Gas phase atomic metals in the circumstellar envelope of IRC+10216

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ABSTRACT

We report the results of a search for gas phase atomic metals in the circumstellar envelope of the asymptotic giant branch carbon star IRC+10216. The search was made using high resolution (λ/Δλ = 50,000) optical absorption spectroscopy of a background star that probes the envelope on a line of sight 35″ from the center. The metal species that we detect in the envelope include Na i, K i, Ca i, Ca ii, Cr i, and Fe i, with upper limits for Al i, Mn i, Ti i, Ti ii, and Sr ii. The observations are used to determine the metal abundances in the gas phase and the condensation onto grains. The metal depletions in the envelope range from a factor of 5 for Na to 300 for Ca, with some similarity to the depletion pattern in interstellar clouds. Our results directly constrain the condensation efficiency of metals in a carbon-rich circumstellar envelope and the mix of solid and gas phase metals returned by the star to the interstellar medium. The abundances of the uncondensed metal atoms that we observe are typically larger than the abundances of the metal-bearing molecules detected in the envelope. The metal atoms are therefore the major metal species in the gas phase and likely play a key role in the metal chemistry.

Key words. Stars: AGB and post-AGB – stars: carbon – stars: mass-loss – stars: individual: IRC+10216 – circumstellar matter

1. Introduction

Metals are important constituents of the interstellar medium. Abundant metals such as calcium and iron are among the most heavily depleted elements in dense clouds, and therefore form a significant component of interstellar dust. The process of dust formation is not well understood (e.g., Draine 2009), but much of the raw material is supplied by mass loss from evolved stars, especially the asymptotic giant branch (AGB) stars.

In the winds of oxygen-rich AGB stars, the dust is primarily in the form of metal silicates, so that the metals play an important role in the formation of dust and the mass loss process. In carbon-rich AGB stars, the situation is less clear. The dust is believed to consist mainly of graphite or amorphous carbon, and silicon carbide. The extent to which metals contribute to dust formation, and the form in which they are returned to the interstellar medium is essentially unknown.

In this paper, we report the first comprehensive search for gas phase atomic metals in a carbon-rich circumstellar envelope. IRC+10216 (CW Leo) is the nearest carbon star with a thick circumstellar envelope, and serves as an archetype for the study of mass loss on the AGB. The star itself is too faint and its spectrum too complex to serve as a useful source for detailed study, although circumstellar C₂ and CN have been observed in this way (Bakker et al. 1997). There are, however, other stars in the field. IRC+10216 is nearby, at a distance of ∼ 120 pc (e.g., Ramstedt et al. 2008), so the circumstellar envelope extends a considerable angle on the sky. It is detected out to a distance of ∼ 3′ from the center in millimeter CO emission (Huggins et al. 1988), and 9″ in infrared dust emission observed with IRAS (Young et al. 1993). Although IRC+10216 is at a relatively high galactic latitude (l = 221°, b = +47°), there are several stars in this region of the sky that are candidates for background sources, as seen in the wide field image in Fig. 1 of Mauron & Huggins (1999).

One of these stars, Star 6 in the UBV photometric sequence of the field by Mauron et al. (2003), is well suited for absorption line studies. It is located behind the envelope at an angular offset of 35″ from the center, and is bright enough for high resolution spectroscopy. This star has been observed with the UVES spectrograph at the VLT by Kendall et al. (2002). Their main objective was to search for diffuse bands that might originate in the circumstellar
gas. No diffuse bands were found, but these authors noted deep absorption lines of Na i and K i, which they attributed to circumstellar gas.

Here we report a comprehensive search for metal lines in the circumstellar envelope of IRC+10216 along this line of sight. Our objectives are to measure the degree of metal depletion in this carbon rich-environment, and to determine the distribution of solid and gas phase metals returned by the star to the interstellar medium. Sect. 2 describes the observational material. Sect. 3 presents the absorption lines, and Sect. 4 the derived column densities and abundances. The results are discussed in terms of dust formation in Sect. 5, and metal chemistry in Sect. 6. Our main conclusions are given in Sect. 7.

2. Observations

The observations of the envelope of IRC+10216 were made using absorption line spectroscopy with Star 6 (USNO 0975-0633-6975) as the background source of illumination. Star 6 lies 35″ from IRC+10216 at position angle 165°. Its visual magnitude is \( V = 16.0 \), and it is the nearest star to the center of the envelope suitable for high resolution spectroscopy. The field is shown in Fig. 1, where the envelope is seen in dust-scattered Galactic light. From its magnitude and spectral type (type G), star 6 is at a distance of \( \sim 1400 \) pc, well beyond IRC+10216, as discussed by Kendall et al. (2002).

The observations were made with the UVES spectrograph at the VLT, and were previously used by Kendall et al. (2002) to search for diffuse bands. The data were obtained over seven nights in December 2000 and January 2001, with an effective exposure of 4 hr at each wavelength. The sky transparency was moderate to good, and the seeing was 0″8. The data consist of 4 spectra, each covering the complete range from 3000 to 10 000 \( \text{Å} \) with a resolving power of 50 000 (\( \Delta \nu = 6 \text{ km s}^{-1} \)). These spectra have been reduced with the ESO UVES pipeline and have been summed after correction to the heliocentric reference frame. All radial velocities given in this paper are heliocentric, except where specified otherwise. The signal-to-noise ratio of the final spectrum at 4200 \( \text{Å} \) is \( \sim 60 \) per resolution element.

