Comment on "Phosphine in the Venusian Atmosphere: A Strict Upper Limit from SOFIA GREAT Observations" by Cordiner et al.

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March 9, 2023

Abstract

Searches for phosphine in Venus' atmosphere have sparked a debate. Cordiner et al. 2022 analyse spectra from the Stratospheric Observatory For Infrared Astronomy (SOFIA) and infer <0.8 ppb of PH3. We noticed that some spectral artefacts arose from non-essential calibration-load signals. By-passing these signals allows simpler post-processing and a 5.7σ candidate detection, suggesting ~3 ppb of PH3 above the clouds. Compiling six phosphine results hints at an inverted abundance trend: decreasing above the clouds but rising again in the mesosphere from some unexplained source. However, no such extra source is needed if phosphine is undergoing destruction by sunlight (photolysis), to a similar degree as on Earth. Low phosphine values/limits are found where the viewed part of the super-rotating Venusian atmosphere had passed through sunlight, while high values are from views moving into sunlight. We suggest Venusian phosphine is indeed present, and so merits further work on models of its origins.

1 Comment on

2 "*Phosphine in the Venusian Atmosphere: A Strict Upper Limit from SOFIA*
3 *GREAT Observations*"

4 by Cordiner et al.

5 **Jane S. Greaves[1], Janusz J. Petkowski[2,3], Anita M. S. Richards[4], Clara Sousa-Silva[5], Sara**
6 **Seager[2,6,7] and David L. Clements[8]**

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23 **Key Points:**

24 • We recover Venusian phosphine in *SOFIA* spectra by reducing contaminating signals; the
25 PH_3 abundance is ~3 part-per billion (ppb).

26 • Six recoveries/limits show PH_3 depleting between clouds and mesosphere, which would
27 require an unknown re-formation process or extra source.

28 • Recoveries and upper limits can instead be reconciled by PH_3 photolysis, as high/low
29 abundances correspond to Venusian mornings/evenings.
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Abstract

Searches for phosphine in Venus' atmosphere have sparked a debate. Cordiner et al. 2022 analyse spectra from the Stratospheric Observatory For Infrared Astronomy (SOFIA) and infer <0.8 ppb of PH_3. We noticed that some spectral artefacts arose from non-essential calibration-load signals. By-passing these signals allows simpler post-processing and a 5.7σ candidate detection, suggesting ~3 ppb of PH_3 above the clouds. Compiling six phosphine results hints at an inverted abundance trend: decreasing above the clouds but rising again in the mesosphere from some unexplained source. However, no such extra source is needed if phosphine is undergoing destruction by sunlight (photolysis), to a similar degree as on Earth. Low phosphine values/limits are found where the viewed part of the super-rotating Venusian atmosphere had passed through sunlight, while high values are from views moving into sunlight. We suggest Venusian phosphine is indeed present, and so merits further work on models of its origins.

Plain Language Summary

Cordiner et al. find no phosphine in Venus' atmosphere, using the airborne SOFIA telescope. By-passing some instrumental effects, we extract a detection with 5.7σ-confidence from the same data. We can resolve the tension between high and low PH_3 abundance values by noticing that the former are from 'mornings' in Venus' atmosphere and the latter from 'evenings'. Sunlight reduces the amount of phosphine in Earth's atmosphere by an order of magnitude, so similarly on Venus, we might expect lower abundances in data taken when the part of the atmosphere observed has passed through sunlight. If the six available datasets can be reconciled in this way, further modelling of possible sources of PH_3 (e.g. volcanic, disequilbrium chemistry, extant life) seem worthwhile.

1 Introduction

Phosphine, if present in Venus' atmosphere, would be unexpected on an oxidised planet. Greaves et al. (2021) searched for PH_3 absorption at 1 mm wavelength, testing the concept that this molecule may be a biosignature when seen in anoxic environments. The unexpected detection-candidates from *JCMT* and *ALMA* have stimulated much community work on robust spectral processing, and on other methods to detect PH_3 at Venus, mostly proving negative except for an in-situ mass-spectrometry recovery (Mogul, Limaye, Way, et al., 2021). Particularly deep (above-cloud) limits have been set by infrared spectroscopy (Encrenaz et al., 2020; Trompet et al., 2020).

We comment here on the findings of Cordiner et al. (2022), hereafter C22, who present a deep upper limit from PH_3 observations with the *GREAT* instrument on *SOFIA*. They propose that *all* the candidate detections of phosphine in Venus' atmosphere could in fact be null results, given the complexity of the challenging observations – although their Figure 1 omits findings by Greaves et al. (2022), our work after calibration and contamination issues were fully resolved, where we find self-consistent 6-8σ detections for the PH_3 J=1-0 line from *JCMT* and *ALMA*.

The C22 observations are of the rotational transitions J=4-3 and 2-1 (around 1 and 0.5 THz), uniquely accessible to the *SOFIA* airborne telescope, and complementary to the existing J=1-0 spectra (at 0.27 THz). From their J=4-3 data processing, C22 find an upper limit of 0.8 ppb of PH_3, applicable to most of the planet and 75+ km altitudes, while their J=2-1 results suggest ~2.3 ppb could be present but only with 1.5σ confidence. These abundances are difficult to

73   reconcile with ~20 ppb levels from the J=1-0 data, without invoking strong temporal-variations

74   or steep gradients over the slightly different altitudes these lines trace.

