

H₂O from R Cas: ISO LWS-SWS observations and detailed modelling*

Truong-Bach¹, R.J. Sylvester², M.J. Barlow², Nguyen-Q-Rieu¹, T. Lim³, X.W. Liu², J.P. Baluteau⁴, S. Deguchi⁵, K. Justtanont⁶, and A.G.G.M. Tielens⁷

¹ Observatoire de Paris, DEMIRM and UMR 8540, 61 avenue de l'Observatoire, F-75014 Paris, France

² Department of Physics and Astronomy, University College London, Gower Street, London WC1E 6BT, UK

³ The LWS Instrument-Dedicated Team, ISO Science Operations Center, P.O. Box 50727, E-28080 Madrid, Spain

⁴ Laboratoire d'Astronomie Spatiale, CNRS, BP 8, F-13376 Marseille Cedex 12, France

⁵ Nobeyama Radio Observatory, NAO, Minamimaki, Minamisaku, Nagano 384-13, Japan

⁶ Stockholm Observatory, S-13336 Saltsjöbaden, Sweden

⁷ Kapteyn Astronomical Institute, P.O. Box 800, 9700 AV Groningen, The Netherlands

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Abstract. We present 29–197 μm spectra of the oxygen-rich Mira variable star, R Cas, obtained with the Long- and Short-Wavelength Spectrometers (LWS and SWS) on board the Infrared Space Observatory (ISO). The LWS grating observations were made during two pulsational stellar phases, $\phi \sim 0.5$ and 0.2 in August 1996 and June 1997 when the stellar luminosity was near its minimum and mean values, respectively. The infrared flux at the latter epoch was ~ 30 –40% stronger than at the former. SWS grating observations were also made in June 1997. The spectrum presents a strong far-infrared (FIR) continuum and is rich in water lines suitable for use as circumstellar diagnostics.

We have constructed a circumstellar model which consistently treats radiative transfer, chemical exchanges, photodissociation, and heating and cooling effects. The overall FIR excitation field was scaled by a factor which varied with the stellar phase. By fitting the model to the observed FIR water line fluxes and continuum while adopting the stellar parameters based on the Hipparcos distance we have found a mass-loss rate of $\dot{M} \sim 3.4 \times 10^{-7} M_{\odot} \text{ yr}^{-1}$ and a total *ortho* and *para* water vapour abundance (relative to H₂) of $f \sim 1.1 \times 10^{-5}$. The kinetic temperature and the relative abundances of H₂O, OH, and O in chemical equilibrium have been derived as functions of radial distance r . H₂O excitation is mainly dominated by FIR emitted by dust grains. The deduced model continuum flux at 29–197 μm for the $\phi \sim 0.5$ phase was 61% of the flux at $\phi \sim 0.2$. Photodissociation by the FUV interstellar field and CO cooling effects operate farther out than the H₂O excitation region. Our derived mass-loss rate of R Cas is similar to the value $6 \times 10^{-7} M_{\odot} \text{ yr}^{-1}$ previously published for WHya, another oxygen-rich AGB star.

Key words: stars: individual: R Cas – stars: mass-loss – stars: abundances – stars: AGB and post-AGB – infrared: stars – molecular processes

1. Introduction

R Cas is a normal red giant, classified as an M star because its chemical composition is oxygen-rich, with an abundance ratio $C/O < 1$. It belongs to the Miras, a group of regularly pulsating cool stars. With a period of light variation larger than 1 year (~ 430 days), it is classified as a long-period variable (LPV) star. It pulsates in the fundamental mode while most of LPV stars with periods shorter than 400 days pulsate in an overtone (van Leeuwen et al. 1997).

Mass loss is a fundamental process that drives the star's further evolution on the AGB branch and beyond. Mass loss by AGB stars also dominates the gas and dust injection budgets of the interstellar medium. The mechanism for exciting the Mira emission is not well understood. Bujarrabal et al. (1989) analysed CO ($J=1-0$) emission from R Cas observed with the IRAM 30m radiotelescope. This single low-excitation millimeter wavelength CO line can partially describe the outer part of the circumstellar envelope. Bujarrabal et al. (1989) estimated a stellar distance of 270 pc, i.e., about 2.5 times the recent Hipparcos measurement, and adopted the envelope characteristics of the carbon star IRC+10216 when deriving the physical parameters for R Cas. Thus, new analyses including observations of other molecules such as water vapour around R Cas are certainly of interest.

Because water vapour is abundant in the terrestrial atmosphere, stellar H₂O spectra are often blended with telluric H₂O emission/absorption rendering the astronomical lines unobservable, even with high-altitude airborne telescopes. Use of the Infrared Space Observatory (ISO) telescope can avoid this problem, which is inherent to ground-based observations. Infrared

Send offprint requests to: Truong-Bach (tbach@obspm.fr)

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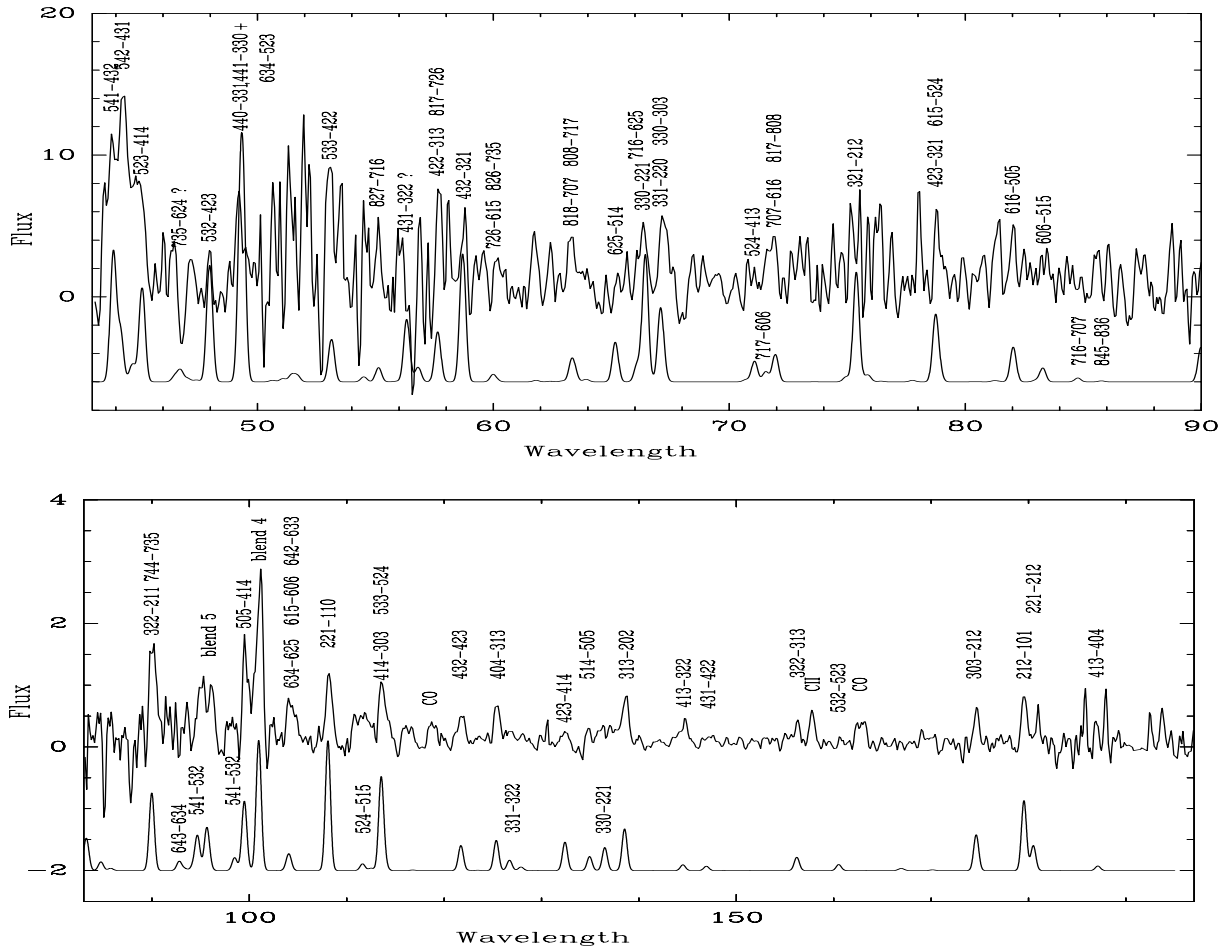


Fig. 1. LWS Observations I (see Table 1 for line fluxes): continuum-subtracted spectrum of R Cas. Our model spectrum is displayed offset for clarity. Units: μm and $10^{-15} \text{ W m}^{-2} \mu\text{m}^{-1}$. For this and subsequent Figures, the model spectrum was produced by convolving the predicted total line fluxes with the appropriate spectral resolution element for the observations

(IR) radiation is emitted mainly from the innermost layers of the circumstellar environments. It is in this hot, high-density and strong radiation field region that atoms and molecules are highly excited. Thus, ISO observations would help us to understand the structure of the circumstellar envelope from the inner layers.

