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Neptune's carbon monoxide profile and phosphine upper limits from Herschel/SPIRE*: Implications for interior structure and formation

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Abstract

On Neptune, carbon monoxide and phosphine are disequilibrium species, and their abundance profiles can provide insights into interior processes and the external space environment. Here we use Herschel/SPIRE (Spectral and Photometric Imaging REceiver) observations from 14.9–51.5 cm⁻¹ to obtain abundances from multiple CO and PH₃ spectral features. For CO, we find that nine CO bands can be simultaneously fitted using a step profile with a 0.22 ppm tropospheric abundance, a 1.03 ppm stratospheric abundance, and a step transition pressure of 0.11 bar near the tropopause. This is in broad agreement with previous studies. However, we also find that the CO spectral features could be fitted, to well within measurement errors, with a profile that contains no tropospheric CO for pressure levels deeper than 0.5 bar,

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which is our preferred interpretation. This differs from previous studies that have assumed CO is well mixed throughout the troposphere, which would require an internal CO source to explain and a high O/H enrichment. Our interpretation removes the requirement for extreme interior O/H enrichment in thermochemical models and can finally reconcile D/H and CO measurements. If true, the lack of lower tropospheric CO would imply a decrease in Neptune's interior water content, favouring a silicate-rich instead of an ice-rich interior. This would be consistent with a protoplanetary ice source with a similar D/H ratio to the current solar system comet population. The upper tropospheric and stratospheric CO at pressures less than 0.5 bar could then be entirely externally sourced from a giant impact as suggested by Lellouch et al. (2005). We also derive a $3-\sigma$ upper limit for PH₃ of 1.1 ppb at 0.4–0.8 bar. This is the most stringent upper limit to-date and is entirely consistent with predictions from a simple photochemical model.

Keywords: Neptune, Atmosphere, Interior, Composition, sub-millimetre

1 1. Introduction

Observations of bulk density, gravity, and moment of inertia show that Neptune and Uranus are significantly enriched in heavy elements compared to Jupiter, Saturn, and the solar composition (Hubbard et al., 1995; Podolak et al., 1995; Owen and Encrenaz, 2006; Irwin, 2009; Helled et al., 2011; Nettelmann et al., 2013). Their interiors are expected to contain a significant proportion of ices due to their formation beyond the ice line in the solar nebula, where cold temperatures permitted formation and subsequent ingestion of icy planetesimals. However, the fraction of rock to ice and the nature of

the original planetesimals are currently not well understood and ice-rich and 10 rock-rich interior models are both compatible with the available bulk den-11 sity, gravity, and moment of inertia observations (Podolak et al., 1995; Helled 12 et al., 2011; Nettelmann et al., 2013). The different formation scenarios and 13 internal structures are consistent with subsets of available spectroscopic ob-14 servations and important discrepancies remain, particularly when trying to 15 reconcile D/H and O/H measurements (Feuchtgruber et al., 2013). Here we 16 consider two of Neptune's disequilibrium species, carbon monoxide (CO) and 17 phosphine (PH_3) , which have the potential to further constrain Neptune's in-18 terior composition, formation, and external planetary environment. 19

CO has been observed on both Uranus (Encrenaz et al., 1996; Cavalié 20 et al., 2014) and Neptune (Marten et al., 1993; Guilloteau et al., 1993; Naylor 21 et al., 1994; Courtin et al., 1996; Lellouch et al., 2005; Marten et al., 2005; 22 Hesman et al., 2007; Lellouch et al., 2010; Fletcher et al., 2010; Luszcz-Cook 23 and de Pater, 2013). On Uranus CO has a stratospheric abundance of 7.1-24 9.0 ppb (Cavalié et al., 2014) and a 3- σ tropospheric upper limit of 2.1 ppb 25 for pressures around 0.1–0.2 bar (Teanby and Irwin, 2013). On Neptune CO 26 is much more abundant, with recent studies suggesting a step-type profile 27 with 1-3 ppm in the stratosphere and up to 0.5 ppm in the troposphere 28 (summarised in Table 1). 29

The CO vertical gradient can be used to determine whether its source is mixing from the deep interior or external supply from comets, micrometeorites, or interplanetary dust. However, there is some disagreement in the published abundances (Table 1). The most recent study by Luszcz-Cook and de Pater (2013) found 1–2 ppm in the stratosphere, 0–0.3 ppm in the

troposphere, and a transition pressure of ~ 0.1 bar (close to the tropopause). 35 This is broadly consistent with Lellouch et al. (2005)'s stratospheric determi-36 nation of 1 ppm, but lower than their value of 0.5 ppm for the troposphere. 37 Conversely, Hesman et al. (2007)'s tropospheric value is consistent with Lel-38 louch et al. (2005)'s, but has a stratospheric abundance 2–3 times higher. 39 There is also considerable uncertainty on the pressure of the transition in 40 the step profiles used in all three studies. The discrepancy of these results 41 could be partly due to the difficulty of observing CO from ground-based tele-42 scopes. The wide CO line wings that probe the troposphere cannot usually 43 be covered with a single observation due to instrument bandwidth limita-44 tions. Therefore, either only the central emission core is observed to obtain a 45 stratospheric abundance, or multiple observations with different local oscil-46 lator tunings must be stitched together. Such observations require baseline 47 matching, which introduces extra uncertainty as observations must be taken 48 at different times with different sky and instrument background levels. There 40 are also differences in the radiative transfer modelling approaches, in particu-50 lar the assumed temperature profile. Therefore, new space-based constraints 51 would be extremely valuable. 52