75   **2 Materials and Methods**

76   C22 note the existence in the *GREAT* spectra of quasi-periodic fringe patterns, due to

77   standing waves between optical elements and to frequency-dependent gain factors used in

78   calibration. Their calibration to antenna temperatures T_A follows the standard method of dividing

79   the power difference of on- and off-Venus spectra by the power difference of hot and cold

80   calibration-load signals, and then multiplying by the temperature difference of the hot and cold

81   loads. We noticed that much of the fringing is introduced because the standing waves differ

82   when observing the sky and the calibration loads. However, calibration to T_A is not essential in

83   measuring the line-to-continuum ratios, *l/c*, from which abundances derive. In the case of the

84   PH_3 J=4-3 line components (seen by the "4G2 pixel"), an alternative is

85
$$l/c = (On_{line} - Off_{line}^*) \, / \, [0.5(On - Off)] \qquad [1]$$

86   where *On* and *Off* are the spectra on Venus and on adjacent blank sky, the subscript *line* indicates

87   the broadband signals have been subtracted, and *Off*$_{line}$* represents the instrumental line-signal

88   that *GREAT* would see for a featureless patch of sky of similar brightness to Venus. *Off*$_{line}$* was

89   generated by multiplying *Off*$_{line}$ by a factor ~1.05 and adjusting this scalar until the residual

90   *(On*$_{line}$ *-Off*$_{line}$*)* was minimised – that is, the procedure tests the null hypothesis, that no

91   absorbing gases are present. Smooth fits to *On* and *Off* (the continuum signals) were used in the

92   denominator of [1] to further minimise noise. The factor of 0.5 arises because *GREAT* is a

93   double-sideband instrument with approximately equal sideband gains (see C22), and so records

94   the planetary continuum twice. The Eq. [1] method has worked well here for PH_3 J=4-3, reaching

95   a similar noise level to C22, but our approach failed for the PH_3 J=2-1 line-pair ("4G1 pixel")

96   because the ripples differ between *On*$_{line}$ and *Off*$_{line}$.

97   Remaining ripples in the J=4-3 spectra were then removed by a one-stage Fourier

98   process, contrasting to the iterative 7-step Lomb-Scargle periodogram approach used by C22 (or

99   traditional polynomial fitting, which is less useful for spectra with many ripples). Both we and

100   C22 similarly "masked" the spectral regions where the four PH_3 components lie, to avoid fitting

101   real lines as if they are ripples. C22's periodogram method works intrinsically on masked data,

102   while we interpolated across the line regions with quadratic fits anchored on adjacent spectral

103   pixels. We used 3-sigma cuts in Fourier space, with features above these cuts inverse-Fourier-

104   transformed to create a family of model sinusoids. Subtracting these model baselines yields well-

105   flattened spectra (Figures 1a, 1b). Finally we "stack" the 24 samples of PH_3 absorption in the

106   data (Figure 1c), namely from the 6 observations and their 4 spectral sections covering the J=4-3

107   components, as these are of similar intrinsic line-depth (see Figure 3 in C22). Two of the PH_3

108   features are close together (Figure 1b), and so to avoid duplications, we replaced the secondary

109   occurrences with noise values from elsewhere in the spectrum, before making the final stack.

110   Several robustness checks were run, exploring possible processing issues.

111   • The net result could be dominated by a few strong artefacts. This was found not to be the

112   case, with the line-integrals from the 24 samples following an approximately normal

113   distribution. The result is also robust to removing a few points. For example, the final

114   observation (#040402) was the noisiest, and removing all these samples from the stack shifts

115   the net line-integral by only -20%.

116   • There could be terrestrial atmospheric signals in the band, that affect the validity of
117   minimising the numerator in Eq. [1]; in particular, C22 note two frequencies where there may
118   be O_3 absorption. We re-calculated the scalar used in Eq. [1] with the O_3 regions blanked,
119   and found negligible change (around 1% reduction in noise).
120   • The interpolation across the PH_3 line-regions might not be following the correct trend,
121   potentially always producing negative residuals that mimic absorption lines. We tested this
122   by re-running the processing identically, but shifting the spectral sections to parts of the band
123   without phosphine features. From 50 tests, none produced a result like Figure 1c, namely a
124   "fake" absorption that is the only feature, centrally placed, and of comparable width to line
125   models (see below).
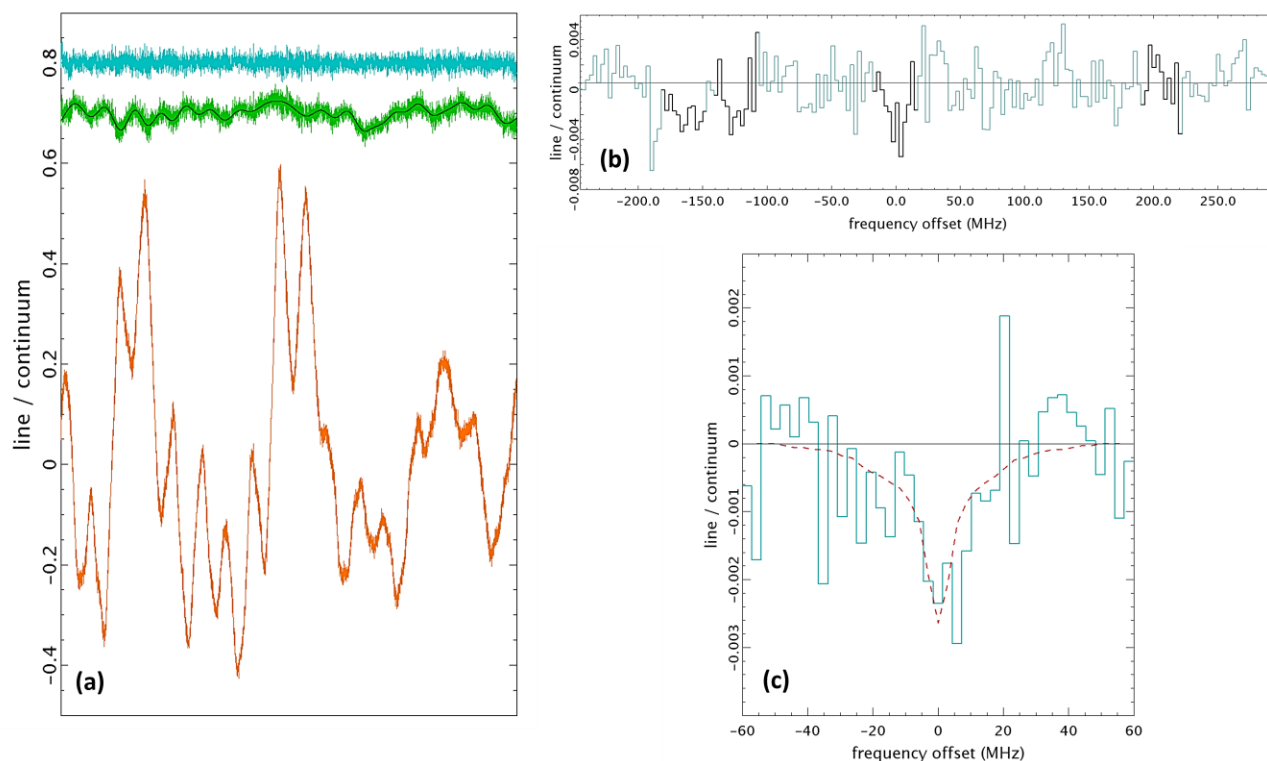
127   **3 Results**