We present here the 29–197 μm ISO (LWS-SWS) spectrum of R Cas obtained in three sets of observations. Our circumstellar model used to fit the observed H₂O line and continuum fluxes enabled us to derive some physical parameters of the stellar envelope: its mass loss rate, radial kinetic temperature, and radial molecular abundances.

2. Observations

Our first set of observations of R Cas, made on 5 August 1996 (hereafter Observations I) during Revolution 263 of the ISO mission, have been reported elsewhere (Truong-Bach et al. 1998, henceforth Paper I). This date corresponds to a stellar phase of 0.5, at minimum stellar luminosity. The spectrum is rather noisy. A second set of observations were obtained on 6 June 1997 (hereafter Observations II) during Revolution 568, using

the same grating LWS01 AOT apparatus and procedure. This date corresponds to a stellar phase of 0.2, when the luminosity was decreasing to its mean value. The total on-target time was 2417 sec for Observations II. As for Observations I, the spectral resolution element is 0.3 μm in second order (SW1–SW5; $\lambda \leq 93 \mu\text{m}$) and 0.6 μm in first order (LW1–LW5; $\lambda \geq 80 \mu\text{m}$) and the spectra were sampled at 1/4 of the resolution element. In order to form a complete spectrum from 43–197 μm , the sub-spectra from the ten LWS detectors were rescaled to give the same flux in regions of overlap. In almost all cases the fluxes were changed by less than 20%.

The merged LWS spectrum, which are continuum subtracted and labelled with some H₂O transitions, are illustrated in Figs. 1–2. The H₂O measured line fluxes for LWS are presented in Tables 1 and 2 with an estimated uncertainty $\sim 40\%$ which increases toward shorter wavelengths. Note that the present Table 1 for Observations I revises and completes that given previously (see Paper I). The FIR emission from R Cas was about 40% stronger in Observations II than in Observations I.

A third set of observations were made during Revolution 564 (on 2 June 1997, hereafter Observations III), using the grating

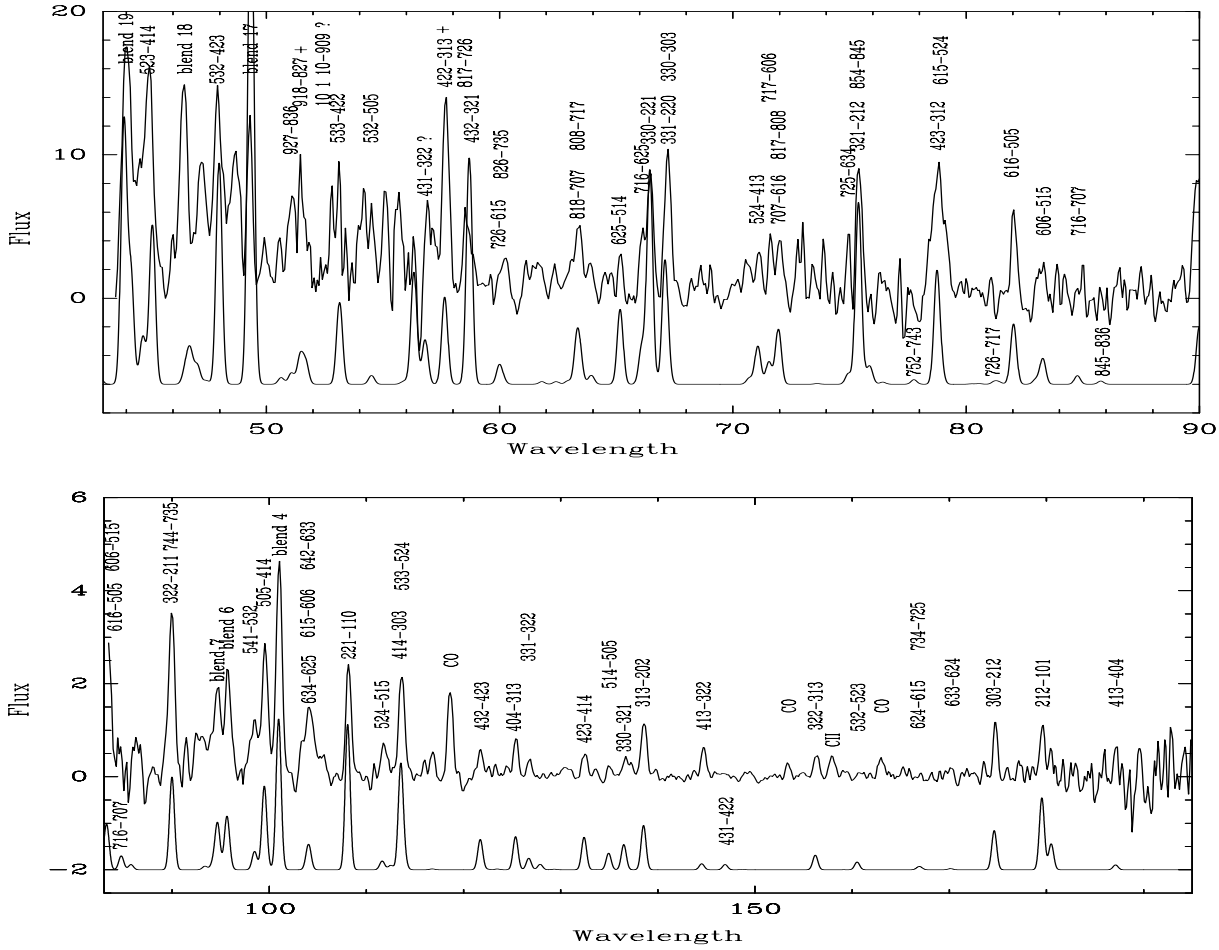


Fig. 2. LWS Observations II (see Table 2 for line fluxes): continuum-subtracted spectrum of R Cas. Our model spectrum is displayed offset for clarity. Units: μm and $10^{-15} \text{ W m}^{-2} \mu\text{m}^{-1}$

