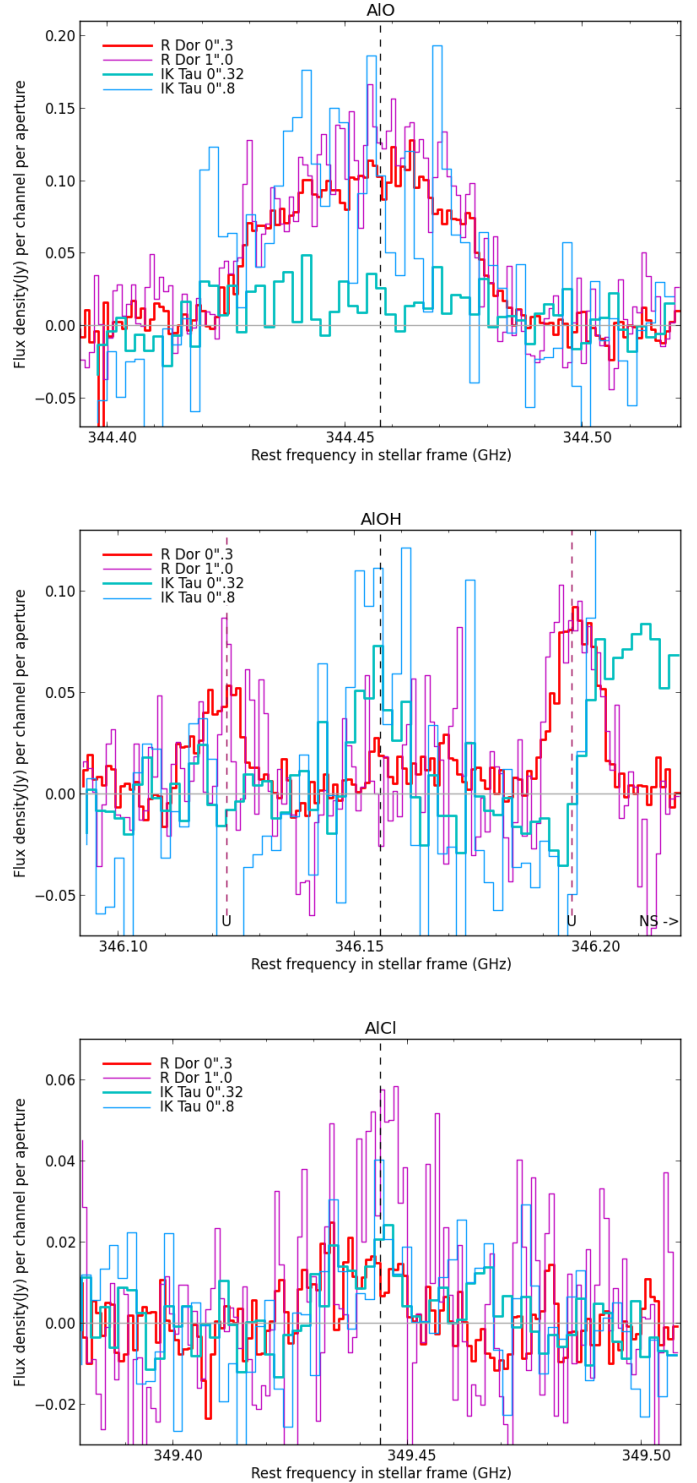


Fig. 1. ALMA AlO, AlOH, and AlCl data extracted for an aperture size of 320 mas (300 mas) and 800 mas (1000 mas) for IK Tau (R Dor). The frequency range is shifted to rest frequency using a local standard of rest velocity, ν_{LSR} , of 35 km s^{-1} for IK Tau and of 7 km s^{-1} for R Dor. Close to the AlOH emission, NS $^2\Pi_{1/2} J = 15/2-13/2$ transition is detected in IK Tau (at rest frequency 346.2201 GHz) and two unidentified lines (“U”) in R Dor at a rest frequencies of 346.123 GHz and 346.196 GHz.

- AlOH: the channel maps of IK Tau show a clear detection of AlOH with a peak strength around 0.07–0.12 Jy (depending on the extraction aperture; Fig. 2). The emission is centred

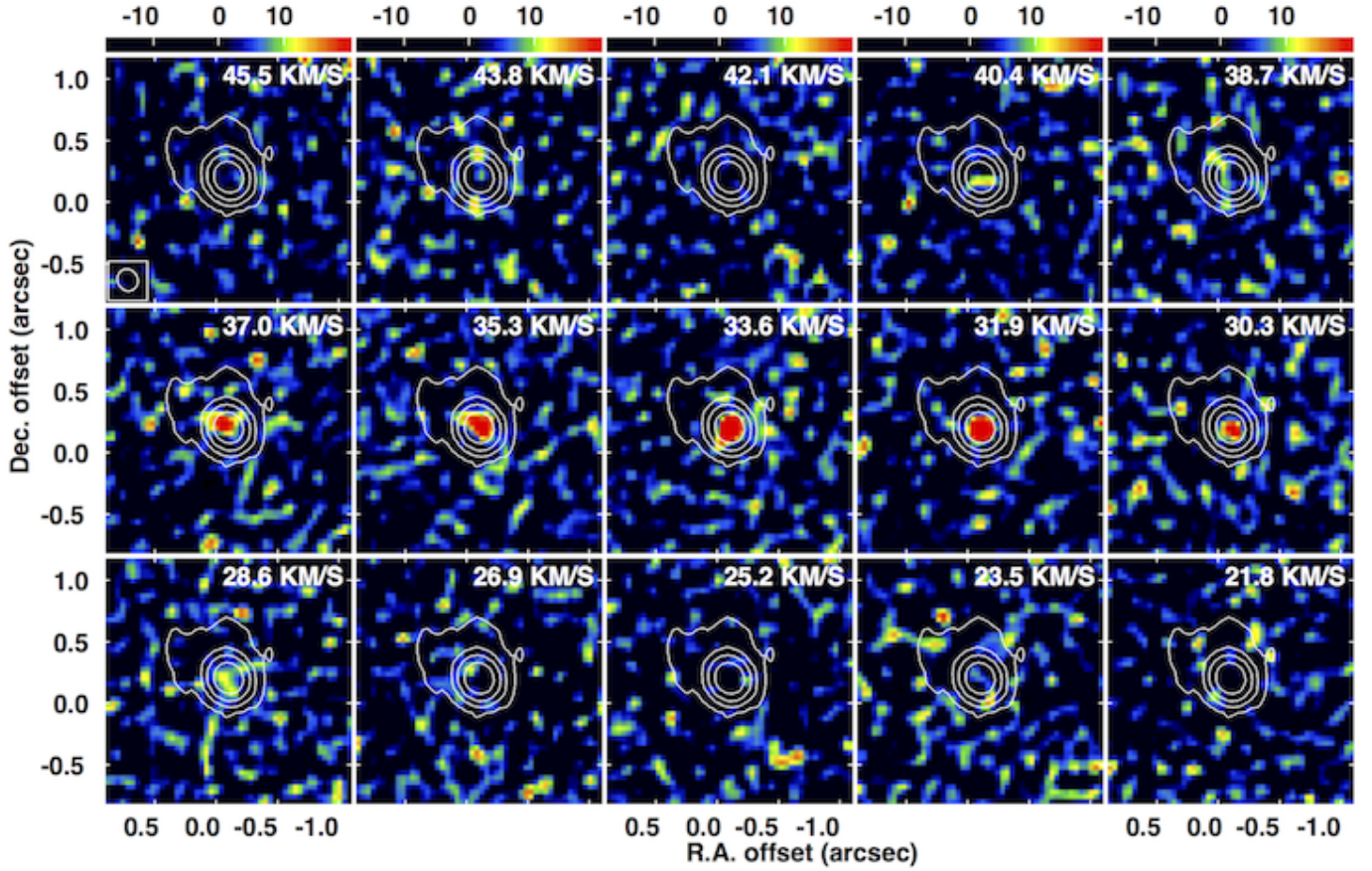


Fig. 2. Channel maps (colour scale in mJy) of the AIOH $J = 11-10$ transition in IK Tau. White contours are the continuum at $(-1, 1, 4, 16, 64, 256) \times 0.3$ mJy. The ordinate and co-ordinate axis give the offset of the right ascension and declination, respectively, in units of arcseconds. The ALMA beam size is shown in the bottom left corner of the *top panel*.

Table 2. Spectral properties of AIO, AIOH, and AlCl as observed with ALMA in IK Tau (*top part of the table*) and R Dor (*bottom part*).

(1) Molecule	(2) Velocity width (km s ⁻¹)	(3) Velocity ^a asymmetry (km s ⁻¹)	(4) Peak flux (Jy)	(5) Integrated flux (Jy km s ⁻¹)	(6) Velocity width (km s ⁻¹)	(7) Velocity ^a asymmetry (km s ⁻¹)	(8) Peak flux (Jy)	(9) Integrated flux (Jy km s ⁻¹)	(10) Angular width (mas)
IK Tau									
	Aperture 320 mas				Aperture 800 mas				
AIO $N = 9-8$	3.40	-0.32	0.036	0.247	32.30	14.98	0.186	3.211	800
AIOH $J = 11-10$	11.84	1.06	0.073	0.466	11.84	1.06	0.121	0.794	640
AlCl $J = 24-23$	6.70	-2.92	0.024	0.228	3.35	-2.92	0.040	0.255	160
R Dor									
	Aperture 300 mas				Aperture 1000 mas				
AIO $N = 9-8$	51.00 ^b	3.67	0.128	4.032	47.60	10.47	0.166	4.570	840
AIOH $J = 11-10$	3.38	-0.36	0.027	0.119	—	—	—	—	240
AlCl $J = 24-23$	3.35	2.65	0.021	0.113	9.22	-1.54	0.058	0.326	240

Notes. Values for the velocity width, velocity asymmetry, angular width, and peak and integrated flux are only given in the case of a clear detection.

^(a) The shift of the centroid w.r.t. the LSR velocity of the star is obtained by dividing the value of the “velocity asymmetry” by two. ^(b) Large velocity width is caused by the hyperfine structure; see Sect. 3.1.

around the peak of the continuum emission and has an angular width of ~ 240 mas.

- AlCl: around the v_{LSR} of ~ 35 km s⁻¹ almost no AlCl emission is seen, but the emission increases in strength at blue-shifted velocities (Fig. A.3). This is reflected in the asymmetric AlCl line profile in Fig. 1.

R Dor

- AIO: the bright AIO $N = 9-8$ is easily picked up by the ALMA instrument with a peak flux of $\sim 0.13-0.17$ Jy for the two different extraction apertures. The emission is smooth and concentrated around the peak of the continuum emission

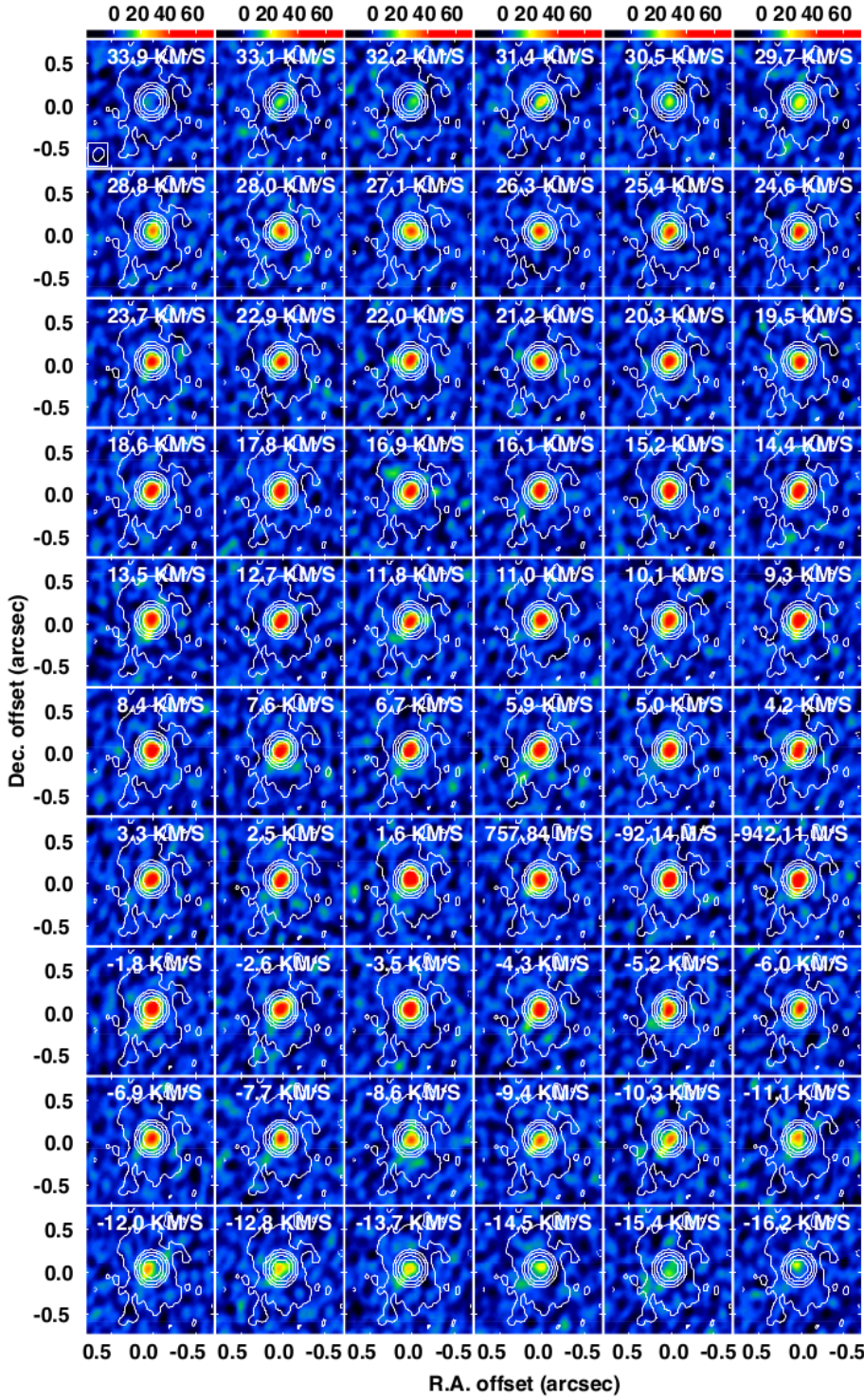


Fig. 3. Channel maps (colour scale in mJy) of the AIO $N = 9-8$ transition in R Dor. White contours are the continuum at $(-1, 1, 4, 16, 64, 256) \times 0.15$ mJy. The ordinate and co-ordinate axis give the offset of the right ascension and declination, respectively, in units of arcseconds. The ALMA beam size is shown in the *bottom left corner of the top panel*.