128   Three of the four expected PH_3 J=4-3 components are visible when all the observations
129   are co-added (Figure 1b), while only one component was apparent in Figure 3 of C22. In Figure
130   1c, our final stacked spectrum indicates an overall 5.7σ detection of PH_3 J=4-3, when integrated
131   over ±17 MHz (the masked region). This confidence level changes marginally (by ±0.4σ) if a
132   different range of spectral pixels is used to calculate the zero-level.

133   We then modelled our net spectrum using the same online tool as C22. We ran a model
134   for 1 ppb of phosphine and scaled it linearly for different abundances, and then calculated a
135   reduced-chi-squared statistic to assess goodness of fit. Figure 1c illustrates that 3 ppb of PH_3
136   provides a good match to the observed line, with χ^2_r of 0.75. (The uncertainties used in χ^2_r were
137   generated per spectral pixel from the internal data dispersion.) We consider that fitting the
138   stacked spectrum better mitigates against artefacts, compared to matching the individual
139   components (Figure 1b) against the model. For example, inserting a single positive "spike" near
140   the highest-frequency component (Figure 1b) was found to significantly reduce the inferred PH_3
141   abundance, as the χ^2_r test attempts to minimise the discrepancy of positive data against a
142   negative model-line. As a positive feature does appears here in this part of C22's spectrum (their
143   Figure 3), this could have driven their upper limit down to the ≤0.8 ppb they obtain. However,
144   we do not rule out an abundance as low as 0.8 ppb, which is at the lower 99% confidence-bound
145   in our χ^2_r tests. We also note that C22 estimated ~2.3 ppb from their PH_3 J=2-1 spectrum, albeit
146   at only +1.5σ confidence, and this estimate is compatible with our 3 ppb result.

147   C22 (Figure S4) find that altitudes around ~80 km are the best-sampled at the PH_3 J=4-3
148   line-frequency; our recovery here is consistent with the model that we both use, and with the
149   predicted short lifetime of PH_3 above ~80 km (Bains, Petkowski, Seager, et al., 2021). Altitudes
150   are however uncertain because the PH_3-CO_2 pressure-broadening coefficient has not been
151   experimentally verified. We also note that all GHz/THz data are limited by the spectral span that
152   can be recovered. Here, any absorption wider than ~200 MHz leads to merged PH_3 J=4-3
153   components, and so any phosphine signatures below ~70 km (roughly cloud-top level) are lost.

154   *Figure 1. The process of extracting the PH_3 J=4-3 signal from the GREAT data is illustrated.*
155   *Panel (a) illustrates processing of 6000 spectral pixels from observation #040093 from the first*
156   *SOFIA flight (the reference spectral pixel is mid-band; vertical offsets are for clarity only). The*
157   *lower orange histogram is from a standard On-Off processing; the middle green histogram is the*
158   *result after applying the Eq.[1] step, and is overlaid with the Fourier-derived trend (black*
159   *curve); the top blue histogram is the flattened output after subtracting this trend. In panel (b),*

160   *the blue histogram is the unweighted average of all six observations made over the three flights,*
161   *with the sections containing the four PH₃ J=4-3 components highlighted in black. The spectral*
162   *pixels are binned in groups of 12 in (b) and (c) to improve clarity, and the thin black lines show*
163   *the zero-level correction made between (b) and (c). Panel (c) shows (blue histogram) the*
164   *unweighted stack of all 24 spectral sections containing PH₃ J=4-3 features. The dashed red*
165   *curve is a model for 3 ppb of phosphine, generated vis the Planetary Spectrum Generator (PSG)*
166   *(https://psg.gsfc.nasa.gov/index.php) following C22. The four line-components from the PSG*
167   *model were masked and stacked similarly to the data.*