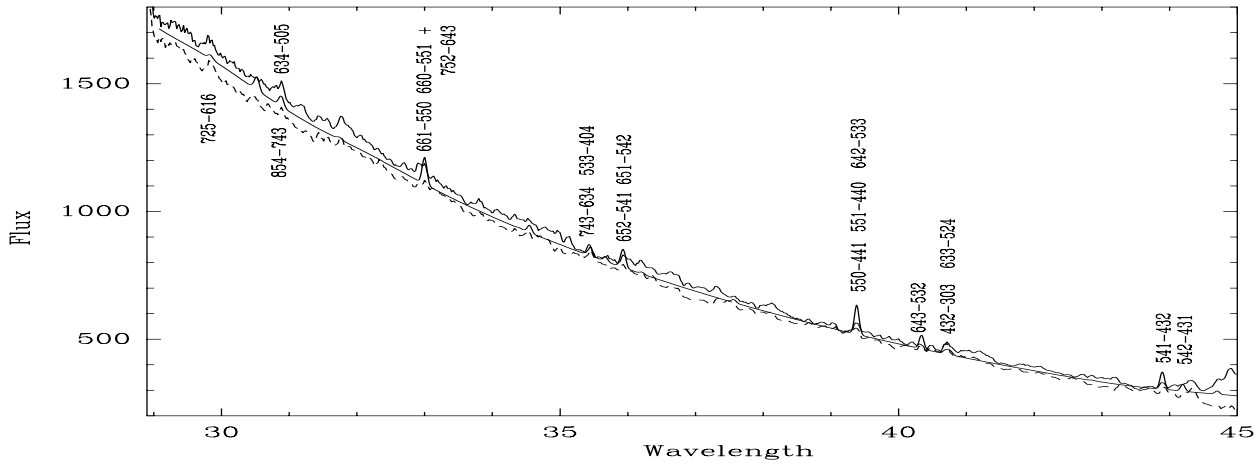


Fig. 3. SWS Observations III: observed forward (*solid*) and backward (*dashed*) spectra; an H₂O model line spectrum is plotted with the average continuum for line identification. Units: μm and $10^{-15} \text{ W m}^{-2} \mu\text{m}^{-1}$

SWS06 AOT. The total on-target time was 1718 sec. The SWS nominal resolution is $\sim 0.03 \mu\text{m}$ in the range 29–45 μm ; however, because of rebinning in data-reduction, the model spectrum was convolved to a resolution lower ($\sim 0.08 \mu\text{m}$) to reproduce the observed linewidths. To judge the SWS data quality, we plot

in Fig. 3 the SWS forward and backward scans with their estimated average continuum superimposed with a set of model H₂O lines for identification. Because of low s/n and blending, the averaged line fluxes (Table 3) can be uncertain by a factor of $\gtrsim 2$.

Table 1. Observations I: calculated (F_{cal1} and F_{cal2} for $r_{\text{min}} = 3.8 \times 10^{14}$ cm and 3×10^{14} cm respectively, see text) and observed (F_{obs}) H₂O line fluxes, for R Cas

J	K_-	K_+	J	K_-	K_+	λ_{vac} (μm)	λ_{obs} (μm)	Det.	F_{cal1} ($10^{-15} \text{ W m}^{-2}$)	F_{cal2} ($10^{-15} \text{ W m}^{-2}$)	F_{obs}	Comments
2	1	2	1	0	1	179.53	179.56	LW4	0.77	0.81	0.60	
3	0	3	2	1	2	174.62	174.64	LW4	0.40	0.43	0.45	
5	2	3	4	3	2	156.27	156.27	LW4	0.02	0.03	0.18	blend 1
3	2	2	3	1	3	156.19	156.27		0.12	0.13		blend 1
4	1	3	3	2	2	144.52	144.60	LW3	0.06	0.07	0.18	noise
3	1	3	2	0	2	138.53	138.59	LW3	0.44	0.46	0.47	
5	1	4	5	0	5	134.93	134.82	LW3	0.15	0.19	0.19	
4	2	3	4	1	4	132.41	132.60	LW3	0.30	0.36	0.12	noise
3	3	1	3	2	2	126.71	126.94	LW2	0.11	0.12	0.10	noise
4	0	4	3	1	3	125.35	125.36	LW3	0.32	0.34	0.40	
4	3	2	4	2	3	121.72	121.76	LW2	0.26	0.33	0.20	
5	3	3	5	2	4	113.95	113.74	LW2	0.06	0.09	0.82	blend 2
4	1	4	3	0	3	113.54			0.98	1.15		blend 2
7	4	3	7	3	4	112.51	112.17	LW2	0.02	0.05	0.17	
5	2	4	5	1	5	111.63	111.22	LW2	0.07	0.10	0.29	
2	2	1	1	1	0	108.07	108.20	LW2	1.43	1.55	1.03	
6	3	4	6	2	5	104.09	103.96	LW2	0.14	0.21	0.60	blend 3 noise
6	1	5	6	0	6	103.94			0.03	0.06		blend 3
6	4	2	6	3	3	103.92			0.02	0.04		blend 3
6	2	4	5	3	3	101.21	101.01	LW1	0.02	0.03	1.96	blend 4
2	2	0	1	1	1	100.98			1.02	1.08		blend 4
5	1	4	4	2	3	100.91			0.36	0.47		blend 4
5	0	5	4	1	4	99.49	99.65	LW1	0.73	0.90	0.84	
4	4	0	4	3	1	95.88	95.42	LW1	0.06	0.08	0.72	blend 5
5	1	5	4	0	4	95.63			0.47	0.51		blend 5
7	4	4	7	3	5	90.05	90.08	LW1	0.01	0.03	1.08	blend 6
3	2	2	2	1	1	89.99			0.81	0.89		blend 6
6	1	5	5	2	4	78.93	78.83	SW4	0.14	0.23	2.08	blend 7
4	2	3	3	1	2	78.74			1.55	1.90		blend 7
3	2	1	2	1	2	75.38	75.40	SW4	2.66	3.01	1.50	
7	0	7	6	1	6	71.95	71.78	SW3	0.65	0.97	2.56	blend 8
7	1	7	6	0	6	71.54			0.24	0.41		blend 8
5	2	4	4	1	3	71.07			0.48	0.65		blend 8
3	3	0	3	0	3	67.27	67.17	SW3	0.09	0.11	1.87	blend 9
3	3	1	2	2	0	67.09			1.70	1.90		blend 9
3	3	0	2	2	1	66.44	66.30	SW3	2.95	3.55	1.74	
8	0	8	7	1	7	63.46	63.35	SW3	0.14	0.31	1.61	blend 10
8	1	8	7	0	7	63.32			0.48	0.86		blend 10
4	3	2	3	2	1	58.70	58.84	SW2	2.97	3.69	1.25	
5	3	3	4	2	2	53.14	53.04	SW2	0.99	1.34	2.51	
7	3	5	6	2	4	46.75	46.43	SW1	0.25	0.51	2.55	blend 11 noise+x?
10	2	9	9	1	8	46.61			0.06	0.25		blend 11
6	3	3	6	0	6	46.54			0.01	0.03		blend 11
6	4	2	6	1	5	46.54			0.01	0.02		blend 11
3	3	1	2	0	2	46.48			0.09	0.10		blend 11
5	4	1	4	3	2	43.89	43.62	SW1	3.03	4.20	3.84	

3. A model for H₂O emission

Our previous modelling of H₂O emission from W Hya (Barlow et al. 1996) (i) neglected excited vibrational states, (ii) did not use the observed FIR continuum as a constraint (iii) adopted the kinetic temperature distribution of Goldreich & Scoville

(1976, hereafter GS) (iv) assumed a two-shell model of H₂O abundances.