The heliocentric velocity of Star 6 measured from numerous photospheric absorption lines is \( +52.4 \pm 0.6 \text{ km s}^{-1} \). For comparison, the systemic heliocentric radial velocity of IRC+10216 is \( -19.3 \text{ km s}^{-1} \), which is derived from millimeter observations of the envelope (\( \text{v}_{\text{LSR}} = -26 \text{ km s}^{-1} \), Loup et al. 1993). Thus a given line in the envelope of IRC+10216 is well separated in velocity from the line formed in the photosphere of Star 6. In addition, the expansion velocity of the envelope is 14.1 \( \text{ km s}^{-1} \) with a small turbulent component (Huggins & Healy 1986), so the absorption lines in the envelope are expected to be wide (\( \sim 30 \text{ km s}^{-1} \)). In contrast, the weak stellar lines are relatively narrow, with widths comparable to the instrumental profile, and the strong stellar lines have characteristic damping profiles with broad wings.

Table 1 lists the metal lines that we searched for in the envelope of IRC+10216. Column 2 gives the laboratory wavelength in air (\( \lambda_{\odot} \)) for each transition, and column 3 gives the oscillator strength (\( f \)) from Morton (1991, 2000). The line list includes the strongest ground-state transitions for the species that are potentially observable in the wavelength range covered. These lines are seen in interstellar clouds and/or in circumstellar environments with similar physical conditions.

At long wavelengths, the spectrum of the background source, Star 6, is relatively free of photospheric lines, so an absorption line arising in the intervening circumstellar envelope is straightforward to identify. At shorter wavelengths, the spectrum is more crowded with photospheric lines, and absorption by the circumstellar envelope is often more difficult to identify. In these cases we use a template technique to extract the envelope signal. We first make a least squares fit to the spectrum of Star 6 using a template spectrum covering a region of \( \sim 15 \text{ Å} \) around (but excluding) the expected envelope line. We then use this as the effective continuum to search for residual absorption from the envelope.

One form of template that we tried was based on the library of synthetic spectra from Coelho et al. (2005), but these did not produce a good match to the photospheric spectrum. A second approach that was successful, was using a scaled solar spectrum. By chance the spectrum of Star 6 is very similar to that of the Sun, and a template based on scaling the solar spectrum gives a close match to the stellar spectrum. The solar spectrum that we use here was published by Delbouille et al. (1973) and covers the wavelength region 3000–10 000 \( \text{Å} \) with a step size of about 0.0125 \( \text{Å} \). In order to fit the stellar spectrum \( F_{\odot}(\lambda) \), we shift the solar spectrum \( F_{\odot}(\lambda) \) to match the radial velocity, rebin it to match the resolution, and scale the flux according to:

\[
F_{\star}(\lambda) = \alpha F_{\odot}(\lambda) + \beta,
\]

where \( \alpha \) and \( \beta \) are constants that are determined from the least-squares fit to the local region of spectrum under consideration.

The use of this template as the effective continuum reveals spectral features of the circumstellar envelope that are...
not otherwise easily observed. Examples of the application of this technique are given in Sects. 3.3 and 3.4.

3. Envelope absorption lines

3.1. K I

Fig. 2 shows the observed spectral regions covering the K I doublet. The 7665 Å line is in the upper panel and the 7699 Å line is in the lower panel. Strong photospheric K I lines of Star 6 are present at the stellar radial velocity of \(V_r = +52.4 \text{ km s}^{-1}\), which is marked with an arrow at the right of each panel. Also present are two weaker photospheric lines (marked +) and four telluric lines (marked o).

The dominant feature in the middle of each spectral region is strong K I absorption from the envelope of IRC+10216. This can be unambiguously ascribed to the circumstellar envelope. It is centered near the systemic radial velocity of IRC+10216 at \(-19.3 \text{ km s}^{-1}\) (marked with an arrow in the figure) and has the broad width expected from the expanding envelope. With the low interstellar absorption in this region of the sky (\(E_{B-V} \lesssim 0.03\)), any interstellar K I would contribute \(\lesssim 1\) percent of the strong lines observed, based on the survey of interstellar K I by Chaffee & White (1982).

We also searched for the much weaker ground state K I line at 4044 Å which has an \(f\)-value \(\sim 110\) times less than the 7664 Å line. The 4044 Å line was not detected, and the upper limit is given in Table 1.

Although the K I doublet lines formed in the envelope are broad, they are clearly composed of several components. The presence of the components is important because it affects the saturation of the lines (see Sect. 4). Assuming gaussian broadening, we found that a best fit synthesis taking into account the instrumental profile requires four components to match the line shapes. A preliminary fit with unconstrained parameters gave consistent estimates for the radial velocities of the individual components in each line. We then fixed the velocities at their mean values, and fit the profiles by varying the relative column densities of the components and their line broadening b-values, which we constrained to be the same for both lines. The final mean parameters are given in Table 2, where the strength is the column density of each component, relative to the strongest component. The uncertainty in the velocities is \(\lesssim 0.5 \text{ km s}^{-1}\) and the relative strengths found from the two lines are the same to within \(\lesssim 5\%\). Thus the results from the two lines are in good agreement, as expected from their similar line profiles.