(Fig. 3). At an offset of ~ 280 mas to the south-east, enhanced AIO emission is seen in many channel maps (Fig. A.4). At the same offset location other molecules (including AIOH, SO, SiO, CO, and SO₂; see Decin et al., in prep.) show excess emission as well with the main peak having a blue-shifted frequency offset of 15–20 MHz.

- AIOH: the AIOH emission in R Dor is very weak and just above $3\sigma_{\text{rms}}$ for the 300 mas extraction aperture (Fig. 1). The main emission around v_{LSR} is slightly off-centre (as compared to the peak of the continuum emission; Fig. A.5). As noted already for AIO, a brighter clump of emission is visible around 280 mas to the south-east (Fig. A.6).

- AlCl: almost no AlCl emission is visible around v_{LSR} for the 300 mas aperture (Fig. A.7), but brighter emission becomes apparent at larger velocities with a peak flux around 0.02–0.06 Jy and an integrated flux of 0.1–0.3 Jy km s⁻¹ (depending on the extraction aperture).

The different velocity offsets for AlCl, on the red-shifted versus blue-shifted sides of the systemic velocity, for R Dor versus IK Tau cautions us to consider a potential blend or misidentification of this species.

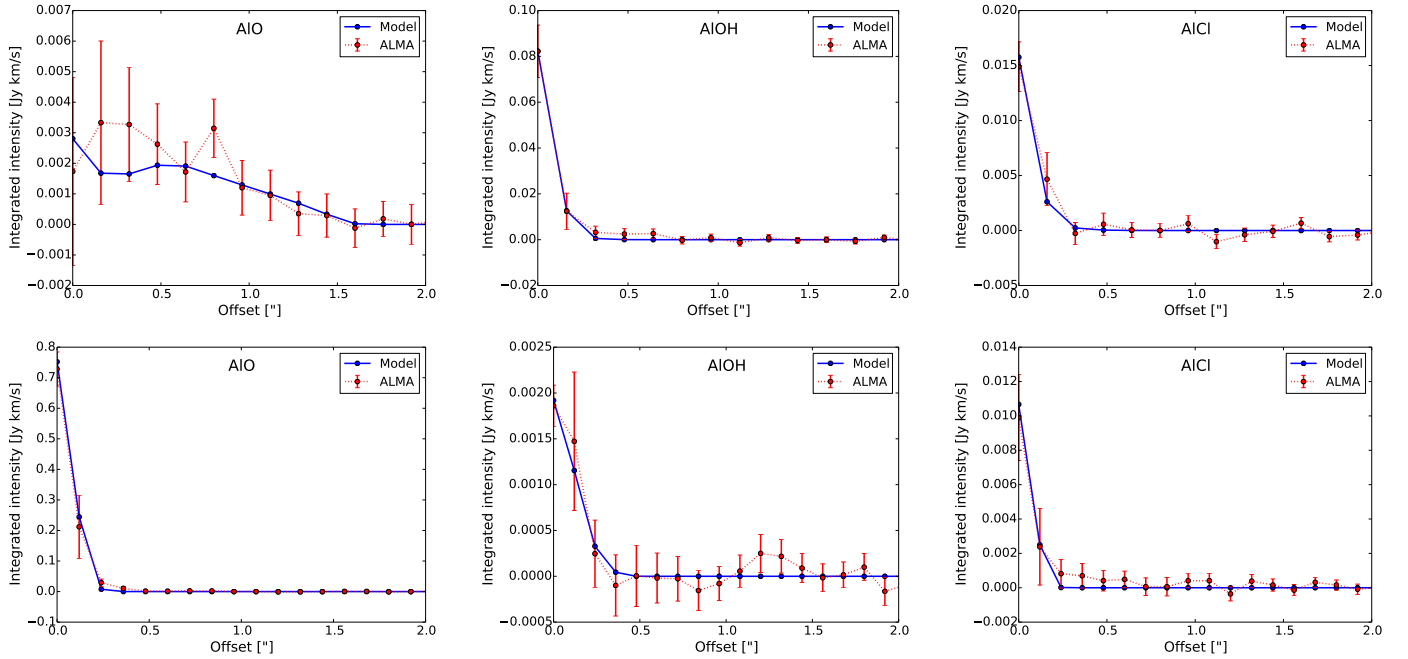


Fig. 4. Azimuthally averaged flux density in function of the angular distance from the central star for IK Tau (*top*) and R Dor (*bottom*), respectively. The ALMA data are shown in red with the vertical bars denoting the uncertainty. The radiative transfer model predictions are shown in blue (see Sect. 3.1).

2.3. Infrared continuum emission

Both R Dor and W Hya were observed with the Short Wavelength Spectrometer (SWS) on board of the Infrared Space Observatory (ISO) using the AOT01 observing mode. R Dor was observed at so-called speed 1, while for W Hya both speed 1 and speed 3 were used. The spectral resolution for the speed 3 data is around 700–1000 and is a factor of ~ 2 worse for the speed 1 data. Interferences in the instrument however cause fringes in the ISO band 3 data (12–29 μm), particularly in the case of the higher resolution observations. Unfortunately, no ISO-SWS data are available for IK Tau, where the only infrared spectrum available is from IRAS-LRS data (see Fig. 5).

The SPHERE/ZIMPOL data of R Dor and W Hya (Khoury et al. 2016; Ohnaka et al. 2016) confirm the earlier result by Norris et al. (2012) that a halo of large transparent grains is present at $\sim 1.5\text{--}3 R_\star$. Estimated grain sizes are around 0.3–0.5 μm and Fe-free silicates or Al_2O_3 are put forward as the possible dust species that produce the scattered light.

Zhao-Geisler et al. (2011) monitored W Hya with MIDI/VLTI (8–13 μm , spectral resolution of 30) during nearly three pulsation cycles. From the spectrally dispersed visibility data, the authors concluded that the Fe-rich silicate emission seen around 10 μm in the ISO-SWS data arises from a region fairly far away from the star, at radii larger than 30 times the photospheric radius. Using image reconstruction techniques, Zhao-Geisler et al. (2015) were able to spatially resolve the dust formation region close to the star and deduced that the data signal the presence of a dust layer of presumably amorphous aluminium oxide (Al_2O_3) at $\sim 2 R_\star$. The observations suggest that the formation of amorphous Al_2O_3 occurs mainly around or after visual minimum (Zhao-Geisler et al. 2012).

VLTI/MIDI data are also available for R Dor. However the fringe-track data do not contain any interferometric signals. The beams were probably not overlapped correctly, as the data were taken for test purposes. No MIDI data are available for IK Tau.

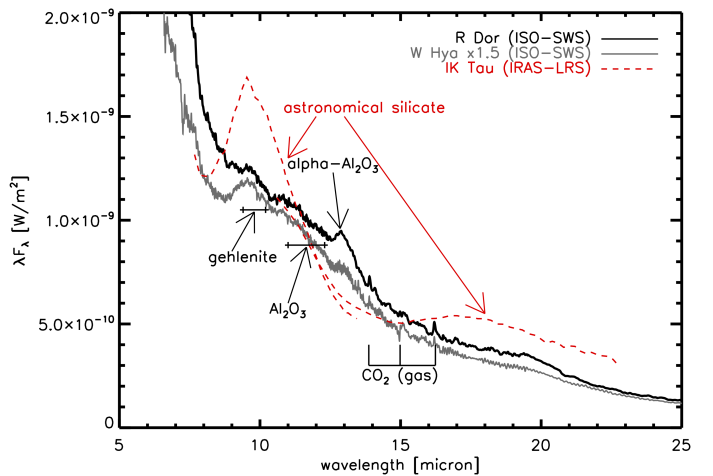


Fig. 5. Infrared spectrum of R Dor (black, full line), W Hya (gray, full line, scaled with a factor 1.5), and IK Tau (red, dashed line). Spectral features arising from dust species such as crystalline and amorphous Al_2O_3 , gehlenite ($\text{Ca}_2\text{Al}_2\text{SiO}_7$), and so-called astronomical silicates (amorphous olivines with different relative magnesium and iron fractions; de Vries et al. 2010) are indicated.

The SED of W Hya together with infrared images and optical scattered light fractions were analysed by Khouri et al. (2015). The only way to explain all observed signatures, and in particular the amorphous Al_2O_3 emission in the SED, was by implementing a so-called gravitationally bound dust shell (GBDS) close to the star at around 1.5 to 3 stellar radii in their radiative transfer model. Observations and models of AGB winds have long shown the existence of dense gas layers that are rich in molecules and dust that are gravitationally bound to the star (Bowen 1988; Willacy & Cherchneff 1998; Tsuji 2000; Ireland et al. 2004; Ohnaka et al. 2005; Cherchneff 2006; Wittkowski et al. 2007).

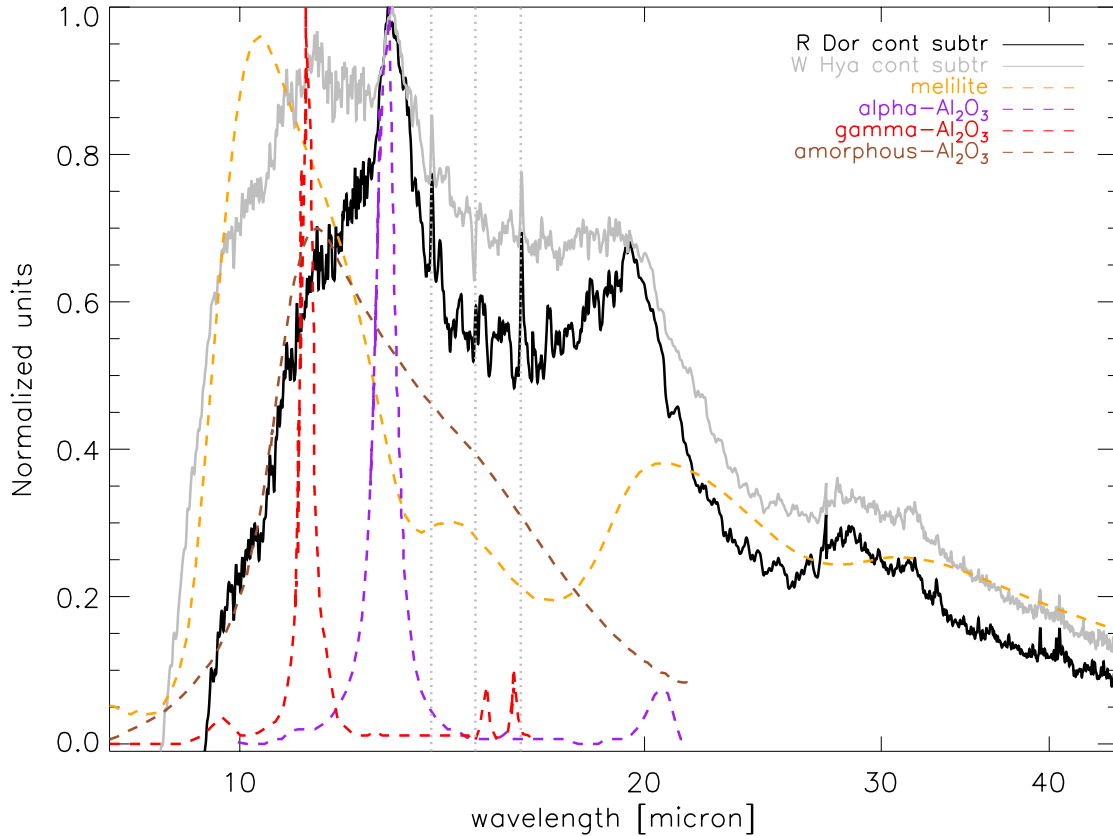


Fig. 6. Continuum subtracted and normalized SED of R Dor and W Hya (in full black and gray lines, respectively). Characteristic (scaled) emission patterns of different dust species are shown in the coloured dashed lines. The gray vertical dotted lines indicate CO₂ molecular emission (Justtanont et al. 1998; Ryde et al. 1999).

In the GBDS, the ratio of the radiation pressure on the grains to the gravitational attraction (called the Γ -factor) is still smaller than 1 so that the dust particles can reside close to the star without being pushed outwards. This allows the material to potentially grow in size, and Γ can become larger than 1 only once their size is larger than 0.1–0.3 μm . This idea of a GBDS was recently shown to fit well with the theoretical dynamical wind models of Höfner et al. (2016), who allowed for the formation of composite grains with an Al₂O₃ core and a silicate mantle at slightly larger distances than the Al₂O₃ GBDS shell, thereby giving the wind a kick start. The location of the GBDS coincides with the region in which large transparent grains are detected in polarized light images (Norris et al. 2012; Ohnaka et al. 2016), albeit this does not imply by necessity that the species responsible for the Al₂O₃ emission in the SED are the same species that cause the stellar light to be scattered. Thus, it is still possible that the scattering agents are Fe-free silicates, since these can remain fairly cold and hence do not produce signatures in the thermal infrared.