The present treatment abandons these assumptions. Here we add the first vibrational state to allow for 4–10 μm photon excitation. The H₂O abundance starts from a guessed value at the inner radius with an initial ortho/para H₂O ratio of 3. We solve

Table 2. Observations II: calculated (F_{cal1} and F_{cal2} for $r_{\text{min}}=3.8 \times 10^{14}$ cm and 3×10^{14} cm respectively, see text) and observed (F_{obs}) H₂O line fluxes for R Cas

J	K_-	K_+	J	K_-	K_+	λ_{vac} (μm)	λ_{obs} (μm)	Det.	F_{cal1} ($10^{-15} \text{ W m}^{-2}$)	F_{cal2} ($10^{-15} \text{ W m}^{-2}$)	F_{obs}	Comments
2	1	2	1	0	1	179.53	179.55	LW5	0.97	1.01	0.82	
3	0	3	2	1	2	174.62	174.66	LW4	0.52	0.55	0.54	
5	2	3	4	3	2	156.27	156.28	LW4	0.04	0.04	0.24	blend 1
3	2	2	3	1	3	156.19			0.17	0.17		blend 1
4	1	3	3	2	2	144.52	144.56	LW4	0.08	0.09	0.29	+ CO 18–17
3	1	3	2	0	2	138.53	138.53	LW3	0.61	0.61	0.74	
3	3	0	3	2	1	136.50	136.62	LW3	0.34	0.38	0.18	
5	1	4	5	0	5	134.94	134.94	LW3	0.23	0.28	0.18	
4	2	3	4	1	4	132.41	132.38	LW3	0.44	0.50	0.35	
3	3	1	3	2	2	126.71	126.74	LW3	0.16	0.17	0.14	
4	0	4	3	1	3	125.35	125.22	LW3	0.45	0.48	0.41	
4	3	2	4	2	3	121.72	121.78	LW2	0.41	0.48	0.31	
7	3	4	6	4	3	116.78	116.79	LW2	0.01	0.02	0.36	+x?
5	3	3	5	2	4	113.95	113.62	LW2	0.11	0.15	1.54	blend 2
4	1	4	3	0	3	113.54			1.42	1.54		blend 2
5	2	4	5	1	5	111.63	111.76	LW2	0.12	0.15	0.27	
2	2	1	1	1	0	108.07	108.04	LW2	1.95	2.07	1.50	
6	3	4	6	2	5	104.09	104.23	LW1	0.25	0.35	0.68	blend 3
6	1	5	6	0	6	103.94			0.07	0.10		blend 3
6	4	2	6	3	3	103.92			0.04	0.07		blend 3
6	2	4	5	3	3	101.21	101.06	LW1	0.04	0.05	2.57	blend 4
2	2	0	1	1	1	100.98			1.50	1.51		blend 4
5	1	4	4	2	3	100.91			0.56	0.67		blend 4
8	2	6	8	1	7	99.98	99.57	LW1	0.02	0.04	1.43	blend 5
5	0	5	4	1	4	99.49			1.14	1.30		blend 5
5	4	1	5	3	2	98.49	98.49	LW1	0.25	0.34	0.36	
4	4	0	4	3	1	95.88	95.73	LW1	0.10	0.13	1.21	blend 6
5	1	5	4	0	4	95.63			0.67	0.78		blend 6
4	4	1	4	3	2	94.71	94.70	LW1	0.33	0.42	0.98	blend 7
6	2	5	6	1	6	94.64			0.31	0.43		blend 7
7	4	4	7	3	5	90.05	89.81	LW1	0.04	0.07	2.54	blend 8 + x?
3	2	2	2	1	1	89.99			1.24	1.31		blend 8
6	0	6	5	1	5	83.28	83.28	SW5	0.57	0.75	0.78	
6	1	6	5	0	5	82.03	82.06	SW4	1.34	1.64	1.36	
6	1	5	5	2	4	78.93	78.81	SW5	0.27	0.40	2.94	blend 9
4	2	3	3	1	2	78.74			2.42	2.74		blend 9
8	5	4	8	4	5	75.50	75.40	SW4	0.06	0.14	3.0	blend 10
3	2	1	2	1	2	75.38			3.92	4.24		blend 10
7	2	5	6	3	4	74.95	74.94	SW4	0.24	0.38	0.64	
8	1	7	8	0	8	72.03	72.01	SW3	0.03	0.07	1.36	blend 11
7	0	7	6	1	6	71.95			1.20	1.61		blend 11
7	1	7	6	0	6	71.54	71.62	SW3	0.50	0.76	1.18	
5	2	4	4	1	3	71.07	71.11	SW3	0.85	1.07	1.01	
3	3	0	3	0	3	67.27	67.20	SW3	0.13	0.14	3.16	blend 12
3	3	1	2	2	0	67.09			2.72	2.94		blend 12
3	3	0	2	2	1	66.44	66.47	SW3	4.68	5.20	3.07	
7	1	6	6	2	5	66.09	66.09	SW3	0.83	1.24	1.08	
6	2	5	5	1	4	65.17	65.17	SW3	1.67	2.21	0.61	noisy
8	0	8	7	1	7	63.46	63.31	SW3	0.36	0.65	1.57	blend 13
8	1	8	7	0	7	63.32			1.03	1.59		blend 13
8	2	6	7	3	5	60.16	60.16	SW3	0.08	0.17	1.59	blend 15 noise
7	2	6	6	1	5	59.99			0.41	0.70		blend 15
4	3	2	3	2	1	58.70	58.58	SW3	4.98	5.70	2.31	
8	1	7	7	2	6	57.71	57.70	SW2	0.21	0.44	2.76	blend 16
4	2	2	3	1	3	57.64			1.76	1.93		blend 16

Table 2. (Continued)

J	K_-	K_+	J	K_-	K_+	λ_{vac} (μm)	λ_{obs} (μm)	Det.	F_{cal1} ($10^{-15} \text{ W m}^{-2}$)	F_{cal2}	F_{obs}	Comments
4	3	1	3	2	2	56.33	56.81	SW2	2.49	2.89		noisy
8	2	7	7	1	6	55.13	55.10	SW3	0.83	1.48	0.85	noisy
5	3	3	4	2	2	53.14	53.08	SW3	1.82	2.30	1.32	
9	1	8	8	2	7	51.69	51.47		0.47	1.04		noisy
6	3	4	5	2	3	49.39	49.31	SW1	3.07	4.12	4.10	blend 17 noisy
4	4	1	3	3	0	49.34			5.89	6.99		blend 17
4	4	0	3	3	1	49.28			3.03	3.74		blend 17
5	3	2	4	2	3	47.97	47.93	SW1	4.92	6.09	3.28	
7	3	5	6	2	4	46.75	46.44	SW1	0.62	1.08	5.60	blend 18 noise/+x?
10	2	9	9	1	8	46.61			0.27	0.75		blend 18
6	3	3	6	0	6	46.54			0.03	0.05		blend 18
6	4	2	6	1	5	46.54			0.03	0.05		blend 18
3	3	1	2	0	2	46.48			0.13	0.14		blend 18
5	2	3	4	1	4	45.11	45.00	SW1	3.53	4.28	3.39	
5	4	2	4	3	1	44.20	43.98	SW1	2.35	3.24	7.37	blend 19
7	4	3	7	1	6	44.05			0.09	0.16		blend 19
5	4	1	4	3	2	43.89			5.75	7.30		blend 19

Table 3. Observations III: calculated (F_{cal1} and F_{cal2} for $r_{min} = 3.8 \times 10^{14}$ cm and 3×10^{14} cm respectively, see text) and observed H₂O line fluxes from R Cas

J	K_-	K_+	J	K_-	K_+	λ_{vac} (μm)	λ_{obs} (μm)	Det.	F_{cal1} ($10^{-15} \text{ W m}^{-2}$)	F_{cal2}	F_{obs}	Comments
5	4	2	4	3	1	44.195	43.98	SWS	2.35	3.24	7.4	blend 1
7	4	3	7	1	6	44.049			0.09	0.16		blend 1
5	4	1	4	3	2	43.893			5.75	7.30		blend 1
6	3	3	5	2	4	40.760	40.71	SWS	1.31	2.01	5.3	blend 2
4	3	2	3	0	3	40.691			1.25	1.48		blend 2
6	4	2	5	3	3	39.399	39.38	SWS	1.90	2.94	3.9	blend 3
5	5	1	4	4	0	39.380			2.46	3.71		blend 3
5	5	0	4	4	1	39.375			5.71	7.73		blend 3
7	4	4	6	3	3	37.566	37.55	SWS	0.98	1.82	2.2	
6	5	2	5	4	1	35.938	35.94	SWS	4.86	7.32	3.5	blend 4 noise
6	5	1	5	4	2	35.904			1.78	3.09		blend 4
5	3	3	4	0	4	35.472	35.45	SWS	0.49	0.65	2.5	blend 5
7	4	3	6	3	4	35.428			3.45	5.61		blend 5
7	3	4	6	2	5	34.549	34.54	SWS	2.40	3.98	2.4	
6	6	1	5	5	0	33.005	33.01	SWS	4.13	6.96	5.6	blend 6
6	6	0	5	5	1	33.005			1.50	2.91		blend 6
7	5	2	6	4	3	32.991			3.55	6.16		blend 6
4	4	1	3	1	2	31.772	31.77	SWS	0.13	0.17	5.1	blend 7
8	4	4	7	3	5	31.738			0.60	1.33		blend 7
6	3	4	5	0	5	30.899	30.89	SWS	1.52	2.22	4.2	blend 8
8	5	4	7	4	3	30.871			2.16	4.48		blend 8
5	4	2	4	1	3	29.885	29.85	SWS	0.10	0.15	5.0	blend 9
7	2	5	6	1	6	29.837			1.20	1.97		blend 9

radiative transfer and thermal balance to derive a radial kinetic temperature $T_K(r)$, H₂O line fluxes, and the FIR continuum consistently in order to compare with the observations.