⁵³ CO is more thermodynamically stable at the higher temperatures of ⁵⁴ Neptune's deep atmosphere ($\gtrsim 1000$ K, $\gtrsim 5000$ bar) and becomes less stable ⁵⁵ (and much less abundant) in comparison to the thermochemical-equilibrium ⁵⁶ favoured molecules methane and water in the colder outer region of Neptune's ⁵⁷ hydrogen-rich atmosphere (Lodders and Fegley, 1994). However, CO de-⁵⁸ struction reactions have a strong temperature dependence and rapid vertical ⁵⁹ mixing can advect CO to pressure levels where the CO destruction timescale

is longer than the mixing timescale, effectively quenching the CO destruction 60 and allowing significant CO to mix into the troposphere and stratosphere. 61 On Neptune this quenching is expected to occur at 2000–10000 bars and 62 850–1100 K depending on the strength of vertical mixing (Luszcz-Cook and 63 de Pater, 2013; Cavalié et al., 2017). Such rapid mixing should result in a 64 uniform CO mixing ratio in the troposphere and stratosphere. For Neptune, 65 plausible estimates of the temperature profile and vertical mixing suggest 66 that O/H must be enriched by at least 280 times relative to solar compo-67 sition in order to reproduce the observed ~ 0.1 ppm CO in the troposphere 68 (Luszcz-Cook and de Pater, 2013; Cavalié et al., 2017). However, such a large 69 enrichment of O/H is not compatible with D/H measurements, which sug-70 gest more modest O/H enrichments of \sim 50–150 (Feuchtgruber et al., 2013) 71 if Neptune's internal water was sourced from protoplanetary ices with D/H 72 comparable to present day comets. CO has not yet been detected in Uranus' 73 troposphere (Teanby and Irwin, 2013), which could be due to less vigorous 74 mixing. 75

The excess of CO in both Neptune and Uranus' stratospheres compared to 76 their tropospheres suggests a significant external CO source for both plan-77 ets (Lellouch et al., 2005; Cavalié et al., 2013). Estimates of the external 78 flux required to explain observed H_2O and CO_2 abundances in Uranus and 79 Neptune's stratosphere were made by Feuchtgruber et al. (1997), suggesting 80 H_2O external fluxes of $0.6-1.6 \times 10^5$ molecules cm⁻²s⁻¹ for Uranus and 1.2-81 150×10^5 molecules cm⁻²s⁻¹ for Neptune. These inferred H₂O influx rates 82 are broadly consistent with those expected from incoming interplanetary 83 dust grains (Poppe, 2016). However, the predicted dust flux for Neptune 84

is roughly two orders of magnitude too small to explain the very large ob-85 served CO abundance in Neptune's stratosphere, even if the oxygen from the 86 dust grains were efficiently converted to CO (see Poppe, 2016; Moses and 87 Poppe, 2017). The large concentration of CO in Neptune's stratosphere led 88 Lellouch et al. (2005) to suggest that the source of Neptune's CO could be 89 a large cometary impact that occurred ~ 200 years ago. This hypothesis is 90 supported by the recent detection of CS by Moreno et al. (2017), an impact 91 product that was also detected after the Shoemaker-Levy 9 impact on Jupiter 92 (Moreno et al., 2003). Moreno et al. (2017) suggest a slightly larger 4 km 93 diameter comet impacting ~ 1000 years ago. 94

Further constraints on Neptune's atmosphere can be obtained by study-95 ing another disequilibrium species, phosphine (PH_3) , which is only stable in 96 the deep atmosphere, but could theoretically be transported to higher at-97 mospheric levels by rapid vertical mixing as on Jupiter and Saturn (Irwin 98 et al., 2004; Fletcher et al., 2009). There is currently considerable uncergc tainty about Neptune's phosphorous enrichment, but a P/H enrichment of 100 50 times solar, similar to that inferred for C/H from methane measurements 101 (Baines et al., 1995), is considered reasonable. However, observing PH_3 is 102 challenging as it condenses in Neptune's cold troposphere for pressures less 103 than ~ 1 bar. Also, PH₃ is photodissociated by solar UV photons in the upper 104 troposphere of giant planets (Kaye and Strobel, 1984; Moses, 2000), leading 105 to additional depletion in the upper troposphere. This should result in sig-106 nificant PH₃ abundance only for pressures greater than $\sim 1-2$ bar, producing 107 spectral features that are wide due to pressure broadening and with negligi-108 ble central emission core due to a lack of significant stratospheric abundance. 109

Nevertheless, an attempt was made to observe the 267 GHz (8.91 cm⁻¹) PH_3 110 feature by Encrenaz et al. (1996) who determined that a 2 ppm deep PH_3 111 abundance ($\sim 3x$ solar P/H) coupled with super-saturation by a factor of 100 112 above the condensation level was inconsistent with observations from the 113 Caltech Submillimeter Observatory. More recently Moreno et al. (2009) de-114 termined a more stringent upper limit of 0.1 times the solar P/H value using 115 the same PH₃ feature observed with the IRAM 30 m telescope, corresponding 116 to an upper troposphere abundance upper limit of ~ 60 ppb (assuming a solar 117 abundance of $P/H=2.81\times10^{-7}$ from Lodders (2010)). Unfortunately, these 118 upper limits are not sufficient to significantly inform photochemical models. 119 Here we use Herschel/SPIRE data to further constrain Neptune's CO and 120 PH_3 profiles. SPIRE is ideally suited to studying these gases as it covers a 121 wide spectral range containing multiple CO and PH₃ features. This allows 122 the line wings to be measured in a single measurement from a space-based 123 platform with no atmospheric interference. This will permit the most sensi-124 tive search of PH₃ to date, more robust constraints on Neptune's CO profile, 125 and insights into Neptune's interior and formation. 126

127 2. Observations

Observations were taken with the SPIRE instrument (Griffin et al., 2010; Swinyard et al., 2010) on board the Herschel Space Observatory (Pilbratt et al., 2010). SPIRE is a Fourier transform spectrometer comprising a long-wave spectrometer (SLW) covering wavenumbers 14.9–33.0 cm⁻¹ (671– 303 μ m) and a short-wave spectrometer (SSW) covering 31.9–51.5 cm⁻¹ (313– 194 μ m). The SPIRE spectrometers have hexagonal arrays of circular pixels; ¹³⁴ 37 pixels for the SLW and 19 pixels for the SSW. The unapodised spec-¹³⁵ tral resolution can be set to low resolution (0.83 cm⁻¹), medium resolution ¹³⁶ (0.24 cm⁻¹), and high resolution (0.04 cm⁻¹). Herschel's 3.5 m diameter pri-¹³⁷ mary mirror combined with the pixel array footprints results in pixel beams ¹³⁸ with a field-of-view of 17–42", which is large compared to Neptune's ~2.3" ¹³⁹ projected diameter, so observations are disc-averaged.