| Species | \(\lambda_c\) (Å) | \(f\)-value | \(W^e_N\) (mÅ) | \(N^e\) (cm\(^{-2}\)) |
|---------|-----------------|-------------|----------------|-----------------|
| Na I    | 5895.924        | 0.3180      | 625            | 4.6(14)\(^b\)  |
|         | 5889.951        | 0.6311      | 625            |                 |
|         | 3392.368        | 0.0090      | 230            |                 |
| Al I    | 3944.006        | 0.1134      | <44            | <2.8(12)        |
| K I     | 7698.974        | 0.3393      | 530            | 4.6(12)\(^c\)  |
|         | 7664.911        | 0.6816      | 635            |                 |
|         | 4044.143        | 0.0061      | <29            |                 |
| Ca I    | 4226.728        | 1.7530      | 300            | 1.9(12)         |
| Ca II   | 3968.468        | 0.3145      | 255            | 7.0(12)         |
|         | 3933.663        | 0.6346      | 390            |                 |
| Ti I    | 3635.462        | 0.2229      | <49            | <1.9(12)        |
| Ti II   | 3383.708        | 0.3401      | <69            | <2.0(12)        |
| Cr I    | 4289.716        | 0.0622      | 37             | 1.4(12)\(^d\)  |
|         | 4274.796        | 0.0839      | 49             |                 |
|         | 4254.332        | 0.1099      | 38             |                 |
|         | 3605.322        | 0.2248      | 27             |                 |
|         | 3593.482        | 0.2897      | 62             |                 |
|         | 3578.683        | 0.3663      | <62            |                 |
| Mn I    | 4034.483        | 0.0257      | <33            | <3.1(12)        |
|         | 4033.062        | 0.0402      | <33            |                 |
|         | 4030.753        | 0.0565      | <33            |                 |
| Fe I    | 3859.911        | 0.0217      | 225            | 8.8(13)\(^e\)  |
|         | 3824.444        | 0.0048      | 80             |                 |
|         | 3719.934        | 0.0412      | 197            |                 |
|         | 3440.606        | 0.0236      | 170            |                 |
| Sr II   | 4077.709        | 0.7010      | <28            | <2.7(11)        |

\(^a\) Upper limits are 3\(\sigma\).
\(^b\) D-lines are saturated, \(N\) based on the 3302 Å line.
\(^c\) Based on the 7699, 7665 Å doublet.
\(^d\) : indicates 2–3\(\sigma\) features; \(N\) based on weighted mean.
\(^e\) Based on the 3860, 3720 Å lines which have better S/N.
Fig. 3. Comparison of the observed (black) and synthesized (grey) profiles of the K\textsc{i} 7698.97 Å line, using the parameters given in Table 2. The individual components are also shown.

To illustrate the quality of the multi-component fit, Fig. 3 shows the results of a synthesis of the 7699 Å line using the parameters of Table 2. It can be seen that the model provides an excellent fit to the observational data.

We interpret the multiple component profiles of the K\textsc{i} lines as the effects of the multiple shell structure in the envelope (Mauron & Huggins 1999, 2000; see also Fig. 1), where the line of sight passes through regions of enhanced density. Although the gaussian line shapes give a good fit to the spectra, we do not know the detailed velocity distribution of the shells along the line of sight, so we cannot reliably determine the gas density contrast between the shell and inter-shell regions. However, the stronger components are clearly separate, and suggest that the contrast is at least a factor of a few. This is consistent with a large shell inter-shell contrast in the dust density derived from images in dust-scattered light by Mauron & Huggins (2000).

Table 2. Parameters of the K\textsc{i} components in IRC+10216

| Component | Strength (rel.) | $b$ (km s$^{-1}$) | $V_r$ (km s$^{-1}$) |
|-----------|----------------|-----------------|-------------------|
| 1         | 0.27           | 2.8             | −32.4             |
| 2         | 0.32           | 2.9             | −28.0             |
| 3         | 1.00           | 5.5             | −21.0             |
| 4         | 0.51           | 3.9             | −09.2             |

3.2. Na\textsc{i}

Fig. 4 shows the spectral regions covering the Na\textsc{i} D lines. The 5890 Å line is in the upper panel and the 5896 Å line is in the lower panel. The photospheric Na\textsc{i} lines of Star 6 (marked with arrows at the right in each panel) are strong, and there is another weaker photospheric line in the lower panel. The dominant feature in the middle of each spectrum is Na\textsc{i} absorption from the envelope of IRC+10216, centered near the systemic velocity. It can be seen that the circumstellar lines are extremely strong. All the components are highly saturated and the residual intensities are close to zero across both lines.

There is an additional, weak Na\textsc{i} component that appears in both spectra near +9.5 km s$^{-1}$ (marked with triangles) which we identify as an interstellar component along this line of sight. Similar weak Na\textsc{i} absorption near this velocity is reported by Kendall et al. (2002) along lines of sight to two other stars in this region of the sky, at angular distances of 153$''$ and 2.5$'$ from IRC+10216.

We also searched for the much weaker ground state Na\textsc{i} line at 3302 Å, which has an $f$-value ∼ 70 times less than the 5890 Å D line. The 3302 Å line lies in a crowded, low signal-to-noise part of the spectrum, but the template approach (Sect. 2) reveals a ∼5 sigma detection, and the equivalent width is given in Table 1. The signal-to-noise ratio is too low to show any details of the profile.

3.3. Ca\textsc{i} and Ca\textsc{ii}

Fig. 5 shows the spectral region around the Ca\textsc{i} line at 4226 Å. The line falls in a very crowded region of the spectrum and provides a good illustration of the template method.
The solid line in the upper panel shows the observed spectrum, and the envelope absorption is not immediately apparent. The dotted line in the upper panel shows the stellar template, scaled to match the observed spectrum. It can be seen that it gives a good overall fit to the data, and reveals the excess absorption from the envelope. The lower panel of Fig. 5 shows the envelope line, using the fitted template as the effective continuum. The small wavelength range around the core of the photospheric Ca\textsc{i} line has been masked (with the horizontal dotted line) because the effective continuum is poorly defined at the low intensity of the line core. The main deviation from a flat continuum in the normalized spectrum is caused by slight differences between Star 6 and the Sun.

3.4. Fe\textsc{i}

Figs. 7 and 8 show the spectral regions around the Fe\textsc{i} lines at 3720 Å and 3860 Å, respectively. These fall in relatively crowded regions of the photospheric spectrum, and the template is needed to determine the envelope absorption. The upper panel in each figure shows the observed spectrum (solid line) and the fitted stellar template (dotted line). The lower panel shows the normalized spectrum with the envelope absorption centered near the systemic velocity of IRC+10216. The signal-to-noise ratio of these spectra is lower than those discussed above, and the template fit is affected by differences in the solar and stellar spectra. The limited quality of the fit is probably responsible for the fact that the equivalent width of the 3720 Å line is slightly less than that of the 3860 Å line, even though the f-value is larger (see Table 1). Nevertheless the lines are well
detected, and the multi-component character of the envelope absorption is similar to that seen in the lines of the other species.