The poor quality of the infrared continuum data of IK Tau led us to decide not to analyse the dust content in this object, but to focus on the dust composition of R Dor and W Hya. We combined the available IR data for the latter two objects and subtracted the stellar continuum (represented as a black body with temperatures of 3000 K for R Dor and of 2500 K for W Hya) to visualize better their dust spectral features; see Fig. 6. The emission bands of some dust aluminium bearing dust species are plotted along with some CO₂ and H₂O molecular emission features.

3. Results

In this section, we first determine the mean fractional abundances of AlO, AlOH, and AlCl in the inner winds of R Dor and IK Tau (Sect. 3.1). These results are then compared quantitatively to the theoretical dynamical-chemical model predicting the inner wind abundance structure (Sect. 3.2). The amount of aluminium locked up in gaseous species turns out to be low, leaving ample room for aluminium to be incorporated into dust grains. However, as we discuss in Sect. 3.3, the estimated grain temperatures for aluminium oxide (Al₂O₃) at $\sim 2R_*$ is high, which suggests that the grains are annealed and hence have a crystalline lattice structure. This outcome implies a tension in the interpretation of the 11 μm spectral feature, present in the SED and interferometric data. We hypothesize that large gas-phase aluminium oxide clusters are the cause of the 11 μm feature and explain how these large clusters can be formed (Sect. 4).

3.1. Abundances of aluminium-bearing molecules

To establish the abundance and distribution of AlO, AlOH, and AlCl, we modelled the ALMA data using a non-LTE radiative transfer model of the circumstellar envelope (CSE). We use a code based on the Accelerated Lambda Iteration (ALI) method (Maercker et al. 2008), which allows us to retrieve the global mean molecular density by assuming a 1D geometry. Collisional excitation rates have not been published for any of these three molecules. For that reason, we used the values for the HCN-H₂ system (Green & Thaddeus 1974) as substitute for AlOH (cf. Tenenbaum & Ziurys 2010) and the values of SiO-H₂

(Dayou & Balança 2006) for AIO and AlCl (cf. Kamiński et al. 2016), scaling for the difference in molecular weight. The collisional rates are used in the form they appear in the LAMDA database² (Schöier et al. 2005). To check for the dependency on collisional excitation rates, we also ran models using the CS-H₂ cross sections (Turner et al. 1992) and found no significant difference in results. The first 30 rotational levels in the ground vibrational state were considered for AIOH and the first 40 such levels were considered for AIO and AlCl. Our treatment of the radiative transfer does not deal with the (hyper)fine structure components, and is therefore limited to the prediction of rotational lines $N \rightarrow N'$ or $J \rightarrow J'$ with ΔN or ΔJ equal to 1. Therefore, all levels treated are described by one quantum number, N or J . The final line profile is computed by splitting the predicted total intensity over the different (hyper)fine components according to their relative quantum-mechanical line strength, S , as given in CDMS³ (Müller et al. 2001, 2005). This approach is only valid under LTE conditions.

For an angular diameter of 20 mas and a distance of 260 pc, the stellar radius of IK Tau is $\sim 3.8 \times 10^{13}$ cm, while the angular diameter of 57 mas and the distance of 60 pc translates into a stellar radius of $\sim 2.5 \times 10^{13}$ cm for R Dor. The gas kinetic temperature is assumed to follow a power law (Justtanont et al. 1994; Groenewegen 1994; Decin et al. 2006)

$$T_{\text{gas}} = T_{\star} \left(\frac{R_{\star}}{r} \right)^{\epsilon}, \quad (1)$$

and the gas velocity profile is parametrized using a β -type law

$$v_{\text{gas}} = v_0 + (v_{\infty} - v_0) \left(1 - \frac{r_0}{r} \right)^{\beta}, \quad (2)$$

with r the distance to the star, v_{∞} the terminal velocity, and v_0 the velocity at radius r_0 at which the wind is launched and which is set to the local sound speed. The gas velocity within the dust condensation radius, $r < r_0$, is assumed to be 2 km s^{-1} . The turbulent velocity for both stars is assumed to be 1 km s^{-1} . The gas density, $\rho(r)$, is calculated from the equation of mass conservation

$$\dot{M} = 4\pi r^2 v(r) \rho(r). \quad (3)$$

The main input parameters for R Dor and IK Tau are listed in Table 3, following Maercker et al. (2016). These properties are simplified estimates, for example the velocity structure inside the dust formation radius is complex, but are adequate for 1D modelling in the regions where these Al compounds have been observed. For all molecular species, the fractional abundance structure with respect to H₂ was varied until a fit with the ALMA azimuthally averaged flux density was achieved (see Fig. 4). In some cases, the fractional abundance declines according to a Gaussian profile centred on the star, and hence is described by following equation

$$f(r) = f_0 \exp \left(- \left(\frac{r}{r_e} \right)^2 \right), \quad (4)$$

with f_0 the initial abundance and r_e the e -folding radius.

The retrieved fractional abundance structures are shown in Fig. 7. Clearly both stars show very different distributions of Al-bearing molecules. For IK Tau, AIOH could be modelled assuming a Gaussian distribution with a peak central abundance

Table 3. Input parameters for the non-LTE radiative transfer modelling.

	R Dor	IK Tau
T_{\star} [K]	2400	2100
R_{\star} [cm]	2.5×10^{13}	3.8×10^{13}
L [L_{\odot}]	6500	7700
\dot{M} [M_{\odot} /yr]	1.6×10^{-7}	5×10^{-6}
r_0 [cm]	5.3×10^{13}	2.38×10^{14}
ϵ	0.65	0.6
$T(r_0)$ [K]	1530	650
v_{∞} [km s ⁻¹]	5.7	17.5
β	1.5	1.2
$v(r_0)$ [km s ⁻¹]	3.9	2.5

of 4.4×10^{-9} and an e -folding radius, r_e , of $22 R_{\star}$. As could be deduced already from Figs. 1 and A.2, the AIO emission might be clumpy and not show a centrally peaked abundance structure. The data could be modelled assuming a constant abundance of 4.4×10^{-8} from 50 – $160 R_{\star}$. If the AlCl identification is correct (see Sect. 2.2), then the data are compatible with a Gaussian distribution with peak central abundance of 9×10^{-10} and r_e of $40 R_{\star}$. The AIOH emission in R Dor can be approximated with a constant slab with abundance of 1.4×10^{-9} from the stellar surface out to $12 R_{\star}$. The AIO and AlCl distribution in R Dor follow a Gaussian distribution with peak central abundance of 8.4×10^{-8} and 2.5×10^{-8} and e -folding radius of $3.8 R_{\star}$ and $2.4 R_{\star}$, respectively. The good correspondence between the observed and predicted emission of AIO in R Dor, allowing for the splitting over the different hyperfine components, is shown in Fig. 8.

While we can trace the fractional abundance above 1×10^{-9} within $\sim 150 R_{\star}$ for IK Tau, the ALMA Al-data of R Dor only cover the first $25 R_{\star}$. The factor ~ 6 difference in spatial extent is in line with the factor ~ 50 in mass-loss rate and hence sensitivity to some particle density assuming a density law, which is proportional to r^{-2} .

For the low-mass loss rate star *o* Ceti, Kamiński et al. (2016) derived AIO abundance from ALMA data in the range 10^{-9} – 10^{-7} and concluded a small amount of aluminium was locked up in AIO. Our AIO abundance for R Dor shown in Fig. 7 is compatible with this value range and we reach a similar conclusion regarding the small Al content in Al-bearing molecules. Indeed, assuming a solar aluminium content of $[\text{Al}/\text{H}] = 3 \times 10^{-6}$, our results imply that for both stars only a small fraction of aluminium ($\lesssim 2\%$) is locked up in AIO, AIOH, and AlCl.

3.2. Gas-phase aluminium budget

The deduced AIO, AIOH, and AlCl abundances of IK Tau can be compared directly to the chemical predictions by Gobrecht et al. (2016). They modelled the synthesis of molecules and dust in the inner wind of IK Tau ($r < 5 R_{\star}$) by considering the effect of periodic shocks induced by the stellar pulsation. The non-equilibrium chemistry in the shocked gas layers was followed between 1 and $10 R_{\star}$ by solving a chemical kinetic network including 100 species and 424 reactions. The condensation of dust is described by a Brownian formalism using the simplifying assumption that the gaseous dimers of forsterite (Mg_2SiO_4), enstatite (MgSiO_3), and alumina (Al_2O_3) are the initial dust seeds and considering coalescence and coagulation of these dimers to form the dust grains.

The aluminium species in the network of Gobrecht et al. (2016) include Al, AlH, AIO, AIOH, AIO₂, AlCl, Al₂, and Al₂O.

² <http://home.strw.leidenuniv.nl/~moldata/>

³ <https://www.astro.uni-koeln.de/cdms/catalog>

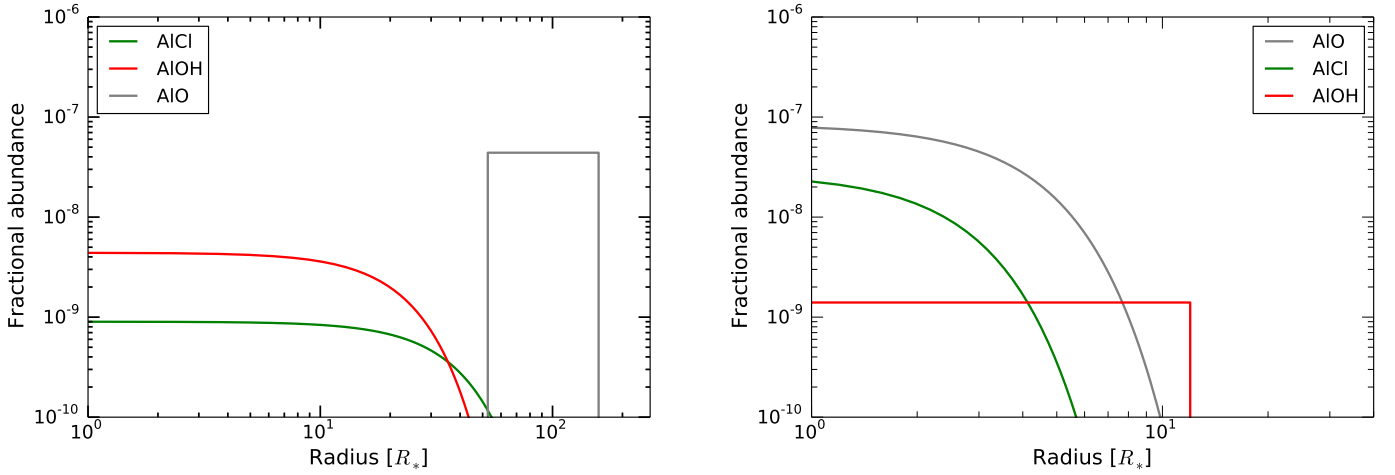


Fig. 7. Retrieved fractional abundance structure $[\text{AIO}/\text{H}_2]$, $[\text{AIOH}/\text{H}_2]$, and $[\text{AICI}/\text{H}_2]$ for IK Tau (*left*) and R Dor (*right*).

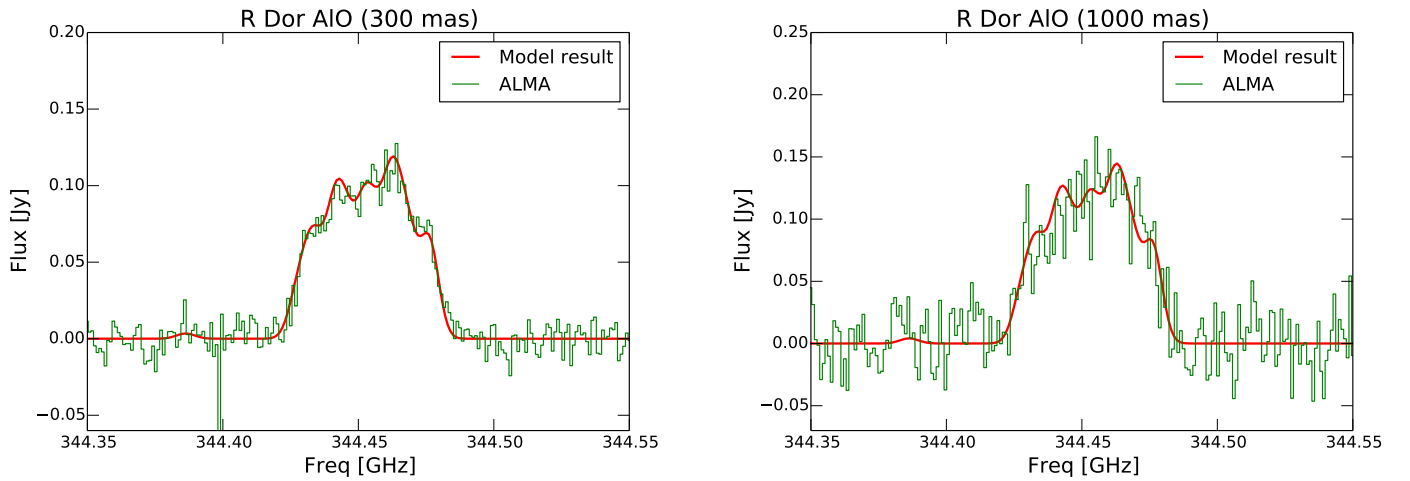


Fig. 8. Comparison between the extracted ALMA AIO spectra of R Dor (green) and the predicted line profile (red) for an AIO abundance structure as given in Fig. 7 for the 300 mas (*left*) and 1000 mas (*right*) extraction aperture.