4 Discussion

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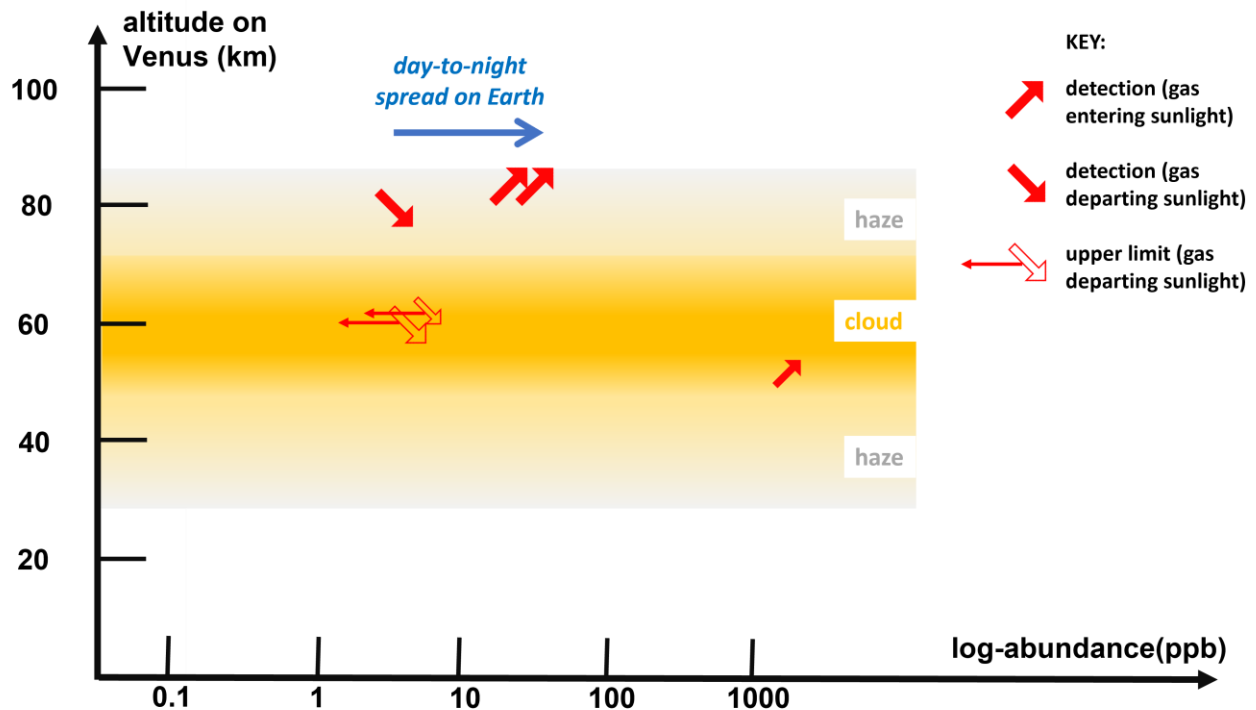169   Debates continue about the best methods to acquire and process deep GHz/THz spectra
170 of Venus. These observations are very challenging in dynamic range, as Venus is so bright,
171 revealing "ripples" in spectral baselines that are not evident in more typical telescope usage.
172 Depending on preferred approaches, different authors argue for between zero and three published
173 detections of rotational (J) transitions of PH₃.

174   We can compare results from the data discussed here with the outcomes of other searches
175 for phosphine at Venus, and assess whether this results in a plausible altitude profile of the
176 molecule (Figure 2). The trend found by connecting the results from six searches for phosphine
177 appears as an upwards decline that then reverses, i.e. PH₃ that is depleted somewhere between
178 ~50 km and ~80 km. This is hard to explain in the absence of a chemical route to reform the
179 molecules, or a new mesospheric source. The order-of-magnitude contrast between some of the
180 candidate detections and the upper limits has led to doubts over the presence of phosphine.

181   However, we noticed that this divide is also between observations made when the
182 'morning' versus the 'evening' sides of Venus' atmosphere were targeted – and this is relevant in
183 gas-mixing processes (e.g. (Lefèvre et al., 2022)). Where the gas observed on Venus has
184 travelled through sunlight and is descending towards the night-side of the planet, we detect at
185 most the ~3 ppb of phosphine estimated here. In contrast, where gas is rising into sunlight, we
186 observe ≥ ~20 ppb of PH_3. Hence photolysis – similar to the observed destruction of terrestrial
187 phosphine by sunlight (Sousa-Silva et al., 2020) – could explain the split between high and low
188 phosphine abundances observed on Venus. It is striking that this evening/morning difference is
189 comparable in magnitude to the factor ~10 difference over night/day for phosphine in the Earth's
190 atmosphere (Glindemann et al. 1996; noting absolute abundances are much lower on Earth).

191 *Figure 2. The trend of phosphine abundances by altitude is sketched. Symbols indicate candidate*
192 *detections plus best upper limits for phosphine abundances. Rising arrows indicate observations*
193 *made where the super-rotating atmosphere was rising into sunlight and falling arrows indicate*
194 *observations made where the atmosphere was descending towards the nightside (see key). Large*
195 *and small symbols indicate that a large fraction of the planet area was observed, or that a small*
196 *region was sampled, respectively. Abundance estimates are, from top: ~20, 25 ppb from J=1-0*
197 *data (via (Greaves et al., 2022) and with altitude proposed by C22); 3 ppb from J=4-3 data (this*
198 *work; beam centred on the evening side); <7 ppb at 62 km from 4 μm spectra (Trompet et al.,*
199 *2020: low-latitude data to best match whole-planet studies); <5 ppb at 60 km from 10 μm*
200 *spectra (Encrenaz et al., 2020: latitudes within ±50°); ~2 ppm at 51 km from Pioneer-Venus in-*
201 *situ sampling during descent (Mogul, Limaye, Way, et al., 2021). The blue arrow indicates the*
202 *ten-fold increase of terrestrial phosphine from day to night (Glindemann et al. 1996) – note the*
203 *arrow's plotted position is arbitrary; Earth hosts much lower PH_3 than Venus.*