Herschel operated from 2009 until 2013, when it ran out of coolant, and 140 all data are now archived. Uranus was used as the primary flux calibrator 141 for the SPIRE spectrometer (Swinyard et al., 2014), but Neptune was also 142 regularly observed for cross-calibration with the SPIRE photometer observa-143 tions (Swinyard et al., 2014; Hopwood et al., 2015). The pipeline radiance 144 calibration used a Uranus reference spectrum combined with darks taken 145 on a relatively empty area of sky, which were used to remove the instru-146 ment self-emission and characterise the instrument (Swinyard et al., 2010, 147 2014; Hopwood et al., 2015). We queried the Herschel Science Archive for all 148 science and calibration observations of Neptune taken with SPIRE in high 140 resolution spectrometer mode so that the CO features could be resolved. The 150 Level 2 unapodised calibrated radiances were extracted and convolved with 151 a Hamming function to give an apodised spectral resolution of 0.07373 cm^{-1} 152 (full-width half-maximum) and a sample spacing of 0.01 cm^{-1} . The CO ab-153 sorption features had widths of $\sim 0.5 \text{ cm}^{-1}$ and the widths of the central 154 emission cores were limited by SPIRE's spectral resolution. The Neptune-155 centred pixel from each observation was extracted to give a disc-average 156 spectrum and associated pipeline uncertainties. After quality control of the 157 observations to reject spectra with excessive noise or anomalously high/low 158

radiances, 17 Neptune observations remained with integration times from
823–13762 s. Observations are summarised in Table 2 and Figure 1a and b.
Total integration time on Neptune was 32687 s (9 hrs 4 mins 47 s), with all
observations taking place during 2009 and 2010.

Prior to further analysis, the 17 individual spectra were combined into 163 a single high signal-to-noise disc-average Neptune spectrum. To account for 164 distance variations and slight calibration differences between observations the 165 weighted average continuum radiances at 20.5-21.5 cm⁻¹ and 39.5-40.5 cm⁻¹ 166 were calculated and individual spectra were rescaled to match the overall av-167 erage. This rescaling was required to allow representative variances on the 168 combined spectrum to be calculated, but did not affect the overall mean ra-169 diance. All 17 observations were then combined into an average spectrum 170 using the error weighted mean (Bevington and Robinson, 1992). To deter-171 mine the uncertainties on the combined spectrum we calculated both the 172 error-weighted variance of the 17 spectra and the theoretical error weighted 173 variance based on the pipeline uncertainties of each spectrum; whichever was 174 the largest was used for the uncertainty. Overall uncertainties (standard er-175 ror) per spectral element were ~ 0.2 Jy for the SLW and ~ 0.3 Jy for the SSW. 176 The combined average SLW and SSW spectra are shown in Figure 1c and d. 177 CO absorption and emission features are clearly visible. However, the spectra 178 still contain slight continuum ripples as noted in previous studies (Teanby 179 and Irwin, 2013; Teanby et al., 2013). Removal of these ripples required 180 comparison to a synthetic spectrum and is discussed in the next section. 181

182 3. Spectral modelling

To fit the observed SPIRE spectrum and retrieve the composition of CO 183 and PH_3 we used the NEMESIS radiative transfer code (Irwin et al., 2008), 184 which we have previously used to analyse SPIRE spectra for Uranus (Teanby 185 and Irwin, 2013) and Titan (Teanby et al., 2013), in addition to extensive 186 use for analysis of Neptune's near-IR spectra (Irwin et al., 2011, 2014, 2016). 187 Our Neptune reference atmosphere had a nominal temperature profile 188 based on Voyager 2 radio occultation analysis by Lindal (1992) for pressures 189 greater than 15 mbar, AKARI spectroscopic analysis by Fletcher et al. (2010) 190 for pressures less than 10 mbar, and a linear interpolation (in log pressure) 191 in between. Temperature was gridded onto 71 levels between 0.6 μ bar and 192 6 bar on a regular logarithmic grid. There is some uncertainty regarding 193 Neptune's tropospheric temperature structure, with differences up to 5 K 194 reported in the literature (for example, see discussion in Hesman et al., 2007; 195 Fletcher et al., 2010; Luszcz-Cook and de Pater, 2013). Therefore, we also 196 consider "Hot" and "Cold" profiles, which differ from the nominal profile 197 by ± 5 K (Figure 2). For the baseline atmospheric composition we assumed 198 a He/H_2 ratio of 0.15 (by volume) and a volume mixing ratio (VMR) of 199 0.003 for N₂ (Conrath et al., 1993). The CH₄ volume mixing ratio was set 200 to a deep value of 0.02, followed the saturation pressure vapour curve in the 201 upper troposphere, and had an abundance of 1.15×10^{-3} in the stratosphere 202 (Lellouch et al., 2015). There is some uncertainty on Neptune's deep CH_4 203 abundance (see discussion in Irwin et al., 2014), but the effect on the SPIRE 204 spectrum is minimal. HCN only has minor spectral features in this range, but 205 we include the abundance profile of Marten et al. (2005) for completeness. 206

²⁰⁷ Spectroscopic parameters were the same as those used in Teanby and Irwin²⁰⁸ (2013).