We also searched for two additional Fe I lines at 3824 Å and 3441 Å. The 3824 Å line is weaker on account of a significantly lower $f$-value, and is marginally detected at $\sim 4$ sigma. The 3441 Å line lies at shorter wavelengths where the spectrum is poorer. It is detected with a strength comparable to the Fe I lines shown in Figs. 7 and 8, but with a lower signal-to-noise ratio.

3.5. Other lines

Searches were also made for lines of several other species, including Al I, Mn I, Cr I, Ti I and Ti II, and Sr II, as listed in Table 1. With the exception of Cr I, no significant envelope absorption was detected. For Cr I, weak ($\sim 2$–$3\sigma$) features are seen at the wavelengths of most of the accessible lines. When appropriately weighted by the oscillator strengths and noise levels they provide an overall 4-sigma detection of this species.

4. Abundances in the envelope

4.1. Column densities

The equivalent widths ($W_\lambda$) of the absorption lines observed in the envelope are listed in column 4 of Table 1. For the non-detected lines, we list 3-sigma limits given by:

$$W_\lambda < 3\sigma \Delta \lambda / \sqrt{n},$$

where $\sigma$ is the noise level and $n$ is the number of resolution elements in the line width $\Delta \lambda$.

The column density ($N$) derived for each species is given in column 5 of Table 1. It is based on the equivalent widths and the $f$-values given in the table. For the weak lines, the column density is directly related to the equivalent width by the optically thin formula:

$$N = 1.13 \times 10^{20} W_\lambda / f \lambda^2,$$

where $W_\lambda$ and $\lambda$ are in Å units.

For the stronger lines, where the optical depths are larger, we determine the relation between $W_\lambda$ and $N$ using the multi-component model used to fit the high signal-to-noise profiles of the K I lines (Table 2). With this model we find that the K I lines are moderately saturated; the column densities derived from the 7699 and 7665 Å lines are larger than the optically thin values by factors of 1.7 and 2.3, respectively. The column densities from the two lines agree to within 20%, and are consistent with a limit of $N$(K I) $\lesssim 3 \times 10^{13}$ cm$^{-2}$ determined from the upper limit to the optically thin 4044 Å K I line. For Na I, the D-lines are extremely optically thick, and therefore insensitive to the column density, although the very low residual intensity across the lines yields a lower limit of $N$(Na I) $\gtrsim 1$–$2 \times 10^{13}$ cm$^{-2}$. Fortunately, we also detect the optically thinner Na I line at 3302 Å line, and we use this with the multi-component saturation curve to determine the column density of Na I given in Table 1. The result is consistent with the lower limit from the D-lines.

The profiles of the ionized lines might be expected to differ from those of the neutral lines because the absorption
can occur at different locations along the line of sight. This does not affect the column density estimates for Sr II and Ti II because the lines are very weak (below the detection level) and are therefore optically thin. For the stronger Ca II lines, the profiles are found to be similar to the K I profiles, but as noted in Sect. 3.3, they are slightly broader with an additional contribution to the absorption. We therefore model the Ca II lines as the sum of two contributions; the fit of the K I multi-component model to the main part of the profile, and an additional, optically thin component to give the total equivalent width. The additional component contributes 32% and 17% of the equivalent widths (22% and 13% of the derived column densities) for the 3933 and 3968 Å lines, respectively. We assume that the additional absorption is circumstellar, but since it is relatively small, the results do not depend sensitively on this assumption.

4.2. Column density of hydrogen

Although there are no direct observations of hydrogen along the line of sight, we can use estimates of the mass loss rate of the envelope to determine the hydrogen column density. Since the early work by Kwan & Hill (1977) there have been numerous estimates of the mass loss rate based on millimeter CO observations, and the results are fairly consistent when the different distances, CO abundances, dustgas heating rates, and He content are taken into account. For a distance of 120 pc, we adopt a mass loss rate \( M_H \) (in hydrogen) of \( 1.25 \times 10^{-5} \, M_\odot \, yr^{-1} \) (corresponding to a total mass loss rate of \( 1.75 \times 10^{-5} \, M_\odot \, yr^{-1} \)), based on the analysis of Schöier & Olofsson (2001) approximately corrected for the effects of He. This value is consistent with other recent estimates.

For \( M_H = 1.25 \times 10^{-5} \, M_\odot \, yr^{-1} \), the column density of hydrogen \( N(H) \), where \( N(H) = N(H_1) + 2N(H_2) \), is \( 1.3 \times 10^{21} \, \text{cm}^{-2} \) along the line of sight 35″ from the center. The uncertainty is a factor \( \sim 2 \), which results from uncertainties in \( \dot{M} \). In addition to a smooth decrease of the column density with distance from the center, there are other variations caused by the multiple shell structure in the envelope. From an analysis of scattered light images (e.g., Fig. 1) we find typical variations in the dust column density of \( \pm 20\% \), and somewhat smaller variations near the region of Star 6. These variations are much smaller than the density contrast of the shells because the column density averages the density along the line of sight through several shell and inter shell regions. The envelope structure is therefore not a major source of uncertainty for \( N(H) \).

4.3. Ionization fraction

The line of sight passes through regions of the circumstellar envelope where the metals are partially ionized. Since most of the observed lines arise from single stages of ionization, we need to consider ionization corrections in order to determine the total gas phase abundances. We estimate the corrections from the relative column densities of the ionization stages of each metal, obtained using the photo-ionization model of an expanding envelope discussed by Glassgold & Huggins (1986).