The aluminium dust precursors are described by Al_2O_2 , Al_2O_3 , and Al_4O_6 . The predicted abundances of the aluminium species as a function of radius and phase ϕ^4 are shown in Figs. 9 and 10, respectively. Al_2 and Al_2O have very low abundances and therefore do not appear in the plots. The plot in Fig. 9 can be interpreted as a snapshot of the inner wind just after the passage of a shock (at phase $\phi = 0.0$) assuming shock fronts every $0.5 R_*$. A full pulsation cycle corresponds to phase 1.0. Phase 0.0 also corresponds also to a snapshot after a full pulsation cycle. However, at this point, the gas densities have been rescaled from the previous radial position and just have been shocked again at the new radial position (see description in Gobrecht et al. 2016).

For a period of 470 days and with T_0 at JD $\sim 2\,453\,025$ (Matsumoto et al. 2008), the ALMA data of IK Tau are obtained around $\phi = 0.98$. The method applied by Gobrecht et al. (2016) implies that a relative comparison between the retrieved and predicted molecular abundance stratifications is best carried out using Fig. 9, while the absolute scaling can be seen in Fig. 10. Taking the predicted abundances at $5 R_*$ (Fig. 10) at phase $\phi = 0.98$ as representative values for the comparison

with the profiles retrieved from the ALMA data (left panel in Fig. 7), we can see that both the observations and chemical models predict a higher abundance for AIOH than for AIO. However, the retrieved AIOH abundance at $5 R_*$ is 2 orders of magnitudes lower than predicted. The ALMA data indeed show that the fractional AIO abundance ($r < 50 R_*$) is very low ($< 10^{-10}$) in accord with the predictions, but remarkably we see that an active Al-chemistry is taking place further out in the wind yielding an increase in AIO emission. We derive a mean AIO fractional abundance of $\sim 5 \times 10^{-8}$ between $50\text{--}150 R_*$ for IK Tau with the emission mainly arising from clumps at that radial distance. The increase in AIO abundance coincides with a strong decrease in AIOH abundance, suggesting that AIOH is converted to AIO. Another possibility is that desorption or sputtering release AIO from grains back into its gaseous form. If no line blend is blurring the AICI identification, then the ALMA data show that the AICI abundance in IK Tau is a factor ~ 40 higher than predicted.

With an inner dust condensation radius around 2.4×10^{14} cm (or ~ 120 mas in diameter), the ALMA observations of IK Tau are not sensitive to the potential presence of transparent grains closer to the star and the ALMA beam is too large to confirm the predicted strong radial variation in molecular abundance within $5 R_*$ (as visible in Fig. 9). The stellar angular diameter of R Dor

⁴ The pulsation phase is defined as the decimal part of $(t - T_0)/P$, where T_0 is the epoch of the start of the pulsation cycle and P is the period.

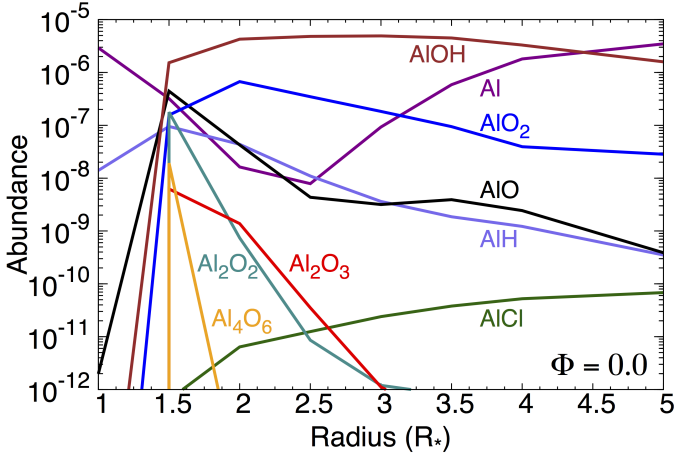


Fig. 9. Predicted abundances of aluminium-bearing molecules with respect to the total gas number density as a function of radius. The chemical models are presented by [Gobrecht et al. \(2016\)](#); only the aluminium-species are shown. Model simulations for a star resembling IK Tau at phase $\phi = 0.0$.

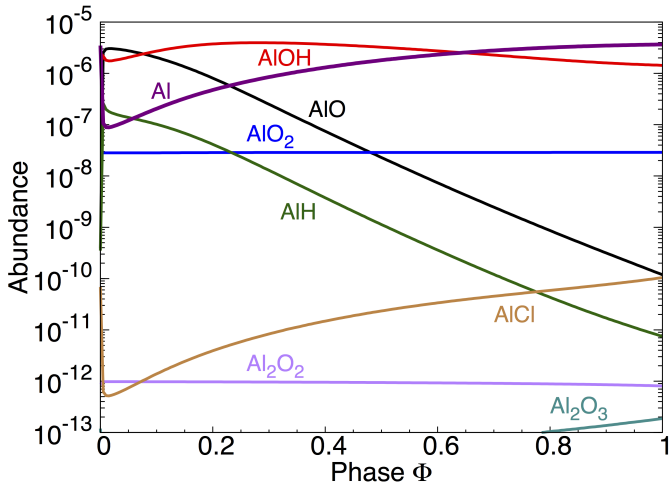


Fig. 10. Predicted abundances of aluminium-bearing molecules with respect to the total gas number density as a function of phase ϕ at $5 R_*$. Chemical models are based on the pulsation-induced shock dynamics presented by [Gobrecht et al. \(2016\)](#) for a star resembling IK Tau; the analogous results at $1 R_*$ are shown in their Fig. 6.

is much larger, $\theta_D \sim 57$ mas. Although a direct comparison with Figs. 9, 10 should be carried out with care since the shock velocities are expected to be smaller for R Dor, our ALMA data do not corroborate any strong radial variation in AIO and AIOH abundance within the first 10 stellar radii. In addition, the ALMA results on R Dor prove that a second phase of dust formation, which is proposed to occur slightly beyond the GBDS, is (almost) not consuming AIO. This supports the scenario suggested by [Höfner et al. \(2016\)](#) that the growth of

silicates (potentially on top of already existing alumina grains) is a trigger for the wind to get launched.

It would be of interest to monitor the changes in abundance structure in function of the pulsation phase and confront such changes with the predictions shown in Fig. 10. This kind of data could then be used to benchmark and improve current chemical models that implement gas dynamics. [Kamiński et al. \(2016\)](#) have reported some variability in the AIO emission of the low mass-loss rate Mira star *o* Ceti, but currently no link with the pulsation cycle is apparent.

Although the current comparison points towards improvements to be made in the chemical model predictions, a conclusion from the ALMA data in combination with these models is that the fraction of aluminium that is locked up in simple aluminium-bearing molecules seems small. That is, the combined fractional abundance of Al-molecules that are thought to form easily in the inner wind region is only $\sim 5 \times 10^{-8}$. Furthermore, since the formation of the more complex Al_2O_3 and its dimer $(\text{Al}_2\text{O}_3)_2$ is described by termolecular recombination and hence requires high densities, these higher order molecules also only consume a tiny fraction of the available aluminium (see Fig. 10).

The most difficult gaseous aluminium form to constrain is that of atomic aluminium. The atomic aluminium abundance is predicted to be high when the parcel of material is close to the star ($r < 2.5 R_*$) for phases $\phi > 0.2$. Exactly at the shock front, phase $\phi = 0$, the high temperature ensures a high Al abundance, but as a consequence of the declining temperature for later phases the atomic Al abundance decreases rapidly. Beyond $3 R_*$, molecular Al-species are transformed in atomic Al because of the decreasing gas densities with distance from the star. We performed an in-depth study of the high spectral resolution optical Mercator/Hermes ([Raskin et al. 2011](#)) data of IK Tau and W Hya (no observations are available for R Dor) to check for the presence of atomic Al in the outer atmosphere/inner wind of both stars. Molecular line veiling complicates the analysis of the data. We could not identify either absorption or emission atomic Al lines. This does not imply that no atomic Al is present, but at least the quantity is too low for it to be detected amongst the molecular lines. An analogous conclusion was drawn by [Kamiński et al. \(2016\)](#) for the low mass-loss rate AGB star *o* Ceti; thus, they concluded that Al is locked in molecules and dust.

This ensemble of considerations supports the conclusion that only a minor fraction of Al resides in Al-bearing molecules and that the atomic Al abundance is difficult to constrain. An important outcome from these ALMA data is that the aluminium condensation process, considered to be the first condensation cycle to start in AGB stars ([Tielens 1990](#)), is for sure not fully efficient since the gaseous precursors of the aluminium grains are detected in and beyond the dust formation region.

3.3. Aluminium depletion into grains

Presolar grains prove that aluminium oxide grains are formed in oxygen-rich AGB stars. The circumstellar Al_2O_3 -grains are relatively abundant in meteorites ([Nittler et al. 1997](#)), albeit perhaps not as abundant as one would expect if all aluminium would condense in these grains ([Alexander 1997](#)). This result from meteorite studies now gets some support by our analysis of the ALMA data, confirming that a fraction of the aluminium is still present in its gaseous form after its passage through the dust formation region. At least for a few silicate grains, the meteoritic studies show that the aluminium oxides are at the core

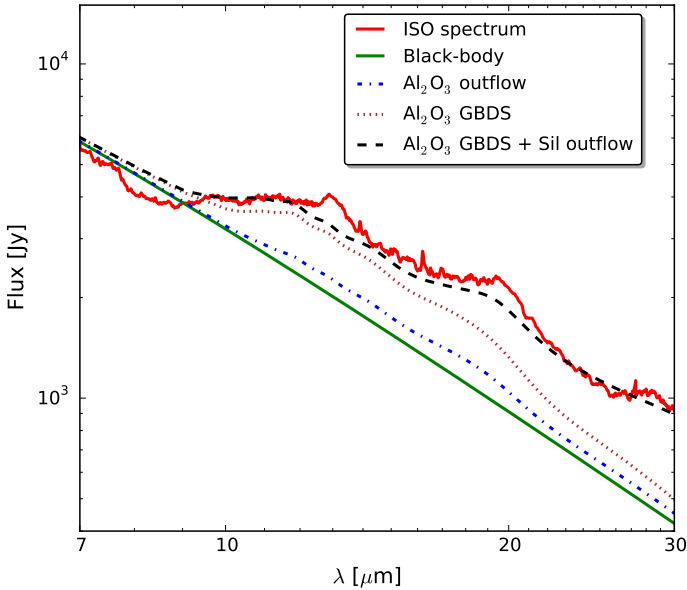


Fig. 11. Infrared SED of R Dor. ISO data are shown in red. The black body represents the stellar emission at a temperature of 3000 K, the blue dash-dotted line indicates a model assuming a 100% condensation of aluminium into amorphous Al_2O_3 that resides in the stellar outflow, the brown dotted line shows the model fit for Al_2O_3 residing in the GBDS, while the black dashed line represents a model fit for amorphous Al_2O_3 that is present in the GBDS and silicates in the stellar outflow.

of the silicate grain suggesting that Al_2O_3 is the first solid to form in the CSE (Nittler et al. 2008; Vollmer et al. 2006). Both crystalline and amorphous Al_2O_3 enclosures have been detected (Stroud et al. 2004). The variation in the crystal structure and Ti content of these presolar grains demonstrate that Al_2O_3 can condense in the absence of TiO_2 seed clusters but that Ti may be important in determining the lattice structure.

The SED analysis of R Dor and W Hya turned out to be challenging in terms of the aluminium content (Khoury 2014; Khouri et al. 2015). Based on constraints from gas-phase models and solar abundances, Khouri et al. were unable to model the SEDs of both stars in the spectral region around $11\,\mu\text{m}$ when considering aluminium condensates only to be present in the stellar outflow (see dotted line in Fig. 11). One can fit the broad $11\,\mu\text{m}$ feature in the SED by invoking the presence of a gravitational bound dust shell (GBDS; see Sect. 2.3 for a more detailed explanation on this terminology) of amorphous Al_2O_3 grains at $\sim 2R_\star$ (see dashed line in Fig. 11). The GBDS can contain both small and large Al_2O_3 grains. The model for R Dor (W Hya) requires an Al_2O_3 dust mass in the GBDS of $\sim 5.7 \times 10^{-10} M_\odot$ ($\sim 1.5 \times 10^{-9} M_\odot$), which would correspond to a gas mass of $7.5 \times 10^{-6} M_\odot$ ($2 \times 10^{-5} M_\odot$) if one would assume full aluminium condensation. Whether this is compliant (or not) with the total aluminium content is difficult to assess since the region where the GBDS resides is known to be highly variable in density due to the stellar pulsations with variations up to one order of magnitude (Höfner et al. 2016).

The GBDS results (Khoury 2014; Khouri et al. 2015), scattered light data (Norris et al. 2012; Khouri et al. 2016; Ohnaka et al. 2016), interferometric results (Zhao-Geisler et al. 2015), and theoretical models (Höfner et al. 2016; Gobrecht et al. 2016) build up evidence for a shell of aluminium oxide grains at $1.5\text{--}3R_\star$. The kinetic gas temperature in that region is around 1500–2000 K. Optical constants are measured

for amorphous Al_2O_3 grains, but only for wavelengths beyond $0.2\,\mu\text{m}$ (Koike et al. 1995) or beyond $7.8\,\mu\text{m}$ (Begemann et al. 1997). Depending on the replica used for the short-wavelength range and under the assumption of thermodynamic equilibrium, the estimated temperature of the amorphous alumina grains at that spatial region would be around 1200–1600 K (Khoury et al. 2015). However, amorphous alumina can only exist below 1000 K (Levin et al. 1998; Levin & Brandon 2005). At temperatures around 1500 K, crystalline α -alumina can form, while below 1300 K crystalline γ -alumina can exist. Because the grain temperature in the $1.5\text{--}3R_\star$ region is higher than the annealing temperature, this would imply a structural transformation of Al_2O_3 to its crystalline form. The thermal infrared spectrum of R Dor and W Hya is not compatible with the presence of Al_2O_3 grains with a crystalline γ -lattice structure (see Fig. 6). As shown in Fig. 6 crystalline α - Al_2O_3 can indeed explain the $13\,\mu\text{m}$ feature in the SED. A crystallinity degree of 5% can explain the $13\,\mu\text{m}$ feature in W Hya, while for R Dor the estimated crystallinity degree is around 12%. These crystalline grains might be the scattering agents of the polarized light emission.