Synthetic spectra were generated using the correlated-k approximation 209 for computational efficiency (Goody and Yung, 1989; Lacis and Oinas, 1991; 210 Irwin et al., 2008). The k-tables included the Hamming instrument function 211 with a FWHM of 0.07373 cm^{-1} . To simulate the disc-averaged spectrum we 212 used the field-of-view averaging technique detailed in Teanby et al. (2013) 213 with 33 field-of-view points; 20 on Neptune's disc and 13 covering the limb. 214 This accounted for limb brightening and limb darkening effects and was suffi-215 cient to reduce the disc-averaged systematic modelling errors, due to emission 216 angle variation across Neptune's disc, to at least five times below the level 217 of the observation uncertainties. 218

Observed spectra were corrected for minor continuum ripples by compar-219 ison with synthetic spectra generated using the reference atmosphere. First, 220 synthetic spectra were created for an atmosphere containing no CO or PH₃. 221 Second, synthetic spectra were created for an atmosphere with the nominal 222 CO profile from Luszcz-Cook and de Pater (2013) and a PH_3 profile with a 223 deep abundance of 1 ppb with the saturation vapour pressure law applied. 224 Third, the difference between spectra with and without CO/PH_3 was used 225 to create a continuum mask where the difference was less than 0.05 Jy, at 226 least a factor of four below the observational uncertainties. Fourth, the mask 227 was applied to the observations to remove spectral regions surrounding the 228 CO and PH_3 line positions, leaving only the continuum points. Fifth, the 229 ratio between masked observation and synthetic was calculated and a smooth 230 cubic b-spline curve fitted using a knot spacing of 3 cm^{-1} for the SLW and 231

 1.5 cm^{-1} for the SSW (Teanby, 2007). These knot spacings were large enough 232 to only remove large scale ripples and did not affect the spectral features. 233 The resulting correction factor $f_{\rm corr}$ is shown in Figure 1e and f and sug-234 gests continuum ripples of order $\pm 1\%$. Finally, the fitted smooth correction 235 factor curve was applied to the observations to give the corrected SLW and 236 SSW spectra shown in Figure 1g and h. This procedure was repeated for 237 the nominal, hot, and cold temperature profiles. Note that a CO feature at 238 15.38 cm^{-1} could not be analysed as it was at the low wavenumber edge of 239 the SLW, so the surrounding continuum level could not be reliably corrected. 240 To determine the information content of our observed spectra we calcu-241 lated the contribution functions, defined as the change in irradiance as a 242 function of gas abundance at each pressure level. The contribution functions 243 for each of the CO bands are shown in Figure 3 assuming the SPIRE spec-244 tral resolution, the nominal temperature profile, and a 0.1 ppm uniform CO 245 profile. In addition to the SPIRE CO bands, we also calculated the contribu-246 tion functions for the lower frequency CO (1-0), (2-1), and (3-2) transitions 247 analysed in previous studies. Therefore, for this calculation the temperature 248 profile was extended to 60 bar using the dry adiabatic lapse profile rate from 249 Luszcz-Cook and de Pater (2013) so that contributions from the troposphere 250 for the lowest frequency CO lines were fully determined. Note that con-251 tribution functions for the CO (1-0), (2-1), and (3-2) transitions are also 252 calculated at the SPIRE spectral resolution, so are representative of the line 253 wings only, not the central emission cores, which can sound to much lower 254 pressure at high spectral resolution. 255



Figure 3 shows that there are two main pressure ranges where the SPIRE

spectra have information on CO abundance: $\sim 1.0-0.1$ bar from the wide 257 tropospheric absorption wings of the CO lines, which are most prominent for 258 the SLW observations; and $\sim 0.001-0.01$ bar from the narrow stratospheric 259 emission cores of the CO lines, which are most prominent in the SSW ob-260 servations. The difference between SLW and SSW CO features is caused by 261 increases in CO line strength and overall atmospheric opacity with increas-262 ing wavenumber across the SPIRE range. There is little or no information 263 in the 0.01–0.1 bar region, which explains the large uncertainty in previ-264 ous determinations of the transition pressure in step-type CO profiles (see 265 e.g. Luszcz-Cook and de Pater, 2013, and Table 1). The SPIRE data are 266 not sensitive to the deep abundance (pressures greater than 1.0 bar), but 267 have excellent coverage of the upper troposphere and stratosphere. The low-268 est frequency bands, including the CO (1-0), (2-1), and (3-2) transitions 269 analysed in previous studies (Table 1), are sensitive to slightly lower deeper 270 levels than our data. Given the limited bandwidth of previous observations 271 (e.g. Luszcz-Cook and de Pater, 2013) Figure 3 shows the maximum pressure 272 probed is ~ 3 bar for the CO (1–0) transition, assuming a 0.1 ppm uniform 273 CO abundance. 274

The PH₃ contribution functions are shown in Figure 4 assuming a nominal deep abundance of 1 ppb and an abundance following the saturation vapour pressure curve in the troposphere, with a maximum stratospheric abundance set to that at the tropopause cold trap. Saturation vapour pressure temperature dependence was based on a fit to the low temperature vapour pressures in Lide (1995), giving the saturation vapour pressure in bars $P_{\text{SVP}}(T) = \exp(a + b/T + cT)$, where a = 11.4600, b = -1974.44 K, and

c = -0.00435846 K⁻¹. The pressure level where PH₃ condenses is a strong 282 function of temperature, anywhere from 0.1-1 bar, which gives very different 283 contribution functions for the nominal, hot, and cold temperature profiles. 284 However, there is very limited information for pressures greater than 1 bar 285 in these data. There are four PH_3 spectral features covered by SPIRE (cen-286 tred on 17.81, 26.70, 35.59, and 44.46 cm^{-1}), but the features at 17.81 cm^{-1} 287 (SLW) and 44.46 $\rm cm^{-1}$ (SSW) are the most favourable for a detection as 288 they are well separated from the CO features. The optimal band for search-289 ing for PH₃ depends on the temperature profile and abundance profile, with 290 the SLW being optimal for the cold and nominal profiles, and the SSW being 291 optimal for the hot profile. 292

Spectra were fitted using NEMESIS' iterative non-linear retrieval scheme (Irwin et al., 2008), which adjusts the composition profiles to minimise the misfit between modelled spectrum and observation. Simple parameterised profiles were used for both CO and PH₃, so it was not necessary to impose apriori constraints on the retrieval scheme.

For fitting the CO spectral features we used three profile types: (1) a 298 three parameter simple step profile, defined by a uniform deep abundance 299 v_1 , a uniform stratospheric abundance v_2 , and a transition pressure p_1 ; (2) 300 a four parameter gradient profile, defined by a uniform deep abundance v_1 , 301 a uniform stratospheric abundance v_2 , and two pressures p_1 and p_2 , which 302 defined a transitional linear gradient region (in log pressure-abundance) from 303 v_1 to v_2 ; and (3) a four parameter external gradient profile with zero deep 304 abundance for pressures greater than p_1 , a linear transition region (in log 305 pressure-abundance) defined by two pressure-abundance pairs (p_1, v_1) and 306

 (p_2, v_2) , and uniform abundance v_2 at pressures less than p_2 . The step profile (1) has been used extensively in the literature to represent a combination of internal and external sources. The gradient profile (2) expands on the step profile slightly by allowing a finite mixing region. The external gradient profile (3) represents the case where there is no significant internal CO source and all tropospheric CO is mixed from above.