For R Cas, the Hipparcos parallax (van Leeuwen et al. 1997) corresponds to a distance of 107 ± 12 pc and $M_{bol} = -3.74$. Adopting the angular diameter E-model of Haniff et al. (1995)

gives: the stellar radius $r_* \sim 2 \times 10^{13}$ cm, stellar effective temperature $T_{eff} \sim 2415$ K, stellar luminosity $L \sim 2420 L_\odot$, and stellar mass $M \sim 1.2 M_\odot$. From CO observations, the outflow terminal velocity is $V_t = 12 \text{ km s}^{-1}$ (Bujarrabal et al. 1989, Loup et al. 1993). We assumed an atomic-to-molecular hydrogen ratio of $f = 1$. Collisional rates of *ortho*- and of *para*-H₂O

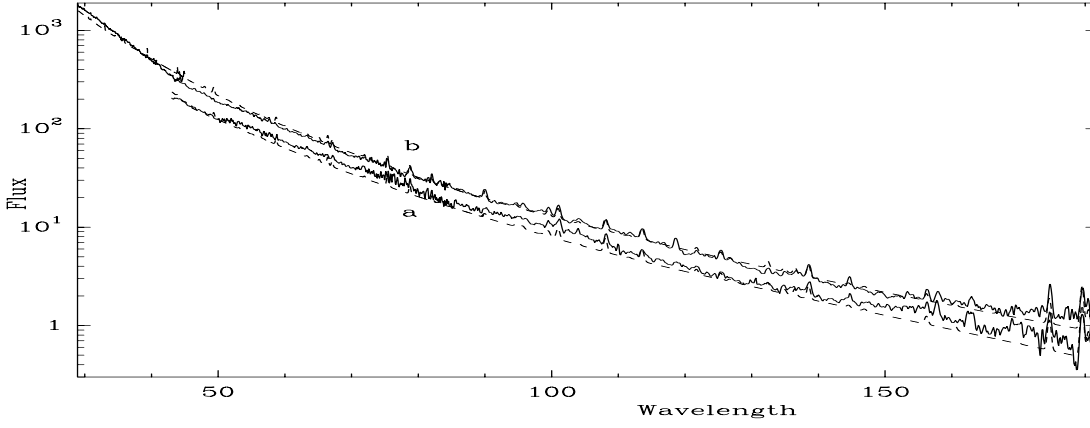


Fig. 4. Merged and smoothed FIR continuum spectra of: (a) Observations I; (b) Observation II-III. *Dotted*, model continuum spectra. Units: μ and $10^{-15} \text{ W m}^{-2} \mu\text{m}^{-1}$

Table 4. A 10-point sample of the observed FIR continuum fluxes used for the fit and their corresponding calculated values (in $10^{-15} \text{ W m}^{-2} \mu\text{m}^{-1}$)

λ (μ)	29	40	43	46	50	62	72	86	108	130	132	155	185	190
Observations I			200	165	125	65	37	16	5.7	2.6		1.3	0.4	
Calculated			240	186	135	58	31	15	5.6	2.5		1.1	0.4	
Observations II-III	1750	500			180	90	50	23	9		3.5	1.75		0.7
Calculated	1559	514			223	95	51	24	9.2		4.0	1.85		0.7

excited by He impact (Green et al. 1993) were corrected by a factor ~ 1.38 to account for H₂ encounters. We treat 45 rotational levels of the ground vibrational state, ν_0 , whose highest energy is ~ 2000 K. To account for $\sim 4\text{--}10 \mu$ photon excitation, as suggested by Barlow et al. (1996), we considered also the 45 lowest rotational levels of the first excited vibrational state, ν_2 . We assumed that ν_2 and ν_0 have the same rotational collision rates, $C(JK_-K_+ - J'K'_-K'_+)$, while between ν_2 and ν_0 the vib-rotational collision rates are lower by a factor ~ 0.01 (Deguchi & Nguyen-Q-Rieu 1990; hereafter D-NQR). This factor was subsequently varied by a factor of 10 to determine its effects. As in D-NQR, we found that uncertainties in the vibrational collisional rates have no effect on the calculated H₂O line fluxes.

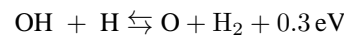
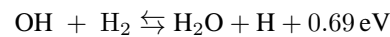
The FIR continuum radiation comes from the central star, the 2.7 K background, and dust. It was scaled by a factor, W_s , to account for the stellar phase dependence. The dust temperature varies as $r^{-0.4}$ (Elitzur et al. 1976). The FIR emissivity of dust grains was taken to vary as $(80/\lambda(\mu\text{m}))^q$ (Bujarrabal et al. 1980), where we have found $q \sim 0.80$ is appropriate to reproduce the observed wavelength dependence of the FIR continuum spectra (Fig. 4). We adopted a grain radius of $r_d=0.1 \mu\text{m}$ and a dust mass density $\rho_d=3 \text{ g cm}^{-3}$. A dust-to-gas mass ratio of 0.005 seems to be appropriate (Groenewegen et al. 1998). If n_d is the grain number density, and σ_d is the grain geometrical cross section, uncertainties in the grain characteristics $n_d\sigma_d$ can be absorbed in the momentum transfer efficiency factor Q , a free parameter in the dust absorption coefficient, which is used to fit the observed H₂O line fluxes and continuum:

$$k(\text{cm}^{-1}) = Qn_d\sigma_d = -\frac{d\tau_d}{dz}.$$

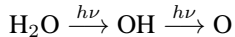
Here $\tau_d(p, z)$ is the averaged optical depth of dust grains at $80 \mu\text{m}$ from $-\infty$ to z along the line of sight at impact parameter p ; p and z are the components of the radius vector \mathbf{r} in the Castor (1970) coordinate system ($p=0$ and $z=r$ for the on-source direction).

In order to reduce the calculation time, we have fitted the model to a sample of 10 points of the observed FIR continuum spectrum (see Table 4). The entire continuum distribution can be obtained by interpolation, using a spline function (see Fig. 4).

Since the *ortho* states do not mix with the *para* states (Herzberg 1962, Townes & Schawlow 1975), we have distinguished two systems whose transition probabilities, collisional rates, and statistical weights [$3(2J+1)$ for an *ortho* and $2J+1$ for a *para* level J] are different. We assumed an initial *ortho*-to-*para* H₂O abundance ratio of 3, which is expected to be equal to the high-temperature limit for H₂O molecules newly formed on a grain surface (Tielens & Allamandola 1987). The gas velocity, $v_e(r)$, was obtained by solving the equation of motion for the ejected gas (GS). The original expression for the drift velocity, $v_d(r)$, of the moving grains relative to the gas particles is not valid close to the stellar atmosphere (GS). We used a modified expression for $v_d(r)$ (Truong-Bach et al. 1991) based on the equation of motion of the grains (Kwok 1975). The molecular densities of H₂O, OH, and O were derived from the exothermic chemical equilibrium reactions



Photodissociation by the interstellar UV field operates in the outer envelope according to



Here the unshielded photodissociation rates were taken to be (Roberge et al. 1991)

$$G(\text{H}_2\text{O}) = 5.9 \cdot 10^{-10} \text{ s}^{-1}$$

$$G(\text{OH}) = 3.5 \cdot 10^{-10} \text{ s}^{-1}$$

H₂O self-shielding is negligible. Dust shielding against the UV interstellar field is represented by a dust opacity at 1000 Å (Morris & Jura 1983). Because the local line width due to turbulence and thermal broadening ($\sim 0.3\text{--}0.7 \text{ km s}^{-1}$) is much lower than the terminal velocity (12 km s^{-1}), one can expect that photons which escape from a local volume are not reabsorbed, and use the escape probability formalism (Castor 1970) to perform the statistical equilibrium calculations. The deduced level populations were used to determine molecular line opacities, excitation temperatures, and fluxes. We have included heating and cooling and photodissociation by FUV interstellar field (e.g. Truong-Bach et al. 1990, 1991) together with chemical reactions (GS) in our calculations.