However, this makes us question how to explain the broad $11\,\mu\text{m}$ feature in the SED of R Dor and W Hya and, in particular, the MIDI data of W Hya, which clearly show that emission arises very close to the star. A potential answer to that question is that the $11\,\mu\text{m}$ feature does not signal the presence of solid-state amorphous aluminium oxide, but – to the contrary – points towards the existence of large neutral gas-phase aluminium oxide clusters (see Sect. 4). Individual $(\text{Al}_2\text{O}_3)_n$ clusters may exist above 1500 K. Our analysis does not exclude the existence of amorphous aluminium oxides in the wind, but only at larger distances from the star where temperatures are cooler.

4. Discussion: indication for the presence of aluminium oxide clusters

If amorphous alumina dust grains are not the carrier of the MIDI and ISO spectral signature at $11\,\mu\text{m}$, then the interesting possibility exists that large aluminium oxide clusters are present, and detected, just above the stellar surface. Indeed, an outcome of the experimental IR-REMPI studies by van Heijnsbergen et al. (2003) is that large $\text{AlO}(\text{Al}_2\text{O}_3)_n$ clusters ($n > 34$) exhibit a spectral signature that is very similar to that of amorphous Al_2O_3 , while smaller $\text{AlO}(\text{Al}_2\text{O}_3)_n$ clusters (around $n = 15$) show the best match with that of crystalline γ - Al_2O_3 (see Fig. 12). Because of the non-linear regime in which the IR-REMPI results are obtained, one cannot however estimate the amount of clusters present in the beam. This renders the impossibility of quantifying the amount of large aluminium oxide clusters that would be needed to explain the $11\,\mu\text{m}$ feature.

If the $11\,\mu\text{m}$ feature indicates the presence of large ($n > 34$) aluminium oxide clusters, then one should wonder if smaller clusters are present as well and if they can be detected. The experiments by van Heijnsbergen et al. (2003) and Demyk et al. (2004) show that clusters with $n = 15\text{--}40$ show two vibrational bands: one around $11.8\,\mu\text{m}$ and another one around $15\,\mu\text{m}$. Comparing this to the emission features of solid-state aluminium oxide lattices identifies these gas-phase clusters as having a structure similar to crystalline γ - Al_2O_3 instead of α - Al_2O_3 (corundum), which is the thermodynamically most stable form of the bulk (see Fig. 12). The infrared observations of W Hya and R Dor do not display an obvious emission component around $15\,\mu\text{m}$. This does not imply by necessity that these $n = 15\text{--}40$ sized clusters are not present, but at least they are not abundant enough to leave any spectral imprint.

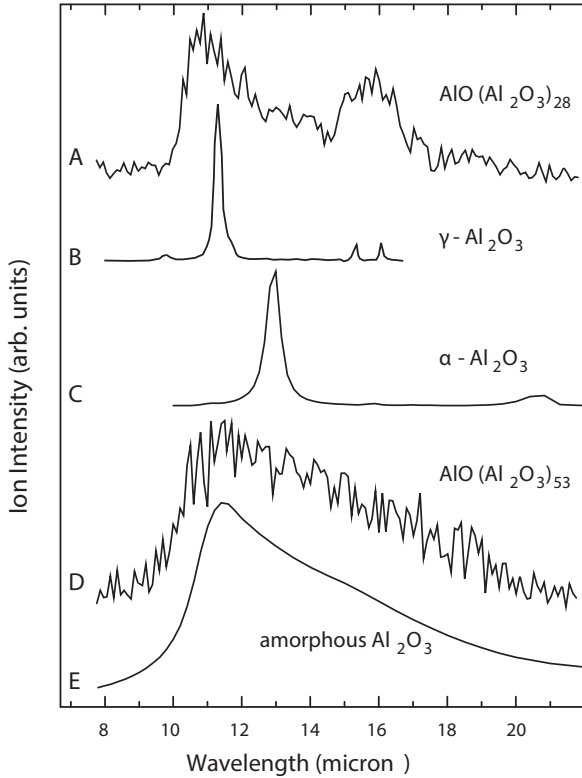


Fig. 12. Comparison of gas-phase vibrational spectra for $\text{AlO} \cdot (\text{Al}_2\text{O}_3)_n$ with $n = 28$ (A) and for $n = 53$ (D) to the infrared absorption spectra of bulk aluminium oxide lattices (B = crystalline $\gamma\text{-Al}_2\text{O}_3$; C = crystalline $\alpha\text{-Al}_2\text{O}_3$; E = amorphous Al_2O_3). This result is obtained by van Heijnsbergen et al. (2003).

A variety of processes including dust production in type II supernovae and AGB stars, dust destruction by supernova shocks, grain growth, and grain-grain collisions determine the grain size distribution in the interstellar medium of galaxies (Asano et al. 2013). Shattering is crucial in producing small grains ($\leq 0.01 \mu\text{m}$). The size distribution of interstellar dust grains in our Milky Way is dominated by small particles, i.e. $n_d(a) da \propto a^{-7/2} da$ (Mathis et al. 1977). Also in our own Earth's atmosphere, fresh aerosols that are created in situ from the gas phase by nucleation display a number distribution heavily skewed towards the very small particles (Seinfeld et al. 1998). It is therefore interesting to look for the presence of very small clusters in AGB winds. Current capacities in quantum chemical computations give us the possibility to test this proposition on the prevailing presence of very small clusters; see next section.

4.1. Emission of very small $(\text{Al}_2\text{O}_3)_n$ ($n \leq 4$) aluminium oxide clusters

A way to test if very small Al_2O_3 polymers are present is via the detection of their vibrational frequencies. These bands arise owing to the presence of internal bending and stretching modes in the cluster. As the energy and geometry, the vibrational spectrum differs from isomer to isomer and thus can be seen as a spectral fingerprint.

Several theoretical and experimental studies on the structure and spectroscopy of small alumina clusters have been performed. Demyk et al. (2004) and Sierka et al. (2007) measured the IR-REMPI spectra of laboratory-produced $\text{AlO} \cdot (\text{Al}_2\text{O}_3)_n$ clusters down to $n = 7$, whereas Li & Cheng (2012) provided

structures for a selection of alumina clusters obtained from density functional theory (DFT) calculations. Because there is no way to currently predict the most likely cluster composition in the outflow and because these clusters would probably cover a wide range in both stoichiometry and atomic sizes, we took the simple approach of calculating the spectral properties of a range of stoichiometric alumina clusters to estimate where observable lines might exist. For $n = 1-4$ we determined the most stable structures of the neutral aluminium oxide clusters using DFT; see Fig. 13. The computations were accomplished by the GAUSSIAN 09 package (Frisch et al. 2009) using the hybrid B3LYP (Becke, three parameter, Lee-Yang-Parr; Lee et al. 1988; Miehlich et al. 1989; Becke 1993) exchange-correlation functional with a 6-311+g* basis set. The binding energy and stability of electronic systems including clusters are pressure and temperature-dependent, respectively. Variations in temperature and density imply that the energy levels of the clusters may shift and that the order of the most stable clusters may change. We calculated how the stability⁵ of the considered small alumina clusters is affected in various ways for conditions occurring in the CSE (with pressure $p = 0.001-50$ Pa and temperature $T = 500-3000$ K). For Al_2O_3 , the first excited structural isomer is a linear chain and has a potential energy 0.1 eV above the kite-shaped triplet ground state. The linear form is slightly more stable above 1000 K and the kite-shaped triplet structure is favoured below 1000 K. Considering $(\text{Al}_2\text{O}_3)_2$, the energy difference of the ground state to the first excited state is relatively large (0.48 eV) and therefore structure 2A (see Fig. 1 in Li & Cheng 2012) is also the most stable in circumstellar conditions. For $n = 3$, we find five stable structures in an energy interval of only 0.13 eV for the alumina trimer. In circumstellar conditions, however, the first excited state 3B (Li & Cheng 2012) is the most stable configuration. For a temperature of 1500 K, the ground and first excited state of $(\text{Al}_2\text{O}_3)_4$ are almost identical in potential energy. Below 1500 K isomer 4A is favoured, above 1500 K structure 4B (Li & Cheng 2012), which is only stable for pressures above 0.1 Pa. Overall, the Gibbs free energy of formation for alumina clusters depends significantly on T , but not much on density or pressure.

For each of the optimized structures, the vibrational spectrum of the ground-state clusters was calculated (see Fig. 13). The predicted spectra compare well with the experimental results by Demyk et al. (2004) and the calculated linear infrared absorption spectra by Sierka et al. (2007) for small cluster sizes (see Fig. 14), with the strongest peaks in the $10-11 \mu\text{m}$ region and an ensemble of smaller peaks around $14 \mu\text{m}$. Implementing the stable clusters in thermo-chemistry calculations for densities down to 1×10^{-4} dyne cm^{-2} (Höfner et al. 2016) and temperatures up to 1800 K and allowing the small clusters to grow via monomer addition indicates that $(\text{Al}_2\text{O}_3)_3$ seems to be the best candidate for possible detection in CSEs (see Table 4).

We tested whether the spectral features present in the infrared SED of R Dor and W Hya could be explained by the vibrational frequencies of these small aluminium oxide clusters (see Fig. 15). However, the low spectral resolution (around 300) of the ISO/SWS speed 1 data for both stars makes it impossible to draw any firm conclusions. Albeit a positive correlation can be noticed between various absorption peaks in the ISO data and the predicted cluster spectra, we caution against over-interpretation

⁵ This value is calculated from the Gibbs free energy. The entropy is given by the partition function (with contributions from electronic, translational, vibrational, and vibrational motion) of which the translational part depends on the pressure.

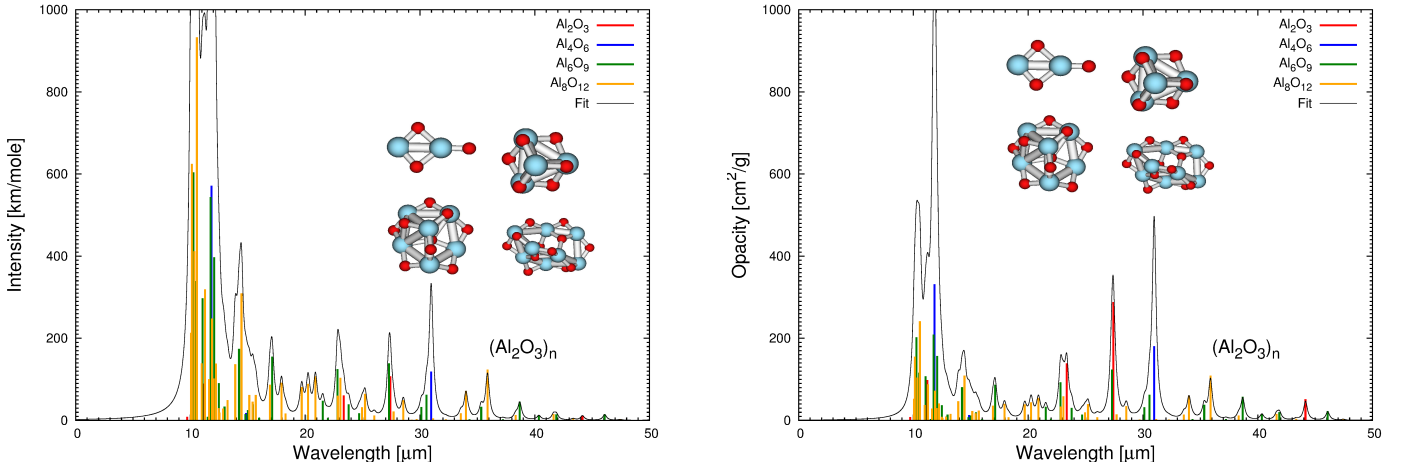


Fig. 13. Vibrational spectrum of the $(\text{Al}_2\text{O}_3)_n$, $n = 1-4$, ground-state clusters and their corresponding structures. The structures (for $n = 1-4$) are depicted with increasing n from left to right and from top to bottom. The black line represents a Lorentzian distribution with HWHM parameter $\gamma = 0.2$ for all peaks. The left panel shows the intensity in units of km/mole; the right panel the opacity in units of cm^2/g .

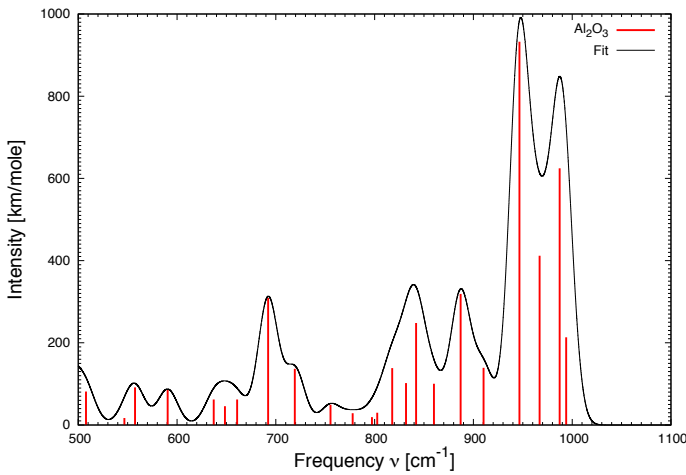


Fig. 14. Intensity of the $(\text{Al}_2\text{O}_3)_4$ ground-state cluster vs. frequency convoluted with a Lorentzian function with FWHM of 24 cm^{-1} . To be compared with Fig. 2d in Sierka et al. (2007).