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5 Conclusions

The question regarding phosphine in Venus' atmosphere is likely to be debated for some time. A further *JCMT* survey is ongoing (see https://www.eaobservatory.org/jcmt/science/large-programs/jcmt-venus-monitoring-phosphine-and-other-molecules-in-venuss-atmosphere/; PI. D. Clements) and is producing open-source data that should yield more definitive answers – in particular, that team is now processing broadband spectra that can sample the cloud decks. The most direct answer regarding phosphine could come from new in-situ sampling, potentially with the addition of one laser channel to the Venus Tunable Laser Spectrometer (VTLS) instrument on-board the *DAVINCI* descent probe (Garvin et al. 2022).

The origins of any phosphine present are also debated, and most scenarios are hard to test for lack of some contextual data. For example, it seems only extraordinary volcanic activity could make ~ppb-level phosphine (Bains et al., 2022) but vulcanism on Venus is not well understood. In some new avenues, (Ferus et al., 2022) discuss abiotic routes to phosphine involving redox disequlibrium, while others (Bains, Petkowski, Rimmer, et al., 2021; Mogul, Limaye, Lee, et al., 2021) explore phototrophic life and the habitability of the clouds. We conclude that establishing an improved PH_3 altitude-profile is worthwhile to test these new models of origins.

Open Research

The SOFIA Level 1 data are available under project id 75_0059_1 through the public data archive at https://irsa.ipac.caltech.edu/applications/sofia. The custom software to generate the data shown in the figures is supplied at https://zenodo.org/record/7692288#.ZAC363bP3IU. The script requires the UK-Starlink software (Currie et al., 2014) which is currently supported by the East Asian Observatory and available at https://starlink.eao.hawaii.edu/starlink/2021ADownload.

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61 2020; Trompet et al., 2020).

62   We comment here on the findings of Cordiner et al. (2022), hereafter C22, who present a
63 deep upper limit from PH_3 observations with the *GREAT* instrument on *SOFIA*. They propose
64 that *all* the candidate detections of phosphine in Venus' atmosphere could in fact be null results,
65 given the complexity of the challenging observations – although their Figure 1 omits findings by
66 Greaves et al. (2022), our work after calibration and contamination issues were fully resolved,
67 where we find self-consistent 6-8σ detections for the PH_3 J=1-0 line from *JCMT* and *ALMA*.

68   The C22 observations are of the rotational transitions J=4-3 and 2-1 (around 1 and 0.5
69 THz), uniquely accessible to the *SOFIA* airborne telescope, and complementary to the existing
70 J=1-0 spectra (at 0.27 THz). From their J=4-3 data processing, C22 find an upper limit of 0.8 ppb
71 of PH_3, applicable to most of the planet and 75+ km altitudes, while their J=2-1 results suggest
72 ~2.3 ppb could be present but only with 1.5σ confidence. These abundances are difficult to

73 reconcile with ~20 ppb levels from the J=1-0 data, without invoking strong temporal-variations
74 or steep gradients over the slightly different altitudes these lines trace.

75 **2 Materials and Methods**

76 C22 note the existence in the *GREAT* spectra of quasi-periodic fringe patterns, due to
77 standing waves between optical elements and to frequency-dependent gain factors used in
78 calibration. Their calibration to antenna temperatures T_A follows the standard method of dividing
79 the power difference of on- and off-Venus spectra by the power difference of hot and cold
80 calibration-load signals, and then multiplying by the temperature difference of the hot and cold
81 loads. We noticed that much of the fringing is introduced because the standing waves differ
82 when observing the sky and the calibration loads. However, calibration to T_A is not essential in
83 measuring the line-to-continuum ratios, l/c, from which abundances derive. In the case of the
84 PH$_3$ J=4-3 line components (seen by the "4G2 pixel"), an alternative is

85 $$l/c = (On_{line} - Off_{line}^*) / [0.5(On - Off)] \qquad [1]$$

86 where *On* and *Off* are the spectra on Venus and on adjacent blank sky, the subscript *line* indicates
87 the broadband signals have been subtracted, and *Off*$_{line}$* represents the instrumental line-signal
88 that *GREAT* would see for a featureless patch of sky of similar brightness to Venus. *Off*$_{line}$* was
89 generated by multiplying *Off*$_{line}$ by a factor ~1.05 and adjusting this scalar until the residual
90 *(On$_{line}$ -Off$_{line}$*)* was minimised – that is, the procedure tests the null hypothesis, that no
91 absorbing gases are present. Smooth fits to *On* and *Off* (the continuum signals) were used in the
92 denominator of [1] to further minimise noise. The factor of 0.5 arises because *GREAT* is a
93 double-sideband instrument with approximately equal sideband gains (see C22), and so records
94 the planetary continuum twice. The Eq. [1] method has worked well here for PH$_3$ J=4-3, reaching
95 a similar noise level to C22, but our approach failed for the PH$_3$ J=2-1 line-pair ("4G1 pixel")
96 because the ripples differ between *On*$_{line}$ and *Off*$_{line}$.