Heating of gas is due to collisions with dust grains and also by exothermic (chemical and photodissociating) reactions. Cooling is due to: (i) the gas adiabatic expansion (GS); (ii) H₂O spontaneous emission between rotational levels (of ν_0 and ν_2) whose fractional level populations are determined by solving the statistical equilibrium equations (see Truong-Bach et al. 1990); (iii) H₂ decay from the first excited to the ground vibrational state (GS). We also estimated cooling by CO emission in a three-level scheme: two rotational levels in the ground vibrational state and one in the excited vibrational state (e.g. GS; Justtanont et al. 1994). With a CO/H₂ abundance of 4×10^{-4} at the inner radius (Mamon et al. 1988), we found that CO cooling operates farther out than the H₂O excitation region. H₂O molecules formed in the vibrationally-excited state (and the reverse, dissociation, reaction) were found to have a negligible effect on the energy balance of the envelope, in agreement with the results of Justtanont et al. (1994).

The net energy variation is related to the temperature gradient throughout the shell. Thus, when the best fit to the observations was achieved, the molecular abundances and densities, the heating and cooling rates, and the kinetic temperature, $T_K(r)$, were simultaneously derived.

4. Results and discussion

In Figs. 1 and 2 we plot the H₂O model spectra superimposed on the observed spectra of observations I and II for comparison. The fit for the long-wavelength portion (i.e. detectors LW1–5) of the LWS spectrum (lower panel of Figs. 1 and 2) seems to be better than for the short-wavelength detectors (upper panels). This is, to a large extent, due to the high continuum/line flux ratio toward short wavelengths, causing the difficulty in determining the baseline among the continuum fluctuations. Fig. 4 represents

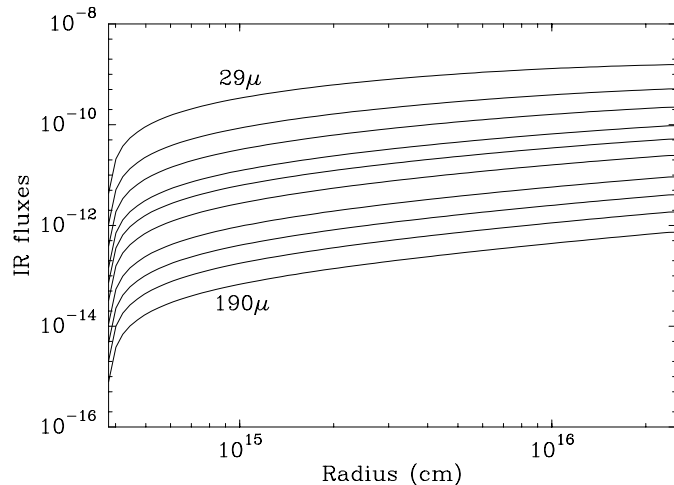


Fig. 5. The FIR fluxes (in $\text{W m}^{-2} \mu\text{m}^{-1}$) at 29, 40, 50, 62, 72, 86, 108, 130, 155 and $190 \mu\text{m}$ as functions of radius (from top to bottom, respectively)

the spectra with continuum: the rapid rise of the continuum toward $29 \mu\text{m}$, the low line/continuum ratios, and blending are the main causes of uncertainty. At some wavelength positions, the forward and backward scans of SWS (Fig. 3, see Sect. 2) do not clearly show the same lines when the continuum fluctuations can be comparable to the line heights. Uncertainties in line fluxes, determined after continuum removal, are estimated to be better than $\sim 40\%$ for LW. They increase toward SW and can reach a factor 2–3 for SWS. The measured line fluxes are listed in Tables 1–3 (see Sect. 2).

Our best fit to the observed FIR continuum and H₂O line fluxes (Observations I–III) was obtained with a mass loss rate of $\dot{M} \sim 3.4 \times 10^{-7} M_{\odot} \text{ yr}^{-1}$, and a H₂O total initial relative abundance of $\sim 1.1 \times 10^{-5}$ (7.9×10^{-6} for *ortho*- and 2.7×10^{-6} for *para*-H₂O). The scaling factor for FIR excitation, W_s (Sect. 3), was found to be ~ 0.61 and 1 for Observations I and II–III, respectively. SW and SWS lines are excited closer to the star ($r \lesssim 1.4 \times 10^{15} \text{ cm}$ for line $\lambda 45.11 \mu\text{m}$) than LW's ($r \lesssim 6.3 \times 10^{15} \text{ cm}$ for line $\lambda 174.62 \mu\text{m}$). Increasing the mass loss rate by 20% would lead to an increase of the line fluxes of $\sim 25\%$ and 40% for the $174.62 \mu\text{m}$ and $45.11 \mu\text{m}$ lines, respectively. In an outward integration over a large range of radius, from $r = 1.0 \times 10^{14} \text{ cm}$ to $2.5 \times 10^{16} \text{ cm}$, we determined the contributions to the line and continuum fluxes at each step. This allowed us to estimate an overall best fit for $r_{\text{min}} = 3.8 \times 10^{14} \text{ cm}$. A smaller value of r_{min} would overestimate the SWS+SW line fluxes, as reported in Tables 1 and 2. Here H₂O line fluxes, F_{calc1} and F_{calc2} , were calculated for 2 values of r_{min} : $\sim 3.8 \times 10^{14} \text{ cm}$ and $3.0 \times 10^{14} \text{ cm}$, respectively. In the following we adopt $r_{\text{min}} = 3.8 \times 10^{14} \text{ cm}$. We note that most of the H₂O line excitation arises at radii $\lesssim 5 \times 10^{15} \text{ cm}$. Within observational uncertainty, an outer radius r_{max} chosen greater than $1 \times 10^{16} \text{ cm}$ makes little difference to the line fluxes. The outer shell from $r = 1 \times 10^{16} \text{ cm}$ to $2.5 \times 10^{16} \text{ cm}$ contributes $\sim 16\%$ to the FIR continuum at $29 \mu\text{m}$, i.e., within the observation uncertainty. The calculated intensities for H₂O vibrationally-excited lines of ν_2 were found

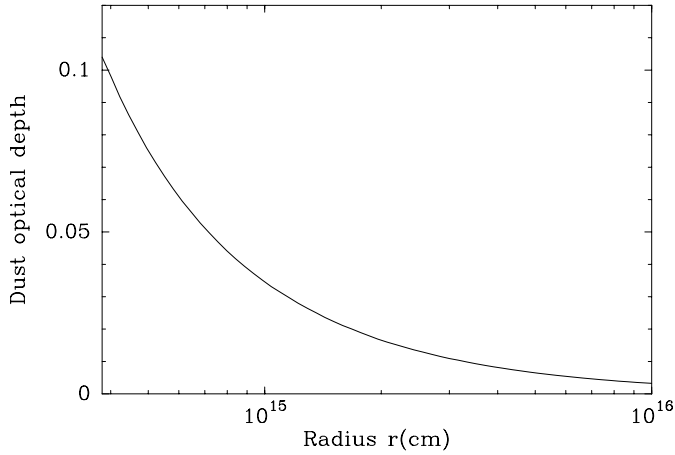


Fig. 6. The averaged optical depth of dust grains at 80 μm from r to ∞

to be weak compared to those in the ground vibrational state, in agreement with the D-NQR model. In order to see the influence of collisional excitation from the ground to the vibrational levels, we multiplied the assumed vibrational collisional rates by a factor of 10. However the calculated H₂O line fluxes in the fundamental state are not appreciably increased. On the other hand, when we decreased the FIR radiation field by a factor of 0.5, we have found that the H₂O line fluxes decrease by e.g., 35% and 46% for the λ 174.62 μm and 99.49 μm lines. If we take out the vibrational levels, the line fluxes would decrease by \sim 61% and 57% for λ 174.62 μm and 45.11 μm , respectively. From this new level configuration, if we vary the collisional rates by a factor of 2 these line fluxes would increase by \sim 6%. These tests indicate that FIR radiative effects are more important than collisions in the excitation of H₂O. This finding opposes that of D-NQR and Barlow et al. (1996) which found that the FIR H₂O lines are mainly excited by collisions. However it is not easy to compare these models with the present because they are fundamentally different and they did not constrain FIR continuum fluxes.