Table 4. Overview of the alumina nucleation by monomer (Al_2O_3) addition at different temperatures T (in K) and pressures p (in dyne cm^{-2}).

$p \backslash T$	1000	1200	1500	1800
1	✓	✓	✓	$(\text{Al}_2\text{O}_3)_3$
0.1	✓	✓	✓	$(\text{Al}_2\text{O}_3)_3$
0.01	✓	✓	✓	x
10^{-3}	✓	✓	$(\text{Al}_2\text{O}_3)_3$	x
10^{-4}	✓	✓	$(\text{Al}_2\text{O}_3)_3$	x

Notes. An energetically feasible nucleation is denoted with a ✓. Suppressed nucleation with (large) energy barriers is denoted with an x. If the nucleation pathway is partially favourable, the largest preferential cluster is given.

of these results. Specifically, molecular line veiling by other molecules, which might be present in the extra-molecular layer created by pulsation-induced density enhancements, result in an ensemble of emission and absorption features that complicates any potential identification. High spectral resolution data in the

10–20 μm regime (as might be obtained in the future using the E-ELT/METIS instrument) are paramount to answer the question on the potential presence and vibrational excitation of these small aluminium oxide clusters.

4.2. Skewed cluster-size distribution

The more pronounced evidence of large ($n > 34$) $(\text{Al}_2\text{O}_3)_n$ clusters warrants a discussion on the possibility of whether a physico-chemical situation might occur resulting in a skewed cluster-size distribution towards large cluster sizes. Clusters can grow via two direct pathways, i.e. by adding monomers and by sticking together after a collision. Since the number of seeds is low, the chance of colliding with monomers is higher than of collision with larger polymers, so the first pathway seems to be preferred. The region around $2 R_*$ coincides with the GBDS, i.e. a higher density region where species can stay longer without being pushed outwards. This might create a favourable condition for clusters to grow to large sizes. These large aluminium oxides can be very stable at high temperatures because of their ability to undergo thermo-ionic emission rather than dissociation (Demk et al. 2004). That is, the clusters preferentially emit an electron, which is a unique property that is only observed for very strongly bound species. This ability might also greatly enhance the growth efficiencies of the clusters, since the transient complex that is formed during the growth reaction process might also stabilize via the emission of an electron.

Growth by adding monomers is hence a potential process for the creation of larger clusters, with a preference for specific cluster sizes initiating coagulation. Energy considerations in this highly turbulent region just above the stellar atmosphere should be taken into account to answer the question regarding the favourable cluster size for coagulation. There are no calculations, including the whole spectrum from small to large aluminium oxide clusters, currently available to answer this question. Generally, the alumina clusters (including the very small clusters) show an ionic character with alternating Al–O bondings. Moreover, some of the low-lying small clusters exhibit a trigonal symmetry (like bulk-corundum). One thus expects that the coagulation might start at rather small cluster sizes ($n = 3, 4$). Our calculations including only the $n = 1-4$ $(\text{Al}_2\text{O}_3)_n$ clusters indeed show a preference for the $n = 3$ cluster size. A similar behaviour is noted for $(\text{SiO})_n$ for which the matrix isolation

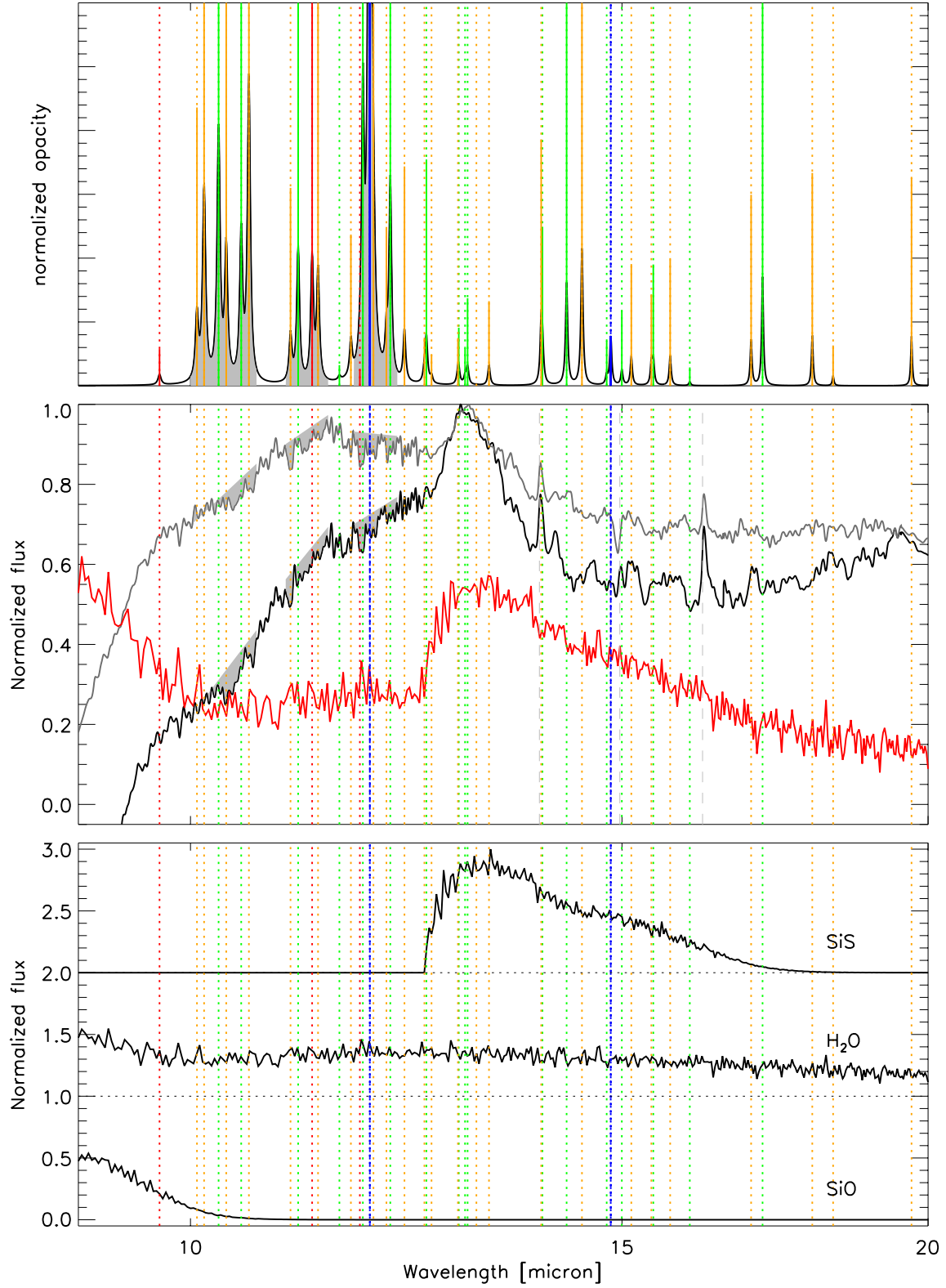


Fig. 15. *Top panel:* normalized opacity of the $(\text{Al}_2\text{O}_3)_n$ clusters convoluted with a Lorentzian profile with a FWHM of $0.03\,\mu\text{m}$. The full coloured lines represent the opacities normalized for each cluster entity separately (with the same colour coding as in Fig. 13), while the dotted coloured lines indicate the wavelength position of all ground-state vibrational frequencies of the stable $(\text{Al}_2\text{O}_3)_n$ ($n = 1-4$) clusters. *Middle panel:* comparison between the (stellar continuum subtracted and normalized) SED of R Dor (full black line) and W Hya (full gray line) with the predicted vibrational frequencies of the stable $(\text{Al}_2\text{O}_3)_n$ ($n = 1-4$) clusters (vertical coloured lines). The gray areas indicate the region with some of the largest $(\text{Al}_2\text{O}_3)_n$ opacities. The dashed gray lines indicate the CO_2 lines as discussed by Justtanont et al. (2004). The full red line represents the (stellar continuum subtracted and normalized) emission for a slab model containing H_2O , SiO , and SiS . The column densities for all species were taken to be $1 \times 10^{20}\,\text{cm}^{-2}$ at $0.5R_*$ above the stellar surface for a temperature of 2000 K. The individual contributions of H_2O , SiO , and SiS are shown in the bottom panel. The column densities of the molecules were not chosen to fit the ISO/SWS data, but serve as illustrative examples on the molecular line veiling.

experiment by Stranz (1980) shows that SiO monomers, dimers, and trimers can be seen, but then the material seems to transition to the bulk phase. If coagulation were to commence from small cluster sizes, then we would not expect to see (much) large clusters, but we assume that these are in fact present, as indicated by the observed $11\ \mu\text{m}$ spectral feature.

Large clusters could, however, be formed by alternative pathways. One possibility is the formation of large clusters by surface processes; i.e. a heterogeneous nucleation process during which large clusters are formed on the surface of existing grains and subsequently ejected. Nozzle experiments show that a substantial quantity of large metal clusters can be formed and ejected that way (Knauer 1987). The NACO and SPHERE data prove that grains (of still unknown composition) are ubiquitous close to the star. These grains might serve as surfaces for cluster growth to occur. Vapourization of solid material might offer another explication for a skewed cluster-size distribution towards large sizes. The Al_2O_3 grain temperatures in the region around $2\ R_\star$ are high (Sect. 3.3) and might lead to the crystallization of the bulk material. If energetically favourable, the material might also get partly vapourized, thereby producing large clusters. This kind of behaviour is noted during formation kinetics experiments and computer simulations for the nucleation and growth rates of metal vapours (Yamada et al. 1987). The recent SPHERE data of Khouri et al. (2016) show that the solid-state material is grouped into irregular, clumpy structures. Backwarming occurring between the dust clumps and the star might locally increase the temperature (Fleischer et al. 1995), favouring vapourization to take place.

5. Conclusions

In this paper we have studied the aluminium content in a low and a high mass-loss rate oxygen-rich AGB star, R Dor and IK Tau, respectively. The aim was to reveal the role of aluminium species in the initiation of the stellar wind. We therefore observed both stars with ALMA at high spatial resolution ($\sim 150\ \text{mas}$) to trace the dust formation region via aluminium bearing molecules (AIO, AIOH, and AlCl). The azimuthally averaged flux densities in function of the angular distance from the central star were fitted using a non-LTE radiative transfer code to retrieve the fractional abundances of the molecules. The gas-phase aluminium chemistry is largely different for both stars, in particular the AIO and AIOH fractional abundances. In IK Tau AIOH is clearly detected, while traces of clumpy structures containing AIO are only apparent beyond $50\ R_\star$. In contrast, AIO is ~ 70 times more abundant than AIOH in R Dor. These three molecules consume only $\lesssim 2\%$ of the total aluminium budget. Hence, there is ample opportunity to form aluminium-bearing dust species. AIO and AIOH are direct precursors of these aluminium-bearing grains and both molecules are detected well beyond the dust condensation radius. This implies that the aluminium dust condensation cycle is not fully efficient.

To assess the aluminium dust content in oxygen-rich AGB stars, we also included the well-studied low mass-loss rate star W Hya. We discuss how the spectral features present in the SED, polarized light signal, and interferometric data point towards the presence of a halo of large (and small) transparent grains close the star. A candidate species for these grains is amorphous Al_2O_3 . However, at a distance of only $1.5\ R_\star$ the estimated grain temperature is $> 1000\ \text{K}$, i.e. too high for the grains to retain their amorphous lattice structure. The grains should anneal and form a crystalline compound. Indeed, the typical $13\ \mu\text{m}$ features signalling the presence of crystalline Al_2O_3 ($\alpha\text{-Al}_2\text{O}_3$) is

seen in the SEDs of W Hya and R Dor. One is then left with the question how to explain the broad $11\ \mu\text{m}$ feature in the SED and interferometric data. On the basis of the experimental work by van Heijnsbergen et al. (2003) and Demyk et al. (2004), we propose that large gas-phase $(\text{Al}_2\text{O}_3)_n$ clusters ($n > 34$) are the species that produce the signatures seen in the SED and interferometric data. Using density functional theory (DFT) we calculate the stability and emissivity of very small $(\text{Al}_2\text{O}_3)_n$ clusters (for $n = 1\text{--}4$). Although we cannot draw firm conclusions on the presence of such small clusters in AGB winds, we notice a positive correlation between various absorption peaks in the ISO data and our predicted spectra. Future E-ELT/METIS observations will be crucial to answer the question of whether very small clusters ($n \lesssim 4$) are present in the inner winds of AGB stars.