97 Remaining ripples in the J=4-3 spectra were then removed by a one-stage Fourier
98 process, contrasting to the iterative 7-step Lomb-Scargle periodogram approach used by C22 (or
99 traditional polynomial fitting, which is less useful for spectra with many ripples). Both we and
100 C22 similarly "masked" the spectral regions where the four PH$_3$ components lie, to avoid fitting
101 real lines as if they are ripples. C22's periodogram method works intrinsically on masked data,
102 while we interpolated across the line regions with quadratic fits anchored on adjacent spectral
103 pixels. We used 3-sigma cuts in Fourier space, with features above these cuts inverse-Fourier-
104 transformed to create a family of model sinusoids. Subtracting these model baselines yields well-
105 flattened spectra (Figures 1a, 1b). Finally we "stack" the 24 samples of PH$_3$ absorption in the
106 data (Figure 1c), namely from the 6 observations and their 4 spectral sections covering the J=4-3
107 components, as these are of similar intrinsic line-depth (see Figure 3 in C22). Two of the PH$_3$
108 features are close together (Figure 1b), and so to avoid duplications, we replaced the secondary
109 occurrences with noise values from elsewhere in the spectrum, before making the final stack.

110 Several robustness checks were run, exploring possible processing issues.

111 • The net result could be dominated by a few strong artefacts. This was found not to be the
112 case, with the line-integrals from the 24 samples following an approximately normal
113 distribution. The result is also robust to removing a few points. For example, the final
114 observation (#040402) was the noisiest, and removing all these samples from the stack shifts
115 the net line-integral by only -20%.

116   • There could be terrestrial atmospheric signals in the band, that affect the validity of
117   minimising the numerator in Eq. [1]; in particular, C22 note two frequencies where there may
118   be O_3 absorption. We re-calculated the scalar used in Eq. [1] with the O_3 regions blanked,
119   and found negligible change (around 1% reduction in noise).
120   • The interpolation across the PH_3 line-regions might not be following the correct trend,
121   potentially always producing negative residuals that mimic absorption lines. We tested this
122   by re-running the processing identically, but shifting the spectral sections to parts of the band
123   without phosphine features. From 50 tests, none produced a result like Figure 1c, namely a
124   "fake" absorption that is the only feature, centrally placed, and of comparable width to line
125   models (see below).
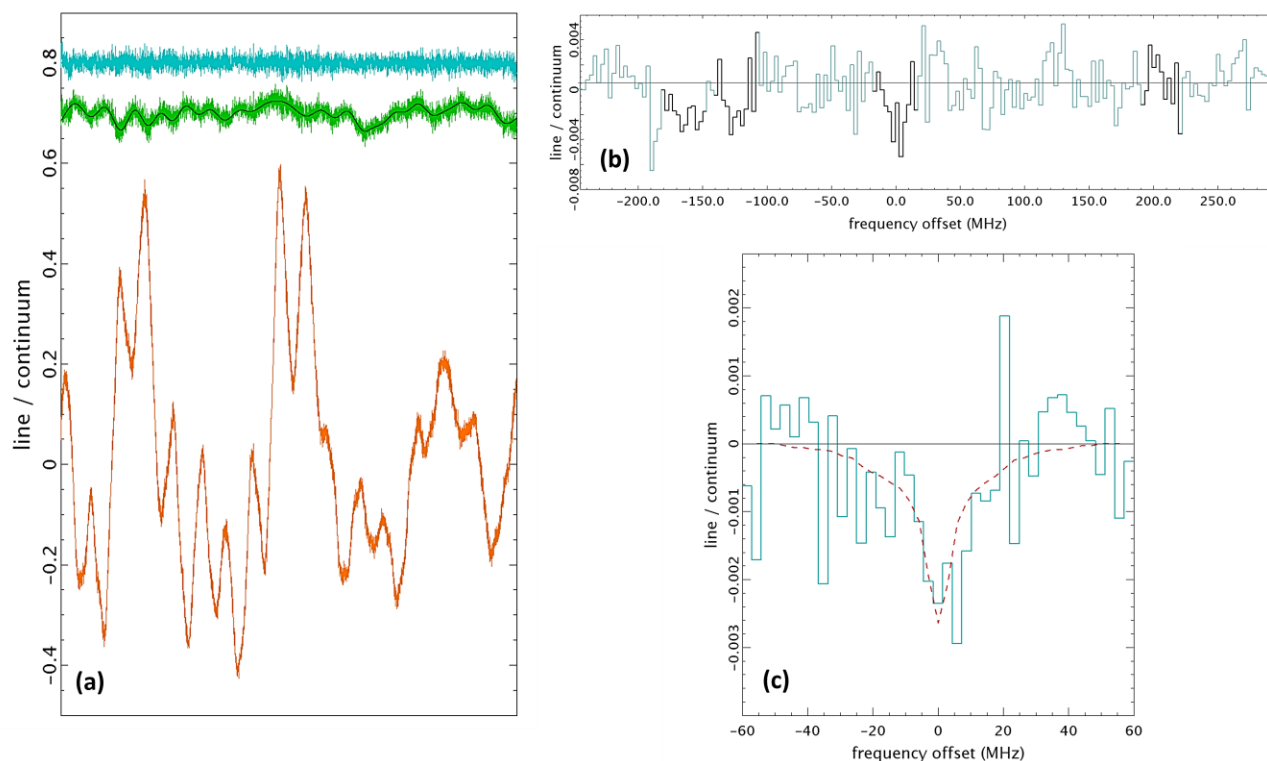
126

127   3 Results

128   Three of the four expected PH_3 J=4-3 components are visible when all the observations
129   are co-added (Figure 1b), while only one component was apparent in Figure 3 of C22. In Figure
130   1c, our final stacked spectrum indicates an overall 5.7σ detection of PH_3 J=4-3, when integrated
131   over ±17 MHz (the masked region). This confidence level changes marginally (by ±0.4σ) if a
132   different range of spectral pixels is used to calculate the zero-level.