For PH₃, we used a single parameter profile based on a uniform deep abundance modified by the saturation vapour pressure and with the stratospheric abundance set to that at the troposphere cold trap.

316 4. Results

All nine CO spectral features were fitted simultaneously for each of the three CO profile types and assuming a nominal, hot, or cold temperature profile. Table 3 summarises the fit parameters for each case. The quality of each fit was assessed using the χ^2 statistic:

$$\chi^2 = \sum_{i=1}^{N} \left[\frac{I_{\text{obs}}(\nu_i) - I_{\text{fit}}(\nu_i)}{\sigma(\nu_i)} \right]^2 \tag{1}$$

where ν_i is the wavenumber, $I_{obs}(\nu_i)$ is the measured spectral irradiance, $I_{fit}(\nu_i)$ is the best fitting synthetic spectrum, $\sigma(\nu_i)$ is the observational error, and N is the number of points in the spectrum. The reduced χ^2 defined by χ^2/N should be ~1 if the synthetic spectrum fits the data within error.

The nominal temperature profile combined with the step CO profile had the lowest χ^2/N (0.63), which we consider our best fitting model. Figure 5 shows the best fit to the nine CO features covered by SPIRE, assuming the nominal temperature profile and using the step profile with deep abundance

of 0.22 ppm, a stratospheric abundance of 1.03 ppm and a transition pressure 329 of 0.11 bar. The reduced χ^2 is less than 1, suggesting the data are well fitted 330 by this profile. In fact, the χ^2/N only exceeds 1 for the hot external gradient 331 profile, but for all other cases the fits can be considered adequate. While 332 the nominal temperature step profile technically provides the best fit to the 333 data, it cannot be statistically distinguished from the other temperature / CO 334 profile cases. Therefore, a wide range of profiles can fit these data, including 335 those with no internal CO source. The fitted profiles are shown in Figure 6a 336 and can all be considered plausible. The range of abundances obtained were 337 0-0.36 ppm for the deep volume mixing ratio and 0.80-1.55 ppm for the 338 stratospheric volume mixing ratio. The transition pressure range is not well 339 determined by these data, but falls between 0.11–0.75 bar. 340

None of the four phosphine spectral features covered by SPIRE were visible in the data. Therefore, we determined upper limits using a modified $\chi^2, \chi^2_r(v)$:

$$\chi_r^2(v) = \frac{\Delta\nu_{\rm obs}}{\Delta\nu_{\rm res}} \sum_{i=1}^N \left[\frac{I_{\rm obs}(\nu_i) - I_{\rm fit}(\nu_i, v)}{\sigma(\nu_i)} \right]^2 \tag{2}$$

where $\Delta \nu_{\rm obs}$ is the observation wavenumber spacing (0.01 cm⁻¹), $\Delta \nu_{\rm res}$ is the observation spectral resolution (0.07373 cm⁻¹), and v is the deep PH₃ volume mixing ratio. The factor $\Delta \nu_{\rm obs} / \Delta \nu_{\rm res}$ accounts for the spectral oversampling. The PH₃ profile is defined by one parameter so the 3- σ upper limit is defined as the abundance which increases χ_r^2 by +9 (Press et al., 1992).

Figure 7 shows the variation of $\Delta \chi_r^2 = \chi_r^2(v) - \chi_r^2(0)$ as a function of PH₃ abundance for the nominal, hot, and cold temperature profiles, along with the observed spectra and 3- σ upper limit synthetics. The SLW 17.81 cm⁻¹ and $_{352}$ SSW 44.46 cm⁻¹ bands were considered independently and the best upper $_{353}$ limits for each assumed temperature profile are summarised in Table 4. The $_{354}$ PH₃ upper limit for the nominal temperature profile was 1.10 ppb. Figure 6b $_{355}$ shows the upper limits and indicates the pressure range of the information $_{356}$ content.

357 5. Discussion

358 5.1. CO profile

Our retrieved CO abundances are broadly consistent with previous re-359 sults and agree with those presented by Luszcz-Cook and de Pater (2013) 360 (Table 1). The step profile results provide the most direct comparison as this 361 profile was also used in the previous studies (Lellouch et al., 2005; Hesman 362 et al., 2007; Lellouch et al., 2010; Luszcz-Cook and de Pater, 2013). Our 363 preferred deep abundance of 0.21–0.24 is most consistent with Luszcz-Cook 364 and de Pater (2013) and Hesman et al. (2007), whereas our stratospheric 365 abundance of 0.80-1.41 is most consistent with Luszcz-Cook and de Pater 366 (2013) and Lellouch et al. (2005). The advantage of our CO profiles over 367 previous studies is that the SPIRE data cover a much larger spectral range, 368 allowing the wide CO line wings and narrow emission cores to be measured 369 at the same time. We also fit nine CO bands simultaneously, providing an 370 extra consistency check compared to previous studies, which fitted between 371 one and three bands (Table 1). However, the SPIRE observations are at 372 relatively low spectral resolution compared to previous microwave studies, 373 so cannot fully resolve the emission cores, resulting in reduced sensitivity 374

at very low pressures (<0.1 mbar). Our observations are also at higher frequency compared to previous studies and do not sound below 1 bar.

One key feature of the SPIRE observations is they do not require significant CO for pressures greater than 0.5 bar, and CO is only required in the upper troposphere (0.1–0.5 bar) to adequately fit the data. This was also apparent in some of the solutions found by Luszcz-Cook and de Pater (2013), which included the J=1-0 line at 115.271 GHz, which has contribution functions that extend slightly deeper than our data (to \sim 3 bar assuming a uniform CO profile (Figure 3)).