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Appendix A: Additional ALMA channel maps in AIO, AIOH, and AICI for IK Tau and R Dor

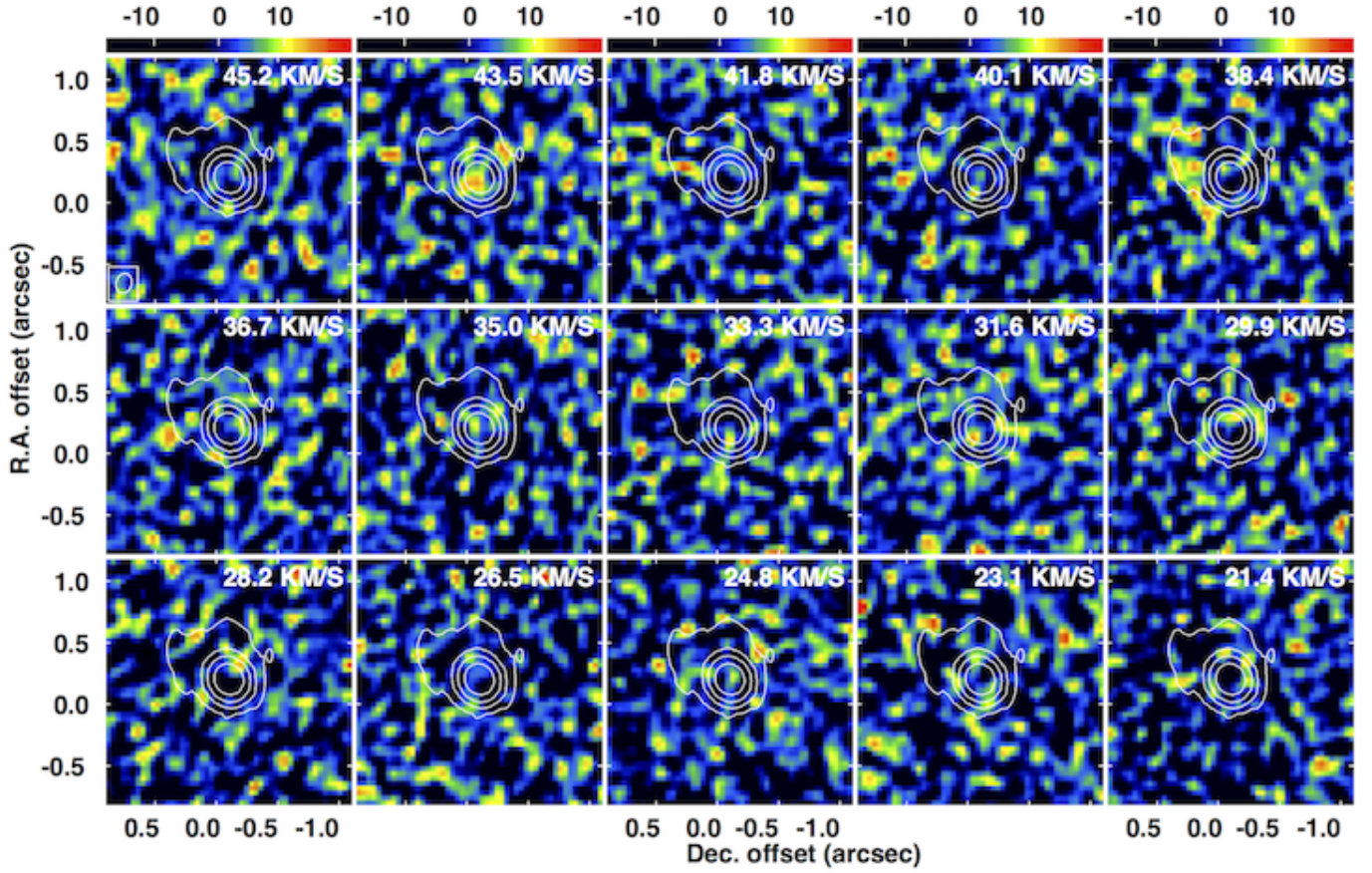


Fig. A.1. Channel maps (colour scale in mJy) of the AIO $N = 9-8$ transition in IK Tau. White contours indicate the continuum at $(-1, 1, 4, 16, 64, 256) \times 0.3$ mJy. The ordinate and co-ordinate axis give the offset of the right ascension and declination, respectively, in units of arcseconds. The ALMA beam size is shown in the bottom left corner of the *top panel*.

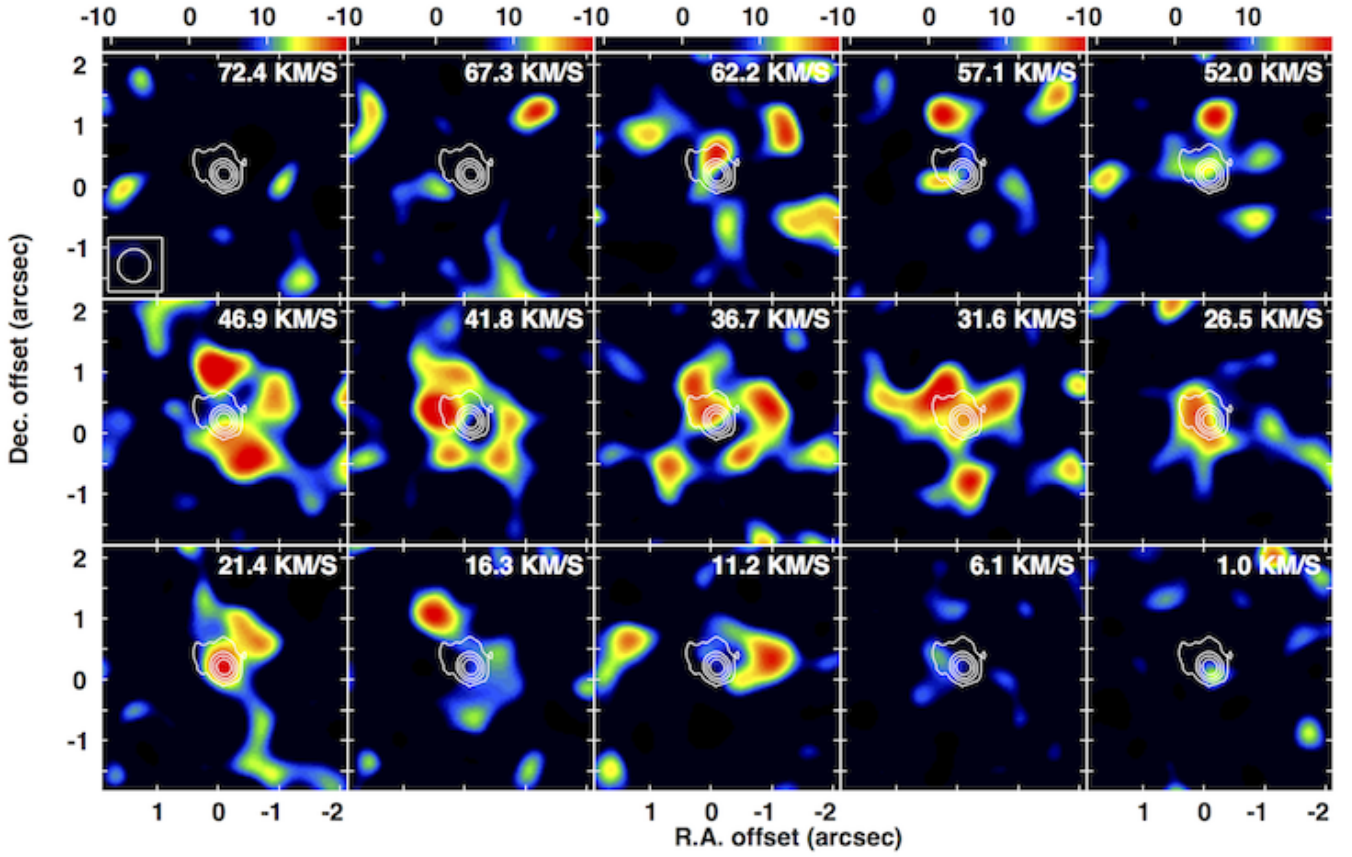


Fig. A.2. Same as for Fig. A.1, but the AlO emission in the wind of IK Tau is averaged over 5 channels and smoothed with 0''.5 Gaussian kernel.

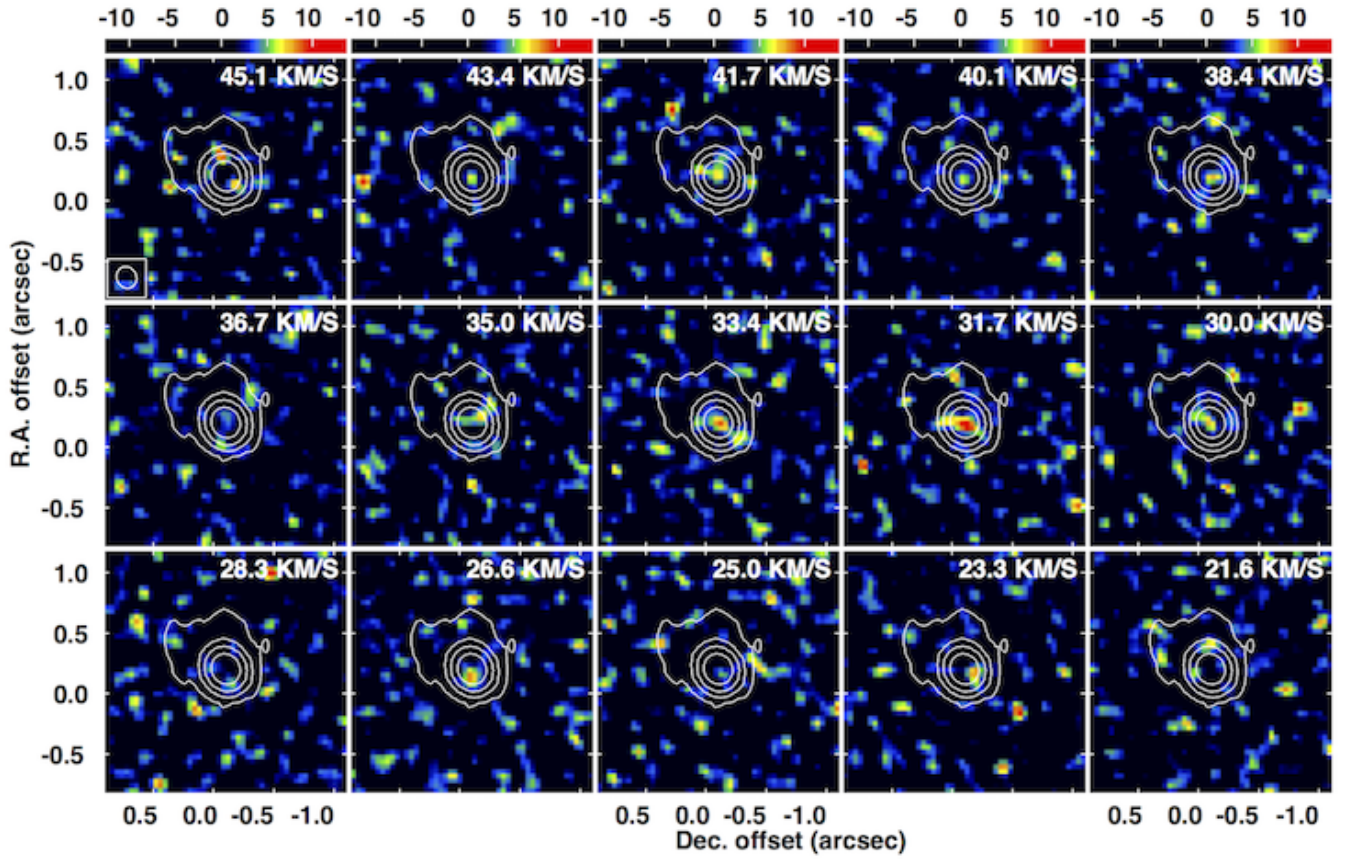


Fig. A.3. Same as for Fig. A.1, but for AlCl $J = 24-23$ in the wind of IK Tau.

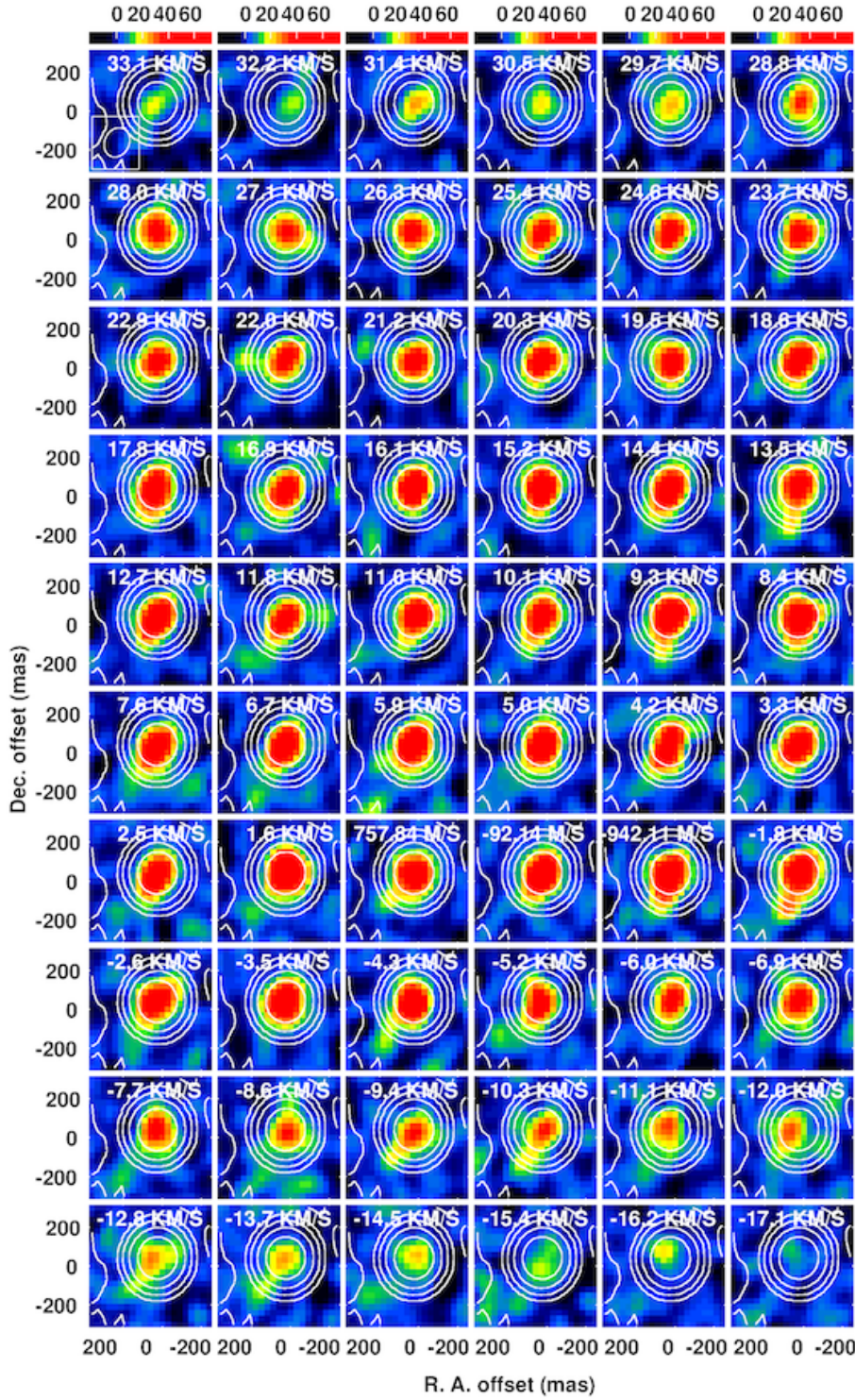


Fig. A.4. Channel maps (colour scale in mJy) of the AIO $N = 9-8$ transition in R Dor, zooming in the inner $0''.2$ region. White contours are the continuum at $(-1, 1, 4, 16, 64, 256) \times 0.15$ mJy. The ordinate and co-ordinate axis give the offset of the right ascension and declination, respectively, in units of arcseconds. The ALMA beam size is shown in the bottom left corner of the *top panel*.