In the model, neutral atoms (which may result from the dissociation of molecules) emerge from the dense, shielded, inner envelope and are photo-ionized by the ambient interstellar radiation field. We calculate the ionization fraction as a function of radius, and use this to calculate the ionization fraction along the line of sight. The ionization of each element is governed by equation 4.9 of Glassgold & Huggins (1986), with no contribution from chromospheric radiation. The ionization depends on the photo-ionization rate (given by the interstellar rate and the envelope shielding) and recombination. In solving the ionization equations we use the interstellar photo-ionization rates and recombination rates from Péquignot & Akdörgazi (1986), except for Cr and Sr which are not included in their compilation; for these the photo-ionization rates are from Glassgold & Huggins (1986) and the recombination rates from Bernat (1976). The radial dependence of the electron abundance is adopted from Cordier et al. (2007), although recombination is relatively unimportant except for Al. For the shielding of interstellar radiation in the envelope we use the standard carbon dust model and dust-to-gas ratio of Cherchneff et al. (1993) and the shielding function of Morris & Jura (1983).

There are a number of uncertainties in the parameters of the model, the most important being the strength of the ambient radiation field (which determines the interstellar photo-ionization rates) and the shielding in the envelope (which includes uncertainties in the dust parameters, the mass loss rate, and the envelope geometry). Fortunately we observe both the neutral and ionized column densities of Ca, and we use this to set the ionization level in the model. For the nominal parameters given above, the predicted ratio \( N(Ca ii)/N(Ca i) \) is a factor 3.6 larger than observed. This is fair agreement considering that the ionization in the envelope has not previously been constrained in this way. However, we can do much better, by fine tuning the ionization level in the envelope to fit the Ca ionization exactly. This can be done by adjusting the ambient radiation field (by a factor of 0.46) or the shielding optical depth (by a factor 1.5). Both variations are within their respective uncertainties. For specificity, we adopt the reduced radiation field to calculate the ionization of the other elements. The resulting ionization corrections \( C_i = N(i + ii)/N(i) \) for neutral species, and \( N(i + ii)/N(ii) \) for singly ionized species), are given in column 4 of Table 3. Adjusting the shielding instead, by the amount given above, produces essentially the same ionization corrections.

The ionization correction for Al in Table 3 is much larger than for the other metals because of its relatively high photo-ionization rate. Our observations are therefore not very sensitive probes of the column density of Al in the gas phase because the Al is nearly completely ionized along the line of sight. The ionization corrections for the other neutral species are much smaller. Some Na is predominantly neutral. This is in contrast to typical interstellar clouds with a similar column density of hydrogen. In the circumstellar envelope, the characteristic outflow time at 35″ is shorter than the photo-ionization time given by Glassgold & Huggins (1986). Hence, even without dust shielding, the Na atoms are expected to be largely neutral, as given in the table.

4.4. Abundances

Except for Ca and Ti, the total gas phase column densities of the metals are determined from the observed column densities and the ionization corrections. The results are given in column 5 of Table 3. Although the \( N(Ca ii)/N(Ca i) \) ratio...
was used to determine the ionization level in the envelope, we use the sum of the ionization stages to obtain the total Ca column density, independent of the ionization. Similarly, the limit for Ti is based on the observed limits for Ti i and Ti ii, and so is independent of the ionization.

The gas phase abundances of the metals relative to hydrogen ($X$) are determined from the total column densities and the value of $N(H)$ from Sect. 4.2, and are given in column 6 of Table 3. For reference, the solar values ($X_\odot$) are given in column 7 of the table, taken from Lodders (2003).

Comparison of the envelope and solar abundances shows that there are large deficiencies in the gas phase abundances in the envelope, which vary from metal to metal.

5. Dust condensation

Several processes affect the state of the circumstellar material as it moves from the stellar photosphere out through the envelope into the interstellar medium. In order of increasing distance from the star these processes include: dust condensation in conditions of approximate thermodynamic equilibrium; gas phase chemical reactions; photodissociation of the molecules; and eventual photo-ionization of the atomic constituents in the outer envelope (e.g., Gilman 1969; Tsuji 1973; McCabe et al. 1979; Huggins & Glassgold 1982; Lafont et al. 1982). In the carbon-rich environment of IRC+10216, the main component of the dust is amorphous carbon with a minor component of SiC (e.g., Martin & Rogers 1987). The state of other elements, especially the refractory metals, is not well understood (e.g., Turner 1995). Hence our observations of the gas phase metals in the outer envelope provide new constraints on the gas phase chemistry and the dust condensation.

5.1. Observed depletions

Comparison of the envelope abundances with the solar abundances in Table 3 shows that most of the gas phase metal atoms in the envelope are “missing”. It is most unlikely that they are in the form of gas phase molecules. The fractional abundance of even the most extreme case of Ca, where $X(Ca)/X_\odot \sim 3 \times 10^{-3}$, is an order of magnitude larger than the largest fractional abundance of any metal bearing molecule detected outside of the core region (see Table 4). In addition, most molecules are dissociated closer to the star than the 35” offset of our line of sight. It is therefore reasonable to infer that the atomic abundances observed are good approximations to the total gas phase metal abundances in the envelope, and that the missing atoms are depleted onto dust grains.

In column (8) of Table 3 we give the conventional measure of depletion $\log_{10}(\delta)$, where $\delta = X/X_\odot$ is the depletion factor. The observational limits for the abundances of Al and Sr are $\gtrsim$ the solar values, so in these cases we have no significant limits for $\delta$, although Sr is an s-process element and may be enhanced in IRC+10216 and depleted.

Based on the measured depletions, there are two immediate conclusions. First, the metals in this carbon-rich archetype are primarily in the form of solids, and this is the dominant form returned to the interstellar medium. Second, in spite of the depletion, a significant residue of metallic atoms remains in the gas phase and varies from metal to metal.