133   We then modelled our net spectrum using the same online tool as C22. We ran a model
134   for 1 ppb of phosphine and scaled it linearly for different abundances, and then calculated a
135   reduced-chi-squared statistic to assess goodness of fit. Figure 1c illustrates that 3 ppb of PH_3
136   provides a good match to the observed line, with χ^2_r of 0.75. (The uncertainties used in χ^2_r were
137   generated per spectral pixel from the internal data dispersion.) We consider that fitting the
138   stacked spectrum better mitigates against artefacts, compared to matching the individual
139   components (Figure 1b) against the model. For example, inserting a single positive "spike" near
140   the highest-frequency component (Figure 1b) was found to significantly reduce the inferred PH_3
141   abundance, as the χ^2_r test attempts to minimise the discrepancy of positive data against a
142   negative model-line. As a positive feature does appears here in this part of C22's spectrum (their
143   Figure 3), this could have driven their upper limit down to the ≤0.8 ppb they obtain. However,
144   we do not rule out an abundance as low as 0.8 ppb, which is at the lower 99% confidence-bound
145   in our χ^2_r tests. We also note that C22 estimated ~2.3 ppb from their PH_3 J=2-1 spectrum, albeit
146   at only +1.5σ confidence, and this estimate is compatible with our 3 ppb result.

147   C22 (Figure S4) find that altitudes around ~80 km are the best-sampled at the PH_3 J=4-3
148   line-frequency; our recovery here is consistent with the model that we both use, and with the
149   predicted short lifetime of PH_3 above ~80 km (Bains, Petkowski, Seager, et al., 2021). Altitudes
150   are however uncertain because the PH_3-CO_2 pressure-broadening coefficient has not been
151   experimentally verified. We also note that all GHz/THz data are limited by the spectral span that
152   can be recovered. Here, any absorption wider than ~200 MHz leads to merged PH_3 J=4-3
153   components, and so any phosphine signatures below ~70 km (roughly cloud-top level) are lost.

154   *Figure 1. The process of extracting the PH_3 J=4-3 signal from the GREAT data is illustrated.*
155   *Panel (a) illustrates processing of 6000 spectral pixels from observation #040093 from the first*
156   *SOFIA flight (the reference spectral pixel is mid-band; vertical offsets are for clarity only). The*
157   *lower orange histogram is from a standard On-Off processing; the middle green histogram is the*
158   *result after applying the Eq.[1] step, and is overlaid with the Fourier-derived trend (black*
159   *curve); the top blue histogram is the flattened output after subtracting this trend. In panel (b),*

160   *the blue histogram is the unweighted average of all six observations made over the three flights,*
161   *with the sections containing the four PH₃ J=4-3 components highlighted in black. The spectral*
162   *pixels are binned in groups of 12 in (b) and (c) to improve clarity, and the thin black lines show*
163   *the zero-level correction made between (b) and (c). Panel (c) shows (blue histogram) the*
164   *unweighted stack of all 24 spectral sections containing PH₃ J=4-3 features. The dashed red*
165   *curve is a model for 3 ppb of phosphine, generated vis the Planetary Spectrum Generator (PSG)*
166   *(https://psg.gsfc.nasa.gov/index.php) following C22. The four line-components from the PSG*
167   *model were masked and stacked similarly to the data.*
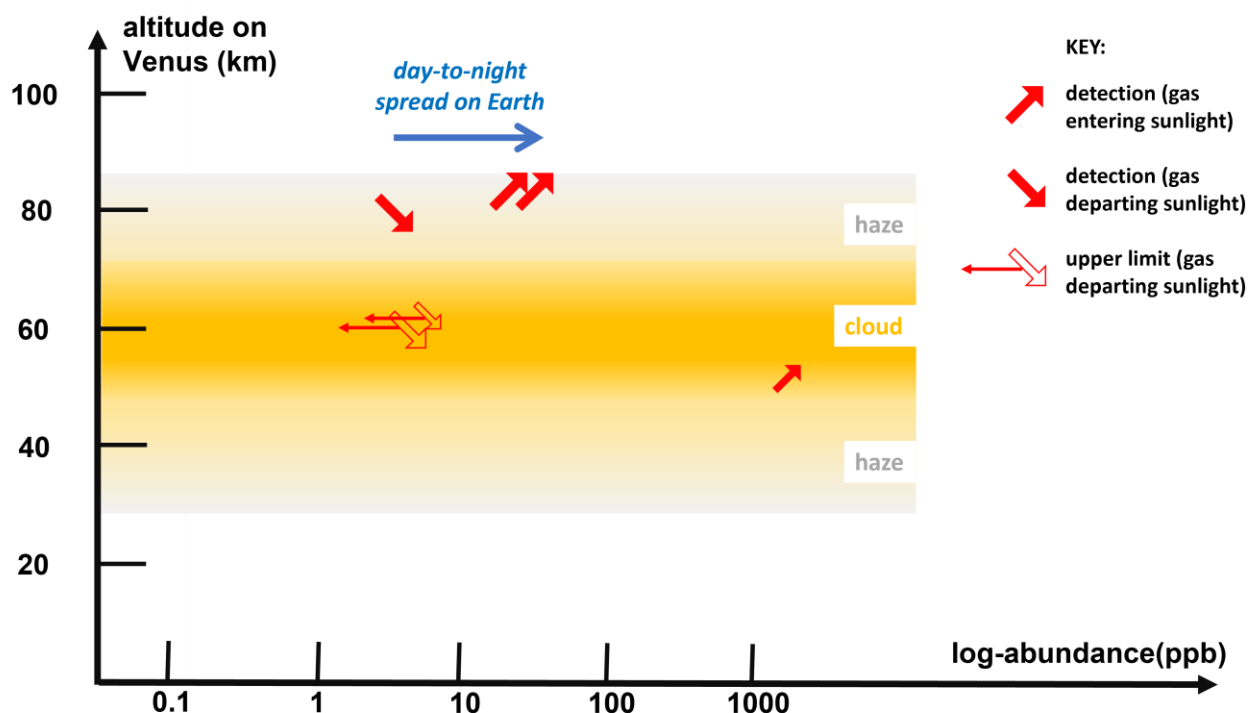