Fig. 6 gives the averaged optical depth of dust grains at 80 μm in the outflow. It decreases from \sim 0.1 to \sim 0.035 as r increases from r_{min} to \sim 1 10^{15} cm.

In Fig. 7, we plot the expansion velocity, $v_e(r)$, of the circumstellar gas, the drift velocity, $v_d(r)$, of dust grains relative to the gas, and the logarithmic velocity gradient, $\epsilon(r) = d \ln v_e / d \ln r$, as functions of radius r . Here we have set $\epsilon = 0.02$ for ϵ lower than this value as discussed by D-NQR for the large velocity gradient approximation, but numerical tests show that the modelling results are insensitive to the adopted limiting value of ϵ .

The H₂O line fluxes were calculated neglecting any possible non-local self-absorption effect (Nguyen-Q-Rieu et al. 1984, Morris et al. 1985). This is a good approximation since the change to the (optically thin/thick) line profile due to that effect is small, as it would enhance the redshifted wing while depressing the blueshifted one (Morris et al. 1985, Truong-Bach et al. 1991). Another characteristic of the self-absorbed line is the appearance of a weak shelf on the blue wing when self-absorption does not reduce the intensity entirely to zero. The width of this

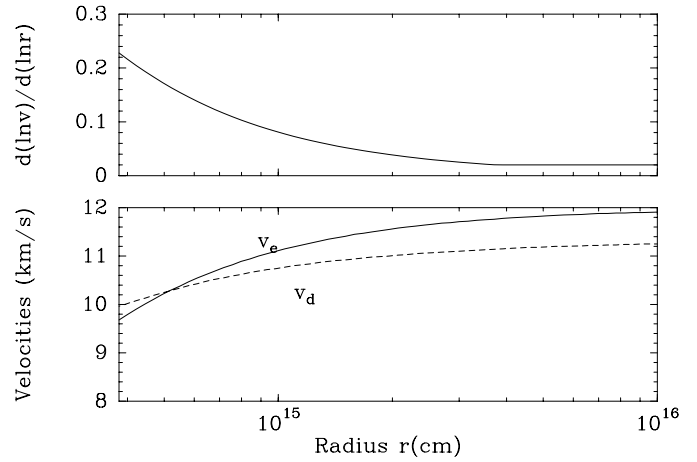


Fig. 7. (Top): the logarithmic velocity derivative $\epsilon(r)$ (in cgs units). (Bottom): the gas expansion velocity v_e and the drift velocity of dust grains v_d as functions of r

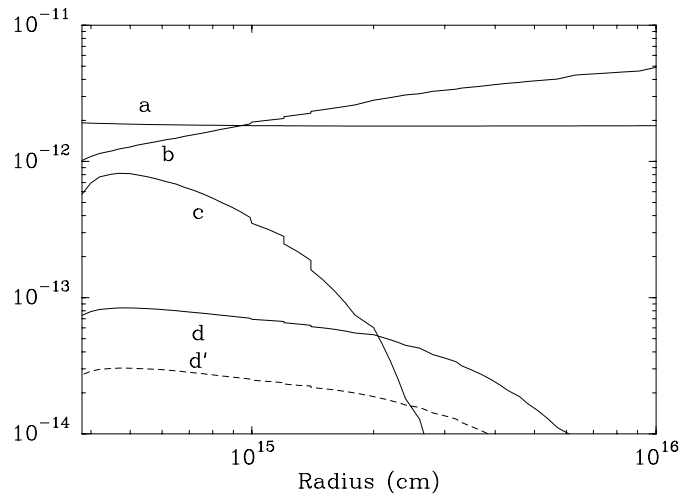


Fig. 8. The heating and cooling rates scaled by $(r/10^{14} \text{ cm})^4$ (in $\text{erg cm}^{-3} \text{ s}^{-1}$). *a*, grain-gas heating. Cooling due to: *b*, adiabatic expansion of gas; *c*, H₂ vibrational excitation; *d*, H₂O rotational excitation. *Solid*, ortho; *dashed*, para H₂O

shelf increases with the line opacity (Morris et al. 1985). The low spectral resolutions of our observations do not allow us to discern the line profiles. It would be of interest to make higher-resolution observations of the IR line profiles. The optical depths at line centre (radial velocity $v_{\text{r}} = 0$, Castor 1970) and along the line of sight at projected distance in the plane of the sky $p \sim 1 \times 10^{15}$ cm are \sim 3.56, 1.40, and 0.15 for the 174.62 μm , 99.49 μm (505–411), and 45.11 μm lines, respectively.

In Fig. 8, we plot the heating and cooling rates. The momentum transfer efficiency used is $Q \sim 0.073$. We recall that Q is linked to the grain characteristic factor $n_d \sigma_d$ in the dust absorption coefficient. A lower (or higher) value of Q means a higher (or lower) value of $n_d \sigma_d$ for the fit. Grain-gas collisions dominate the heating effects. Cooling by gas adiabatic expansion is important throughout the envelope. Cooling by H₂ $v = 1-0$ pho-

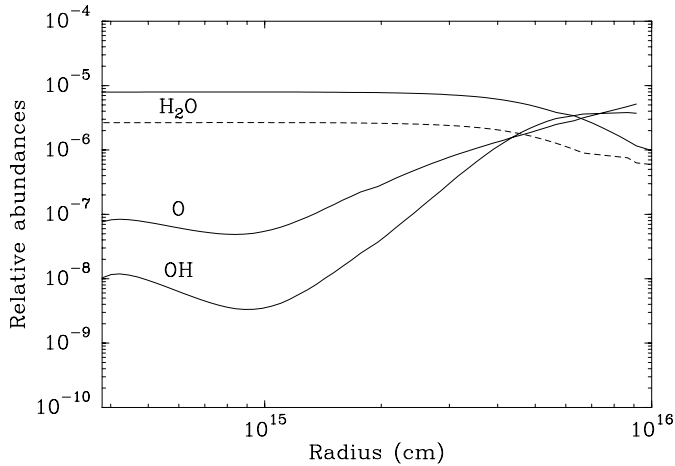


Fig. 9. The radial abundances of H₂O, O, and OH as functions of r for Observations II. *Solid*, ortho; *dotted*, para H₂O

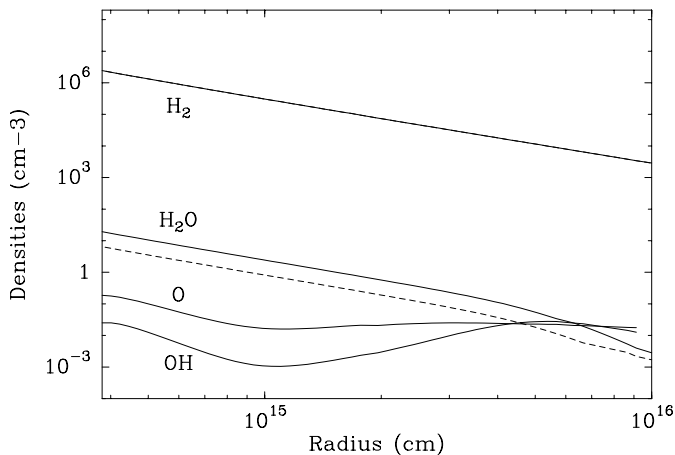


Fig. 10. The radial density distributions of H₂ and H₂O, O, and OH for Observations II. *Solid*, ortho; *dotted*, para H₂O

tons is important close to the central star. CO cooling is weak and operates outside of the H₂O excitation region.