5.2. Condensation and adsorption

There are no detailed predictions for the depletion of metals in IRC+10216, but there are some important considerations that bear on the issue. A commonly used approach to the condensation of solids in circumstellar envelopes is the assumption of thermodynamic equilibrium in the warm, dense, inner envelope, where the chemical time scales are rapid compared with the expansion time scale. Under these conditions the formation of solid particles is controlled by the condensation temperature of the primary condensate of each species.

For a carbon-rich envelope, the condensation sequence depends somewhat on the C/O ratio and the gas pressure. The following sequence, for C/O = 1.1 and a pressure of $10^{-6}$ bar (from Lodders & Fegley 1995, updated for Fe by Lodders & Fegley 1999) is representative: C (1670 K), TiC (1640 K), SiC (1460 K), FeSi (1230 K), AlN (1170 K), CaS (1150 K), MgS (960 K), with other more volatile metals such as Na and K at lower temperatures. Cr and Mn probably form sulphides but their location in the sequence is uncertain. Thus the metals Ti, Fe, Al, Ca, and then K and Na, are expected to be removed from the gas phase successively. Those with lower condensation temperatures are less likely to go to completion because of the decreasing density with radius.

This qualitative picture is largely consistent with the observed depletion pattern. The observations provide a firm upper limit on the gas phase abundance of Ti (from the Ti i and Ti ii lines); Fe and Ca are strongly depleted, although Ca is more depleted than Fe (in reverse order to the condensation sequence); and Na and K are less depleted.

The simplifying assumption of thermodynamic equilibrium is not a complete physical theory of condensation because kinetic effects must play a role near freeze-out. In addition, once formed, the grains can act as sites for the adsorption of gas phase species further out in the envelope (Jura & Morris 1985). The adsorption depends on the binding energy of the species to the grain surface, and on the sticking probability ($p$), which is essentially unknown. Using the analysis of Jura & Morris (1985), Turner (1995) finds that for IRC+10216, a volatile species such as K is depleted by adsorption to a fractional abundance of 0.84 (for $p = 0.1$) and 0.17 (for $p = 1$), and a more refractory species such as Al is depleted to a fractional abundance of 0.05 (for $p = 0.1$) and $10^{-10}$ (for $p = 1$). Thus adsorption of metals may be as important as the initial condensation.

Our observations of a residual atomic component in the gas phase constrain the efficiency of both condensation and adsorption. Below about 1150 K, the phase diagrams of Lodders & Fegley (1995) show that Ca is essentially completely removed from the gas phase, but our observations show that even for this refractory metal there is a residual component in the gas phase. Metal bearing molecules with Na and K (albeit with low fractional abundances) are seen in the core region (see Sect. 6) but not in the extended envelope. This may be the result of adsorption. On the other hand, adsorption of the more refractory species to levels of $10^{-10}$ are clearly ruled out by the observations.

We conclude that current theoretical ideas are qualitatively in accord with our observations of metal depletion, but improved models with specific quantitative predictions are needed to discriminate the underlying processes.
has been photo-ionized, and the dust grains are exposed to cumstellar material has undergone major changes. The gas descendants of carbon-rich AGB stars, in which the cir-

5.3. Comparison with PNe

Carbon-rich planetary nebulae (PNe) are the immediate descendants of carbon-rich AGB stars, in which the circumstellar material has undergone major changes. The gas descendant of a carbon-rich AGB star, in which the circumstellar material has undergone major changes. The gas has been photo-ionized, and the dust grains are exposed to intense radiation fields and high ($\sim 10^8$ K) temperatures. A comparison of depletions in AGB stars and PNe may therefore reveal some aspects of grain evolution.

Element abundances have been extensively measured in the ionized gas in PNe. The abundances are, however, subject to systematic, and sometimes, large uncertainties, and relatively few metals have accessible lines covering the appropriate stages of ionization. For comparison with IRC+10216 we focus on NGC 7027, which is one of the more similar. Na and K are the least depleted, at compara-

5.4. Comparison with the ISM

It is also of interest to compare our results for IRC+10216 with depletions in the ISM. We take the line of sight towards $\zeta$ Oph as representative of the ISM, bearing in mind that the overall level of IS depletions varies with location but the general pattern remains the same. The depletions towards $\zeta$ Oph, from Savage & Sembach (1996), are listed in column (10) of Table 3.

In spite of the different physical and chemical environments that lead to grain formation in IRC+10216 and the ISM, it can be seen that the depletion patterns are qualita-

Table 3. Abundances in the envelope of IRC+10216

| El. | Ion | $N$ (cm$^{-2}$) | $C_i$ | $N$(I+II) (cm$^{-2}$) | $N$(I+II)/$N$(H) | $X_\odot$ | $\log_{10} \delta$ | $\log_{10} \delta_{Oph}$ |
|-----|-----|---------------|-------|----------------------|------------------|----------|-----------------|-----------------|
| Na  | i   | 4.6(14)       | 1.22  | 5.6(14)              | 4.2(−7)          | 2.00(−6) | −0.68           | −0.06           | −0.95           |
| Al  | <2.8(12) | 3.5(3)      | <9.8(15) | <7.3(−6)            | 2.88(−6)         | ...      | ...             | ...             |
| K   | i   | 4.6(12)       | 1.60  | 7.4(12)              | 5.5(−9)          | 1.29(−7) | −1.37           | −0.17           | −1.09           |
| Ca  | i   | 1.9(12)       | 4.70  | 8.9(12)b            | 6.6(−9)          | 2.19(−6) | −2.52           | −0.75           | −3.73           |
| Ti  | i   | <1.9(12)      | 4.26  | <3.9(12)b            | <2.9(−9)         | 8.32(−8) | <−1.45          | ...             | −3.02           |
| Cr  | <2.0(12) | 1.31         | ...   | ...                 | ...              | ...     | ...             | ...             |
| Mn  | <3.1(12) | 2.21         | <6.9(12) | <5.1(−9)          | 3.16(−7)         | <−1.79  | −1.45           | ...             |
| Fe  | i   | 8.8(13)       | 2.58  | 2.3(14)             | 1.7(−7)          | 2.95(−5) | −2.24           | −1.68           | −2.27           |
| Sr  | <2.7(11) | 3.82         | <1.0(12) | <7.6(−10)       | 8.13(−10)        | ...     | ...             | ...             |

a. Ionization correction $G_i = N(I + II)/N(I)$ for neutral species and $N(I + II)/N(II)$ for singly ionized species.