Therefore, by combining our results with those of Luszcz-Cook and de Pater (2013) we infer that while some CO is required in the upper troposphere (0.1–0.5 bar) to fit the observations, it is not a requirement to have significant amounts at deeper pressures. This has important implications for the formation and composition of Neptune's deep atmosphere and could resolve some of the apparent discrepancies between CO and D/H measurements.

³⁹⁰ 5.2. Neptune's formation and internal structure

D/H in Neptune's atmosphere has been measured to be $4.1\pm0.4\times10^{-4}$ 391 from Herschel/PACS observations (Feuchtgruber et al., 2013). When this is 392 combined with water-rich interior models fitted to mass, moment of inertia, 393 and gravitational coefficients from the Voyager 2 flyby (Hubbard et al., 1995; 394 Podolak et al., 1995; Helled et al., 2011; Nettelmann et al., 2013), this value 395 of D/H suggests the icy planetesimals that formed Neptune had D/H ratios 396 of $5.1-7.7 \times 10^{-5}$. This is ~2-3 times less enriched that any known source 397 material in the present day solar system, with comets having a D/H ratio of 398 $\sim 1.5 - 3.0 \times 10^{-4}$ (Hartogh et al., 2011). 390

One potential solution proposed by Feuchtgruber et al. (2013) is to in-400 crease the silicate content of the pre-Neptune planetesimals to 68–86% rock 401 an 14-32% ice, i.e. similar to the estimated bulk rock fraction of Pluto (Si-402 monelli and Reynolds, 1989). This would give Neptune a more rock-rich 403 interior compared to more conventional water-rich internal models (Hubbard 404 et al., 1995) and would allow water to be sourced from ices with D/H in the 405 range of current comets $(1.5-3.0\times10^{-4})$. The reduced ice content then puts 406 the O/H enrichment contribution due to Neptune's water ice content to be 407 50–150 times solar (Feuchtgruber et al., 2013). This solution is appealing as 408 the resulting inferred rock content is also consistent with predictions from 409 the Solar Composition Icy Planetesimals (SCIPs) model of planet forma-410 tion (Owen and Encrenaz, 2006) and simple formation models using either 411 comet-like or clathrate-hydrate planetesimals (Ali-Dib and Lakhlani, 2018). 412 The inferred O/H enrichment is also comparable with C/H enrichment, which 413 is estimated at ~ 50 times solar based on methane measurements by Baines 414 et al. (1995). 415

However, a problem arises when comparing this potential formation and 416 interior model with thermochemical schemes for CO, which require highly 417 enriched O/H ratios of 280–650 times solar to allow sufficient mixing of CO 418 into the troposphere to explain the previously derived ~ 0.1 ppm abundances 419 (Lodders and Feglev, 1994; Luszcz-Cook and de Pater, 2013; Cavalié et al., 420 2017). For example, Cavalié et al. (2017) used a 1-D thermochemical kinetics 421 and transport model to predict the CO mixing ratio profile on Neptune for 422 different assumptions about the deep O/H abundance. They found that a 423 deep atmospheric O/H ratio of ~ 540 times the solar ratio was required to 424

explain a tropospheric CO mixing ratio of 0.2 ppm, if they assumed the fast 425 $CO \rightarrow CH_4$ chemical conversion scheme described in Venot et al. (2012). 426 However, as is discussed by Visscher et al. (2010), Moses et al. (2011), and 427 Moses (2014), the adoption of a very large rate coefficient for the reaction H 428 + CH₃OH \rightarrow CH₃ + H₂O in the Venot et al. (2012) scheme likely leads to 429 an overestimate in the required deep O/H abundance on the giant planets. 430 Using the Moses et al. (2011) rate coefficient for this reaction, Cavalié et al. 431 (2017) derive a deep O/H ratio of \sim 280 for Neptune in order to produce an 432 upper tropospheric CO mixing ratio of 0.2 ppm. This O/H value is still too 433 large to be compatible with the D/H ratio, as discussed above. 434

One potential solution to reconcile the CO and D/H measurements is to 435 form Neptune (and Uranus) on the CO ice line (Ali-Dib et al., 2014). In this 436 model CO pebbles are concentrated near the ice line due a combination of 437 rapid outward diffusion of CO gas and slow inward migration of pebbles due 438 to gas drag. If Neptune forms close to this ice line then the pre-Neptune 439 planetesimals are largely composed of CO instead of H_2O , resulting in the 440 bulk of Neptune's water originating from transformation of CO into H_2O 441 in the planets interior. This would then be compatible with a higher more 442 comet-like D/H ratio for the water-rich planetesimals contributing to Nep-443 tune's formation. 444

We propose an alternative and perhaps simpler solution – that there is in fact no significant tropospheric CO for pressures greater than ~ 0.5 bar – and that the majority of Neptune's observable CO is externally sourced. This hypothesis would be consistent with our observations and those of Luszcz-Cook and de Pater (2013) and relaxes the requirement for extreme O/H enrich-

ment in the deep interior. A requirement of this scenario is that the eddy 450 mixing coefficient K in the upper troposphere is reduced so that sufficient 451 CO from a comet impact can be maintained in the 0.1-0.5 bar pressure range 452 to fit the observations. The current estimate of $K=10^8 \text{ cm}^2 \text{s}^{-1}$ in the deep 453 troposphere is based on mixing length theory and internal heat flux (Moses, 454 1992; Moses et al., 1992) and in the absence of other constraints is generally 455 applied to the whole troposphere. However, this heat flux is effectively emit-456 ted from the radiative convective boundary, which is likely to be somewhat 457 below the tropopause. Comparing Neptune's observed brightness tempera-458 ture of 60 K near the $\sim 100 \ \mu m$ peak of its infrared emission (Burgdorf et al., 459 2003) to the temperature profiles in Figure 2 suggests a pressure level of 460 ~ 0.5 bar for this emission level. The lapse rate is also somewhat reduced in 461 the upper troposphere region, suggesting a more stable atmosphere close to 462 the trop pause than deeper in the atmosphere. Therefore, a reduced K in 463 the 0.1-0.5 bar range is plausible, and is also in line with requirements from 464 recent photochemical modelling (Moses et al., 2018). In this case the external 465 CO source would also still be compatible with the giant comet impact pro-466 posed by (Lellouch et al., 2005) and supported by (Moreno et al., 2017). The 467 silicate-rich Neptune suggested by Feuchtgruber et al. (2013) would then be 468 more compatible with the available observations than a water-rich Neptune. 469