Fig. 9 shows the radial abundances of H₂O in chemical equilibrium with O and OH. The *ortho*-to-*para* abundance ratio which starts from the value ~ 3 in the centre remains unchanged through the entire shell. Fig. 10 represents the densities of H₂, H₂O, OH, and O.

Fig. 11 shows the kinetic temperature distribution, $T_K(r)$, in the envelope. It decreases outward from ~ 1300 K. At radius $r \sim 10^{15}$ cm, the kinetic temperature is still ~ 950 K. At $r \sim 5 \times 10^{15}$ cm, it becomes ~ 380 K. From here the excitation becomes too low to contribute significantly to the observed line fluxes.

In Fig. 12 we plot the radial abundances of H₂O (*ortho*), O, and OH for two cases: with and without photodissociation. The weak dust shielding would allow interstellar FUV to penetrate deeply into the inner shell, since photoproduction of O and OH begins at $r \sim 7 \times 10^{14}$ cm. However their abundance variations are weak compared to the H₂O abundance itself. The figure shows that the H₂O abundance is perceptibly decreased

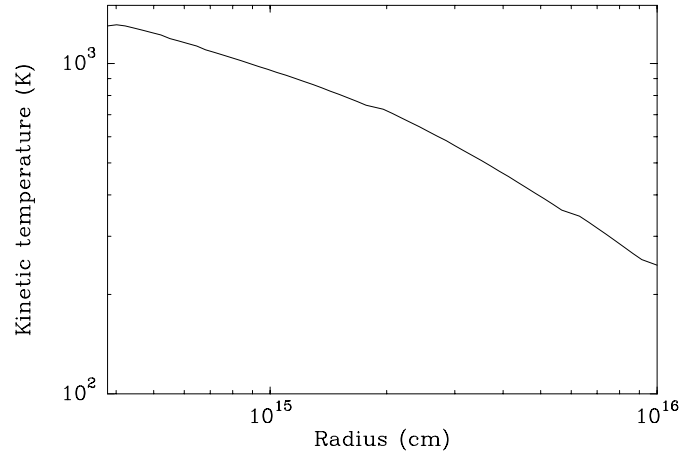


Fig. 11. The radial kinetic temperature distribution $T_K(r)$

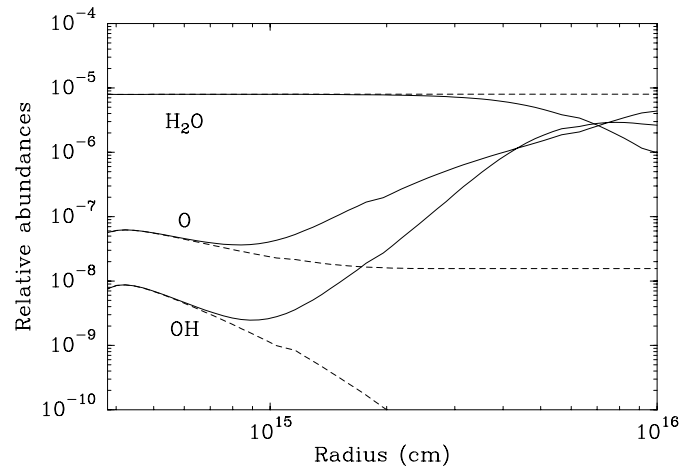


Fig. 12. The radial abundances of H₂O (*ortho*), O, and OH. *Solid*, with UV photodissociation; *dotted*, without photodissociation

by photodissociation from $r \gtrsim 2.2 \times 10^{15}$ cm. Here H₂O excitation becomes low and the outer layers contribute typically ~ 14 – 20% to LW (λ 174.62 μm and 108.07 μm), but insignificantly to SWS (λ 45.11 μm and 29.87 μm) line fluxes.

Uncertainty of the derived mass loss and H₂O abundance due to uncertain thermodynamics can be estimated. We recall that the kinetic temperature depends on the drift velocity which varies as $(Qv_e)^{0.5}$. If we varied this quantity, for example by decreasing Q by $\delta Q = 50\% Q$, the calculated line fluxes would noticeably decrease by, e.g., 80%, 92%, and 98% for the LW 174.62 μm , SW 82.03 μm , and SWS 34.549 μm lines, respectively. For a moderate decrease of Q , $\delta Q = 30\% Q$, the latter line intensities would decrease only by 30%, 31%, and 50%, respectively, i.e., by about their observed uncertainties. We tried to compensate this line flux diminution by increasing the mass-loss rate by $\delta \dot{M} = 25\% \dot{M}$, and the intensity deviations were reduced to 0.6%, 8%, and 19%, respectively. This means that the new fit obtained is still good for LW but worse for shorter wavelengths. Therefore the upper limit of uncertainty for the mass loss can be estimated as 25%. Accordingly, the uncertainties for the H₂O abundance and the kinetic temperature at the inner

radius were derived to be 47% and 33%, respectively. Naturally the final uncertainty should include that from the Hipparcos distance determination (van Leeuwen et al. 1997).

5. Conclusion

Within the uncertainties encountered in the observations as discussed, Figs. 1, 2 & 4 and Tables 1–3 show acceptable comparisons between the predicted and observed integrated H₂O spectra and FIR continua. The error bars for observed SWS line fluxes are nevertheless high by a factor 2–3; and it is interesting to perform new observations in this wavelength region. Some CO and CII lines are also found in the H₂O spectra but their investigation is outside the scope of this paper. Observations I show a spectrum about ~ 30 – 40% less intense compared to Observations II's where new or stronger lines appear and some components of a blend become clearly visible. This light variation due to stellar pulsations of the Mira variable between ~ 300 days corresponds to a phase variation from 0.5, at minimum luminosity, to 0.2 where the luminosity was falling down to its mean value. A best fit to the overall spectra were obtained with the dilution factor for the FIR field scaled by a factor ~ 0.61 and 1, respectively. This FIR field is mainly dominated by dust. Photodissociation of H₂O by the interstellar UV field does not affect significantly the FIR H₂O line fluxes, since the effect operates in the outer region ($\gtrsim 10^{16}$ cm) where H₂O excitation temperatures become weak. The derived H₂O abundance and density, the FIR continuum, and the kinetic temperature vary as functions of radius across the circumstellar envelope of R Cas. The derived mass-loss rate is $\dot{M} \sim 3.4 \times 10^{-7} M_{\odot} \text{ yr}^{-1}$. The deduced total (ortho and para) H₂O abundance is $\sim 1.1 \times 10^{-5}$ and the kinetic temperature is $T_k \sim 1300$ K at the inner radius of the envelope, and they decrease with distance from the star. Compared to W Hya, another O-rich AGB star, the initial H₂O abundance of R Cas is about one order of magnitude lower while the mass-loss rate is nearly the same; the values found for W Hya (1.2×10^{-4} and $6 \times 10^{-7} M_{\odot} \text{ yr}^{-1}$; see Barlow et al. 1996) were determined, however, by using a different model. Similar mass-loss rates ($10^{-7} M_{\odot} \text{ yr}^{-1}$) have been derived for a number of LPV Miras using CO radio observations (e.g. Knapp et al. 1982, Loup et al. 1993, Kahane & Jura 1994, Gonzalez-Alfonso et al. 1998), and optical K_I/Na_I scattering (e.g. Guilain & Maunon 1996).

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