b. Based on observed values of $N(I) + N(II)$; independent of $C_i$.

c. Upper limit on $\delta > X_\odot$, no useful limit on $\delta$.
Table 4. Metal bearing molecules in IRC+10216

| Species   | θ_c (″) | X^b_{mol} | X_{mol}/X_⊙ | Ref. |
|-----------|---------|------------|--------------|------|
| NaCl      | 2.5     | 1.3(−09)   | 6.3(−4)     | 1    |
| NaCN      | 2.5     | 6.8(−09)   | 3.4(−3)     | 1    |
| KCl       | 2.5     | 3.8(−10)   | 2.9(−3)     | 2    |
| AIF       | 2.5     | 4.6(−08)   | 1.6(−2)     | 1    |
| AlCl      | 2.5     | 2.8(−08)   | 9.8(−3)     | 1    |
| AINC      | 5−15    | 2.3(−10)   | 7.9(−5)     | 3    |
| MgNC      | 10−20   | 6.3(−09)   | 1.8(−4)     | 1    |
| MgCN      | 10−20   | 2.9(−10)   | 8.0(−6)     | 4    |

a Source radius.
* Abundance relative to H, for d = 120 pc, M(H) = 1.25 × 10^-5 M_⊙ yr^-1.

References: (1) Highberger & Ziurys (2003). (2) Assuming X(KCl) = 0.3X(NaCl) Cernicharo & Guélin (1987).
(3) Ziurys et al. (2002). (4) Assuming X(MgCN) = 0.045X(MgNC) Ziurys et al. (1995).

6. Metal chemistry

The circumstellar envelope of IRC+10216 exhibits a remarkable gas phase chemistry, with more than 50 molecular species detected to date, mainly through their rotational transitions at millimeter wavelengths. The majority of the molecules are formed of the abundant elements H, C, N, and O, several include Si and S, but a small number unexpectedly include metals, Na, K, Mg, and Al. The first of these were discovered by Cernicharo & Guélin (1987). Our observations of gas phase metal atoms in IRC+10216 constrain some aspects of the metal chemistry in the envelope.

The metal bearing molecules detected in the envelope are listed in Table 4. In addition to these, numerous other metal bearing species have been searched for at comparable levels, but not detected (e.g., Turner 1995). There are also other metal species whose rotational frequencies have not yet been measured in the laboratory (e.g., Ziurys 2006b).

For comparison with the atomic abundances reported here, we list updated abundances for the metal molecules in column (3) of Table 4. These are based on the column densities given in the references in the table, but are derived for the distance and mass loss rate used in this paper, and are relative to hydrogen. The fractional abundance X_{mol}/X_⊙ is the fraction of each metal in a particular molecular species, θ_c is the measured, or inferred, angular radius of the molecular distribution, adopted from the references.

The metal bearing molecules divide naturally into two groups. The first is confined to the dense core region represented by θ_c ≲ 2.5″ in the table. The second, which includes only Mg and Al bearing molecules, is found in distinct shells in the envelope, in the photo-dissociation region. The general level of incorporation of the metals into molecules is very small in both groups.

In the outer envelope, the fractional abundances of the metal bearing molecules are ≲ 0.02 per cent, while the atoms that we detect have fractional abundances of 0.3–20 per cent. This we used in Sect. 5.1 to justify the assumption that the observed atoms are the dominant gas phase metal carriers along the line of sight.

Even in the core region the molecules do not seem to be dominant. The observed fraction of Na in the core in the form of molecules is ∼ 0.4 per cent, and we detect 20 percent in the form of atoms farther out. Similarly, the fraction of K in the core in the form of molecules is ∼ 0.3 per cent, and we detect 4 per cent in atoms farther out. For these species the numbers are consistent with the idea that apart from condensation or adsorption onto dust grains, the dominant gas phase species is atomic throughout the envelope. There is lack of specific molecular information for the other metals, but the absence of detected molecular species argues that this applies to them as well.

Our finding that residual metals are present in the envelope in the form of neutral atoms and ions provides an observational foundation for understanding certain aspects of the metal chemistry. For example, it has been proposed that metal cyanides are formed by reactions of metal ions with cyanopolyynes (Dunbar & Petrie 2002), and the model of Cordier & Millar (2009) shows that this can account for the observed abundance of MgCN if enough Mg⁺ is present in the gas phase. We have not observed Mg, but we expect that it behaves like the other metals. Its condensation temperature is significantly less than that of Ca and Fe, so it is likely to be less depleted in the envelope. The gas phase abundances of the metals that we observe are given in column (6) of Table 3. These may undergo reactions with cyanopolyynes and other abundant neutral molecules, such as unsaturated hydrocarbons, and lead to a variety of metal bearing species. The largest abundances that we observe in the gas are those of Na and Fe (whose large depletion is balanced by its high cosmic abundance). Thus Fe offers interesting possibilities for a potentially observable Fe chemistry.

7. Conclusions

The observations of IRC+10216 reported in this paper represent the first comprehensive study of atomic metals in a carbon-rich circumstellar envelope. We detect lines of Na, K, Ca, Cr, and Fe, and obtain upper limits for Al, Ti, Mn, and Sr. Combined with a simple model of the ionization, the observations provide estimates of the gas phase metal abundances in the outer envelope.