470 5.3. Phosphorous chemistry

Our PH₃ upper limits of 0.192-5.52 ppb for 0.1-1.2 bar are one to two orders of magnitude more stringent that previous determinations (Encrenaz et al., 1996; Moreno et al., 2009) and show that PH₃ can be considered negligible in the upper troposphere and stratosphere. If the tropospheric ⁴⁷⁵ PH₃ profile is determined entirely by condensation of a deep abundance then ⁴⁷⁶ our upper limits are significantly sub-solar; corresponding to enrichments ⁴⁷⁷ of 3.93×10^{-4} – 1.13×10^{-2} times the Lodders (2010) solar values (Table 4). ⁴⁷⁸ This is at least three orders of magnitude below the expected ~50 times ⁴⁷⁹ enrichment expected from methane measurements and formation models, so ⁴⁸⁰ suggests other loss processes must be active.

To test whether the lack of PH_3 in this region of Neptune's atmosphere 481 has a photochemical origin, we developed a simple, global-average, one-482 dimensional model for tropospheric and stratospheric chemistry on Neptune 483 that includes PH₃ photochemistry, using the Caltech/JPL KINETICS code 484 (Allen et al., 1981; Yung et al., 1984). The model inputs are similar to those 485 discussed by Moses and Poppe (2017), but we add nitrogen and phospho-486 rous species and reactions based on the Jupiter and Saturn studies described 487 in Kaye and Strobel (1983b), Kaye and Strobel (1983a), Kaye and Strobel 488 (1984), Visscher et al. (2009), Moses et al. (2010), and Fletcher et al. (2018). 489 We assume that the PH_3 mixing ratio at the model lower boundary (8 bar) 490 is either 2.0×10^{-5} or 4.6×10^{-5} (~30x or ~70x solar, based on the protosolar 491 abundances of Lodders et al. (2009)). The eddy diffusion coefficient profile 492 adopted in the model is shown in Fig. 4 of Moses et al. (2018), and is based on 493 the hydrocarbon photochemical modelling discussed in Moses et al. (2005). 494 Although the eddy diffusion coefficients increase with increasing altitude in 495 the stratosphere of Neptune, Moses et al. (2005, 2018) find that the eddy 496 diffusion coefficient must be small (assumed $400 \text{ cm}^2 \text{s}^{-1}$) in the upper tropo-497 sphere and/or lower stratosphere of Neptune in order for the large observed 498 C_2H_6 abundance to be reproduced. This is consistent with the qualitative 499

radiative arguments given in Section 5.2. Our models include absorption
of solar radiation by atmospheric gases and multiple Rayleigh scattering by
gaseous H₂, He, and CH₄, but we do not consider aerosol extinction.

The results from this photochemical model indicate that PH₃ should be 503 confined to pressure greater than ~ 1 bar on Neptune (Figure 6b), consis-504 tent with our derived PH_3 upper limits. Phosphine is photolysed by solar 505 ultraviolet photons with wavelengths less than 230 nm. The main products 506 are $PH_2 + H$. The resulting atomic H can also react with PH_3 to produce 507 $PH_2 + H_2$, and two PH_2 radicals can combine to form P_2H_4 , which then con-508 denses (Ferris and Benson, 1981; Kaye and Strobel, 1984). In this way, the 509 PH₃ is relatively efficiently converted to diphosphine and other phosphorus-510 bearing aerosols in Neptune's troposphere. Solar photons with wavelengths 511 less than 230 nm do not penetrate past the $\sim 2-3$ bar level in Neptune's 512 troposphere, but multiple Rayleigh scattering allows significant photolysis of 513 PH_3 in the ~0.5–2.5 bar region, limiting the vertical extent of PH_3 . The low 514 eddy diffusion coefficient in this region prevents PH₃ from being transported 515 up faster than it can be destroyed by photolysis. This result could change if 516 the eddy diffusion coefficient in the upper troposphere were greater than we 517 have assumed or if aerosols (not included in the model) shield the PH₃ from 518 photolysis. Our strong upper limits on the PH_3 mixing ratio in the 0.5–1 bar 519 region will allow for useful constraints on the strength of atmospheric mixing 520 in the upper troposphere of Neptune using future models that include aerosol 521 extinction. 522

523 6. Conclusion

We used all available Herschel/SPIRE observations to constrain Nep-524 tune's CO and PH₃ abundances. A simultaneous fit of all nine CO bands 525 was possible using conventional step profiles, gradient profiles, and profiles 526 with zero deep abundance for pressures greater than 0.5 bar. The fitting 527 of multiple CO bands simultaneously improved the robustness of our abun-528 dance results. Our abundances ranged from 0.80–1.55 ppm in the strato-529 sphere (<0.1 bar) and 0.21–0.36 in the upper troposphere (1.0–0.1 bar), in 530 broad agreement with previous studies. 531

Importantly, the Herschel/SPIRE data to not require tropospheric CO 532 to exist at pressure levels deeper than 0.5 bar. This is also true of previous 533 observations of longer wavelength CO lines by Luszcz-Cook and de Pater 534 (2013) that probe slightly deeper (\sim 3 bar for the 115 GHz line depending on 535 profile assumptions). Therefore, both our observations and those previously 536 published are not very sensitive to the deep CO abundance, meaning that we 537 cannot confirm or deny the presence of CO for pressures deeper than 1 bar. 538 Caution must then be used in assuming that abundances measured in the 539 upper troposphere are representative of the deep interior. 540

⁵⁴¹ We suggest that the $\sim 0.1-0.2$ ppm deep tropospheric abundances previ-⁵⁴² ously reported could in fact be a result of extrapolating an idealised profile ⁵⁴³ to pressures where the data do not constrain the CO abundance. If this is ⁵⁴⁴ the case then extreme O/H enrichments would no longer be required in ther-⁵⁴⁵ mochemical interior models to explain a CO abundance throughout the tro-⁵⁴⁶ posphere, which allows the CO abundance profile and D/H measurements to ⁵⁴⁷ be reconciled. This means that current solar system water reservoirs become ⁵⁴⁸ a plausible water source, especially if the rock content of Neptune's interior ⁵⁴⁹ can be increased to dilute the high D/H ratio in these sources. The bulk of ⁵⁵⁰ Neptune's stratospheric and tropospheric CO can then be considered entirely ⁵⁵¹ externally sourced from a large cometary impact within the past few hundred ⁵⁵² years. We propose that a rock-rich Neptune as proposed by Feuchtgruber ⁵⁵³ et al. (2013) is more compatible with the current geophysical/spectroscopic ⁵⁵⁴ observations and possible formation scenarios.