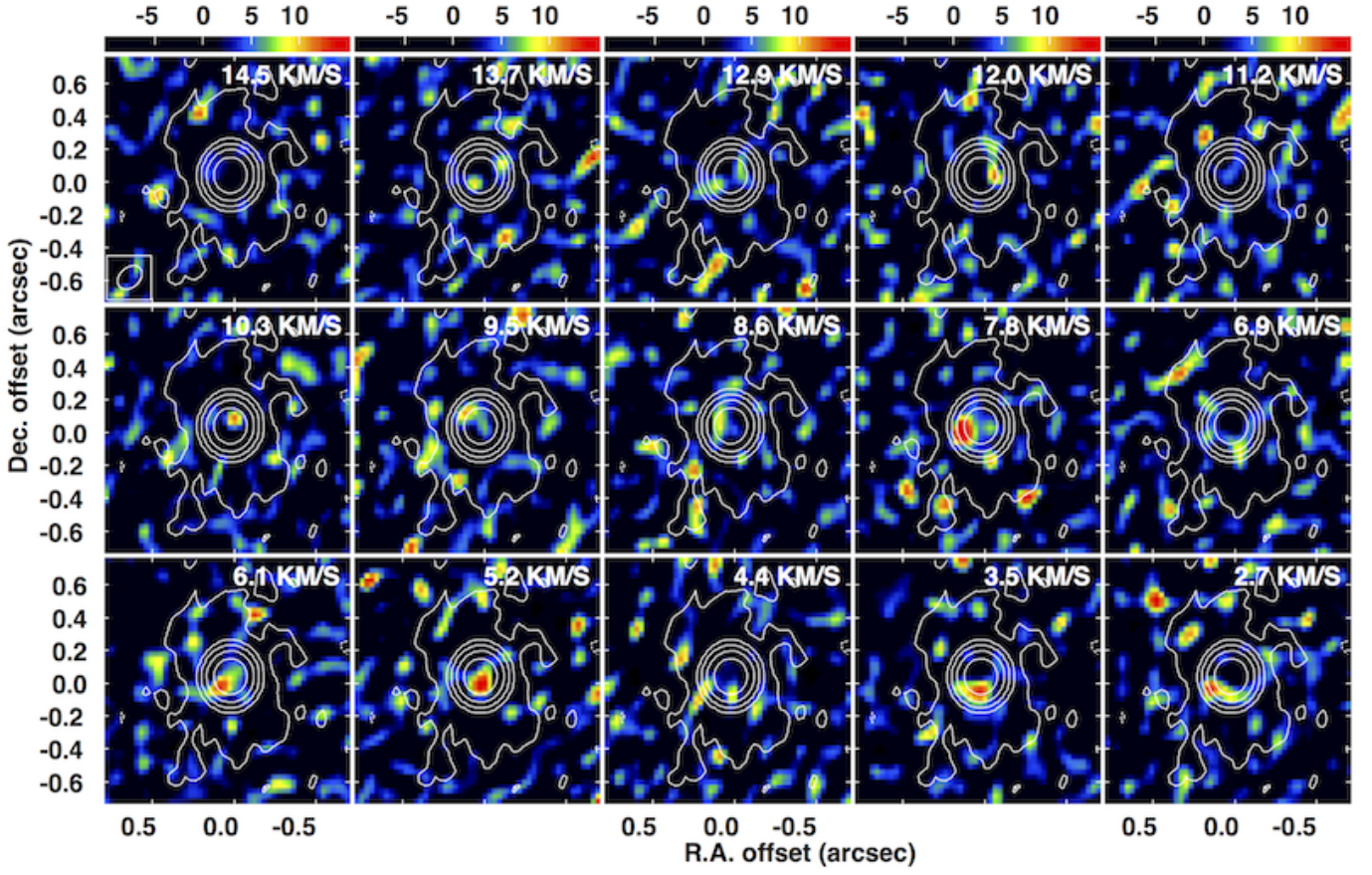


Fig. A.5. Same as for Fig. A.4, but for AlOH $J = 11-10$.

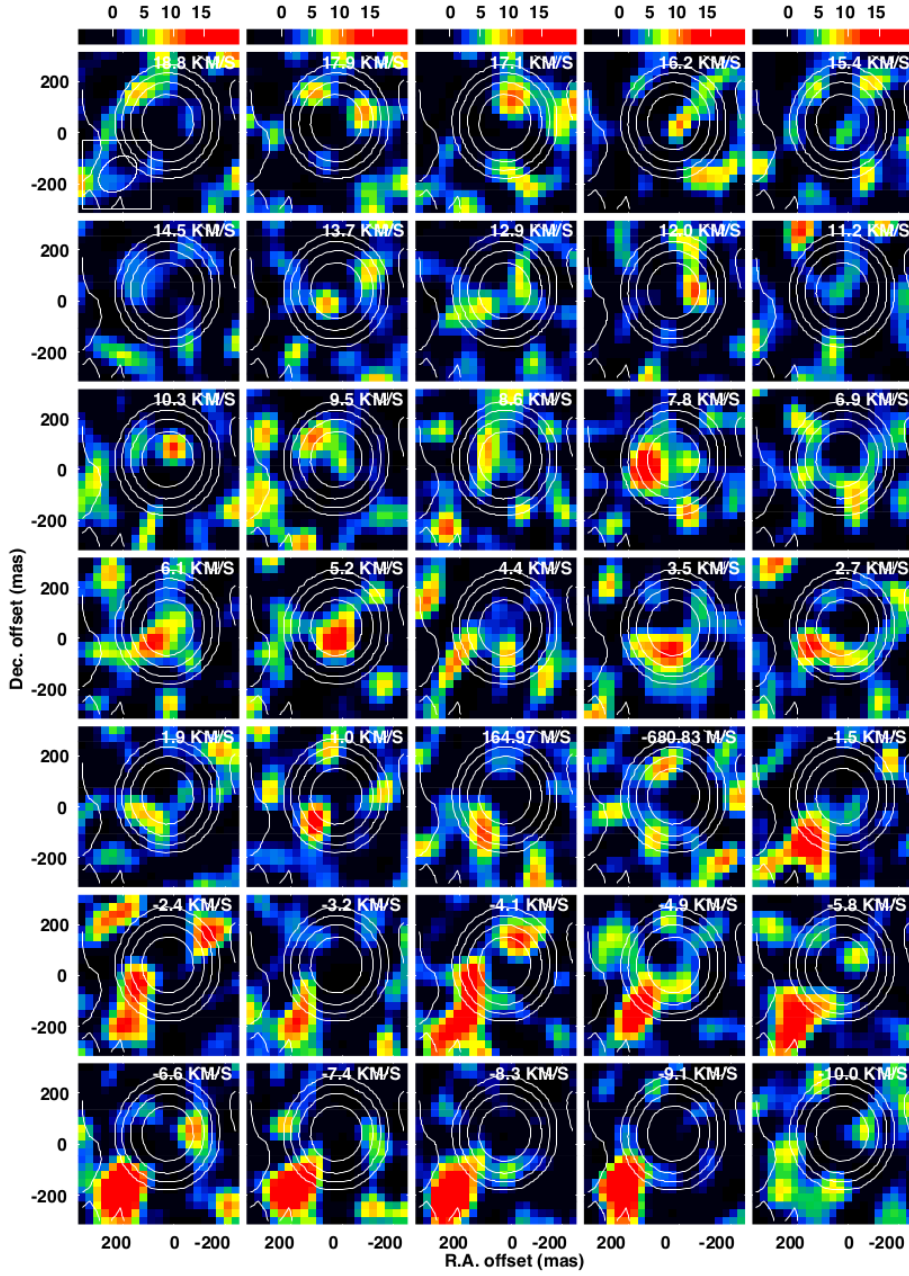


Fig. A.6. Same as for Fig. A.4, but zooming into the AIOH emission in the inner 0''.2 region in the wind of R Dor.

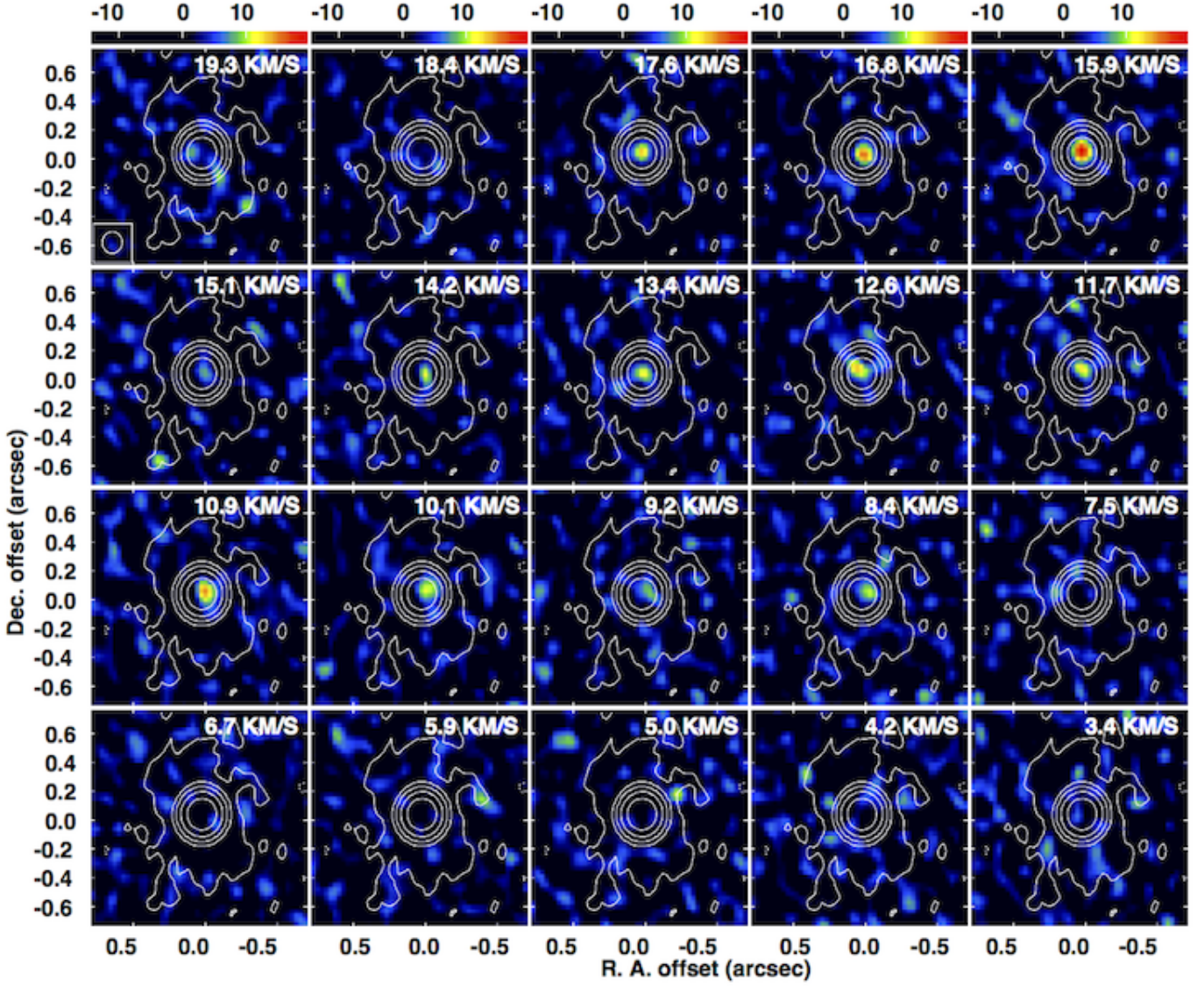


Fig. A.7. Same as for Fig. A.4, but for AlCl $J = 24-23$ in the wind of R Dor.