The results show that the metals, especially Ca and Fe, are significantly depleted onto dust grains in the circumstellar envelope, and this is the dominant form returned to the ISM. The depletion pattern has some similarity with depletion in the ISM, and is roughly consistent with expectations of dust condensation in a carbon-rich envelope.

Although the metals are depleted in the envelope, atomic metals in the form of neutral atoms and ions appear to be the major metal species in the gas phase. As such, they likely play a key role in the metal chemistry of the envelope.

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References

Baeker, E. J., van Dishoeck, E. F., Waters, L. B. F. M., & Schoenmaker, T. 1997, A&A, 323, 469
Bernard Salas, J., Pottasch, S. R., Beintema, D. A., & Wesselius, P. R. 2001, A&A, 367, 949
Bernat, A. P. 1976, Ph.D. Thesis, Cherchneff, I., Glassgold, A. E., & Mamon, G. A. 1993, ApJ, 410, 188
Cernicharo, J., & Guelin, M. 1987, A&A 183, L10
Chaffee, F. H., Jr., & White, R. E. 1982, ApJS, 50, 169
Coelho, P., Barbuy, B., Melendez, J., Schiavon, R. P., & Castilho, B. V. 2006, A&A, 443, 735
Cowley, M. A., & Millar, T. J. 2009, ApJ, 697, 68
Cordiner, M. A., Millar, T. J., Herbst, E., Chuínin, R. N., & Walsh, C. 2007, Molecules in Space and Laboratory, eds J.L. Lemaire & F. Combes, p98
Cox, P., Huggins, P. J., Maillard, J.-P., Habart, E., Morisset, C., Bachiller, R., & Forveille, T. 2002, A&A, 384, 603
Delbouille, L., Neven, L., Roland, G. 19973, Photometric Atlas of the Solar Spectrum $\lambda_{3000}$ to $\lambda_{10000}$ (Institut d’Astrophysique, University of Liège)
Delgado Inglada, G., Rodríguez, M., Mampaso, A., & Vironen, K. 2009, ApJ, 694, 1335
Draine, B. T. 2009, arXiv:0903.1658
Dunbar, R. C., & Petrie, S. 2002, ApJ, 564, 792
Field, G. B. 1974, ApJ, 187, 453
Gillman, R. C. 1969, ApJ, 155, L185
Glassgold, A. E., & Huggins, P. J. 1986, ApJ, 306, 605
Highberger, J. L., & Ziurys, L. M. 2003, ApJ, 597, 1065
Huggins, P. J., & Glassgold, A. E. 1982, ApJ, 252, 201
Huggins, P. J., & Healy, A. P. 1986, ApJ, 304, 418
Huggins, P. J., Olofsson, H., & Johansson, L. E. B. 1988, ApJ, 332, 1009
Jura, M., & Morris, M. 1985, ApJ, 292, 487
Kendall, T. R., Mauron, N., McCombie, J., & Sarre, P. J. 2002, A&A, 387, 624
Keyes, C. D., Aller, L. H., & Feibelman, W. A. 1990, PASP, 102, 59
Kingdon, J., Ferland, G. J., & Feibelman, W. A. 1995, ApJ, 439, 793
Kwan, J., & Hill, F. 1977, ApJ, 215, 781
Lafont, S., Lucas, R., & Omont, A. 1982, A&A, 106, 201
Lodders, K. 2003, ApJ, 591, 1220
Lodders, K., & Fegley, B., Jr. 1999, Asymptotic Giant Branch Stars, 191, 279
Lodders, K., & Fegley, B., Jr. 1995, Meteoritics, 30, 661
Loup, C., Forveille, T., Omont, A., & Paul, J.F. 1993, A&ASS, 99, 291
Martin, P. G., & Rogers, C. 1987, ApJ, 322, 374
Mauron, N., & Huggins, P. J. 1999, A&A 349, 203
Mauron, N., & Huggins, P. J. 2000, A&A, 359, 707
Mauron, N., De Laverny, P., Lopez, B. 2003, A&A, 401, 985
McCabe, E. M., Smith, R. C., & Clegg, R. E. S. 1979, Nature, 281, 263
Middlemass, D. 1990, MNRAS, 244, 294
Morris, M., & Jura, M. 1983, ApJ, 264, 546
Morton, D. C. 1991, ApJS, 77, 119
Morton, D. C. 2000, ApJS, 130, 403
Olofsson, H. 2005, ESA Special Publication, 577, 223
Péquignot, D., & Aldrovandi, S. M. V. 1986, A&A, 161, 169
Ramstedt, S., Schöier, F. L., Olofsson, H., & Lundgren, A. A. 2008, A&A, 487, 645
Savage, B. D., & Sembach, K. R. 1996, ARA&A, 34, 379
Schöier, F. L., & Olofsson, H. 2001, A&A, 368, 969
Tsuji, T. 1973, A&A, 23, 411
Turner, B. E. 1995, Ap&SS, 224, 297
Young, K., Phillips, T. G., & Knapp, G. R. 1993, ApJSS, 86, 517
Zhang, Y., Liu, X.-W., Luo, S.-G., Péquignot, D., & Barlow, M. J. 2005, A&A, 442, 249
Ziurys, L. M. 2006a, Proceedings of the National Academy of Science, 103, 12274
Ziurys, L. M. 2006b, Astrochemistry - From Laboratory Studies to Astronomical Observations, 855, 231
Ziurys, L. M., Apponi, A. J., Guelin, M., & Cernicharo, J. 1995, ApJ, 445, L47
Ziurys, L. M., Savage, C., Highberger, J. L., Apponi, A. J., Guelin, M., & Cernicharo, J. 2002, ApJ, 564, L45