4 Discussion

169   Debates continue about the best methods to acquire and process deep GHz/THz spectra
170 of Venus. These observations are very challenging in dynamic range, as Venus is so bright,
171 revealing "ripples" in spectral baselines that are not evident in more typical telescope usage.
172 Depending on preferred approaches, different authors argue for between zero and three published
173 detections of rotational (J) transitions of PH₃.

174   We can compare results from the data discussed here with the outcomes of other searches
175 for phosphine at Venus, and assess whether this results in a plausible altitude profile of the
176 molecule (Figure 2). The trend found by connecting the results from six searches for phosphine
177 appears as an upwards decline that then reverses, i.e. PH₃ that is depleted somewhere between
178 ~50 km and ~80 km. This is hard to explain in the absence of a chemical route to reform the
179 molecules, or a new mesospheric source. The order-of-magnitude contrast between some of the
180 candidate detections and the upper limits has led to doubts over the presence of phosphine.

181   However, we noticed that this divide is also between observations made when the
182   'morning' versus the 'evening' sides of Venus' atmosphere were targeted – and this is relevant in
183   gas-mixing processes (e.g. (Lefèvre et al., 2022)). Where the gas observed on Venus has
184   travelled through sunlight and is descending towards the night-side of the planet, we detect at
185   most the ~3 ppb of phosphine estimated here. In contrast, where gas is rising into sunlight, we
186   observe ≥ ~20 ppb of PH_3. Hence photolysis – similar to the observed destruction of terrestrial
187   phosphine by sunlight (Sousa-Silva et al., 2020) – could explain the split between high and low
188   phosphine abundances observed on Venus. It is striking that this evening/morning difference is
189   comparable in magnitude to the factor ~10 difference over night/day for phosphine in the Earth's
190   atmosphere (Glindemann et al. 1996; noting absolute abundances are much lower on Earth).

191   *Figure 2. The trend of phosphine abundances by altitude is sketched. Symbols indicate candidate*
192   *detections plus best upper limits for phosphine abundances. Rising arrows indicate observations*
193   *made where the super-rotating atmosphere was rising into sunlight and falling arrows indicate*
194   *observations made where the atmosphere was descending towards the nightside (see key). Large*
195   *and small symbols indicate that a large fraction of the planet area was observed, or that a small*
196   *region was sampled, respectively. Abundance estimates are, from top: ~20, 25 ppb from J=1-0*
197   *data (via (Greaves et al., 2022) and with altitude proposed by C22); 3 ppb from J=4-3 data (this*
198   *work; beam centred on the evening side); <7 ppb at 62 km from 4 µm spectra (Trompet et al.,*
199   *2020: low-latitude data to best match whole-planet studies); <5 ppb at 60 km from 10 µm*
200   *spectra (Encrenaz et al., 2020: latitudes within ±50º); ~2 ppm at 51 km from Pioneer-Venus in-*
201   *situ sampling during descent (Mogul, Limaye, Way, et al., 2021). The blue arrow indicates the*
202   *ten-fold increase of terrestrial phosphine from day to night (Glindemann et al. 1996) – note the*
203   *arrow's plotted position is arbitrary; Earth hosts much lower PH_3 than Venus.*

204

5 Conclusions

The question regarding phosphine in Venus' atmosphere is likely to be debated for some time. A further *JCMT* survey is ongoing (see https://www.eaobservatory.org/jcmt/science/large-programs/jcmt-venus-monitoring-phosphine-and-other-molecules-in-venuss-atmosphere/; PI. D. Clements) and is producing open-source data that should yield more definitive answers – in particular, that team is now processing broadband spectra that can sample the cloud decks. The most direct answer regarding phosphine could come from new in-situ sampling, potentially with the addition of one laser channel to the Venus Tunable Laser Spectrometer (VTLS) instrument on-board the *DAVINCI* descent probe (Garvin et al. 2022).

The origins of any phosphine present are also debated, and most scenarios are hard to test for lack of some contextual data. For example, it seems only extraordinary volcanic activity could make ~ppb-level phosphine (Bains et al., 2022) but vulcanism on Venus is not well understood. In some new avenues, (Ferus et al., 2022) discuss abiotic routes to phosphine involving redox disequlibrium, while others (Bains, Petkowski, Rimmer, et al., 2021; Mogul, Limaye, Lee, et al., 2021) explore phototrophic life and the habitability of the clouds. We conclude that establishing an improved PH_3 altitude-profile is worthwhile to test these new models of origins.

Open Research

The SOFIA Level 1 data are available under project id 75_0059_1 through the public data archive at https://irsa.ipac.caltech.edu/applications/sofia. The custom software to generate the data shown in the figures is supplied at https://zenodo.org/record/7692288#.ZAC363bP3IU. The script requires the UK-Starlink software (Currie et al., 2014) which is currently supported by the East Asian Observatory and available at https://starlink.eao.hawaii.edu/starlink/2021ADownload.

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