⁵⁵⁵ PH₃ upper limits are sensitive to the temperature profile assumptions, ⁵⁵⁶ but fall in the range 0.192–5.52 ppb for the 0.1–1.2 bar pressure range, with ⁵⁵⁷ a nominal value of 1.10 ppb. This corresponds to 3.93×10^{-4} – 1.13×10^{-2} ⁵⁵⁸ times the solar value, which requires loss processes other than condensation ⁵⁵⁹ to explain. This is most likely due to photochemistry and the derived upper ⁵⁶⁰ limits are consistent with our simple photochemical modelling, which predicts ⁵⁶¹ negligible PH₃ in this pressure region.

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Study	Telescope	Freq.	Wavenumber	Uniform†	Step‡		
					v_1	v_2	p_1
		(GHz)	(cm^{-1})	(ppm)	(ppm)	(ppm)	(bar)
Luszcz-Cook and de Pater (2013)	CARMA	115.271	3.845	-	0 - 0.3	0.93 - 1.9	0.0025 - 0.2
	CARMA	230.538	7.670	-	"	"	"
	CARMA	345.796	11.535	-	"	"	"
Lellouch et al. (2010)	Herschel/PACS	$\sim \! 1750$	~ 60	-	~ 0.5	~ 1	~ 0.01
Hesman et al. (2007)	JCMT	345.796	11.535	-	$0.6{\pm}0.4$	$2.2^{+0.6}_{-0.4}$	0.006
Lellouch et al. (2005)	IRAM	230.538	7.670	-	$0.5 {\pm} 0.1$	$1{\pm}0.2$	0.02
Marten et al. (2005)	IRAM	230.538	7.670	$1.0 {\pm} 0.2$	-	-	-
	JCMT	345.796	11.535	"	-	-	-
	JCMT	461.040	15.379	"	-	-	-
Naylor et al. (1994)	JCMT	230.538	7.670	$1.0 {\pm} 0.3$	-	-	-
Guilloteau et al. (1993)	IRAM	115.271	3.845	0.6 - 1.5	-	-	-
Marten et al. (1993)	CSO	230.538	7.670	$1.2 {\pm} 0.4$	-	-	-
	JCMT	345.796	11.535	"	-	-	-
Rosenqvist et al. (1992)	IRAM	230.538	7.670	$0.65 {\pm} 0.35$	-	-	-

Table 1: Summary of previous microwave and far-IR CO studies. \dagger Early studies used a profile with a uniform mixing ratio throughout the troposphere and stratosphere. \ddagger Recent studies assume a step-type profile defined by a transition pressure p_1 , a high pressure uniform abundance v_1 , and a low pressure uniform abundance v_2 .

Obs. ID	Start Time	Integration Time	RA	DEC	Dist.	Disc
	(UT)	(s)	(°)	$(^{\circ})$	(AU)	(")
1342187090	2009-11-19 05:52:45.0	823.0	326.06	-14.06	30.07	2.27
1342187883	2009-12-09 00:26:43.0	823.0	326.34	-13.97	30.40	2.25
1342187884	2009-12-09 00:40:54.0	823.0	326.31	-13.98	30.40	2.25
1342187887	2009-12-09 01:23:18.0	876.0	326.32	-13.97	30.40	2.25
1342195348	2010-04-21 19:57:15.0	1145.0	330.47	-12.53	30.46	2.24
1342195771	2010-05-03 19:03:43.0	1145.0	330.68	-12.46	30.27	2.26
1342196617	2010-05-17 04:43:33.0	1145.0	330.83	-12.42	30.05	2.27
1342197362	2010-05-31 11:27:26.0	1145.0	330.90	-12.41	29.80	2.29
1342197363	2010-05-31 11:46:49.0	1145.0	330.88	-12.42	29.80	2.29
1342197364	2010-05-31 12:06:10.0	1145.0	330.89	-12.42	29.80	2.29
1342197365	2010-05-31 12:25:33.0	1145.0	330.90	-12.39	29.80	2.29
1342197366	2010-05-31 12:44:56.0	1145.0	330.87	-12.40	29.80	2.29
1342197367	2010-05-31 13:04:17.0	1145.0	330.88	-12.39	29.80	2.29
1342197368	2010-05-31 13:23:38.0	3597.0	330.89	-12.40	29.80	2.29
1342198429	2010-06-09 19:52:13.0	13762.0	330.87	-12.42	29.65	2.30
1342208385	2010-11-08 00:51:49.0	839.0	328.16	-13.40	29.82	2.29
1342210841	2010-12-05 15:26:40.0	839.0	328.39	-13.32	30.29	2.25
Total		32687.0				

Table 2: Summary of SPIRE observations extracted from the Herschel Science Archive (http://archives.esac.esa.int/hsa/whsa/). RA, right ascension; DEC, declination; Dist., distance between Neptune and Herschel; and Disc, projected diameter of Neptune on the sky.

Temperature	CO profile	Pre	Pressure VM		ЛR	χ^2/N
profile	type	p_1	p_2	v_1	v_2	
		(bar)	(bar)	(ppm)	(ppm)	
Cold	Step	0.14		0.21	0.80	0.69
Cold	Gradient	0.74	0.0009	0.23	0.91	0.74
Cold	External gradient	0.48	0.0018	0.35	0.93	0.79
Nominal	Step	0.11		0.22	1.03	0.63
Nominal	Gradient	0.75	0.0013	0.23	1.19	0.67
Nominal	External gradient	0.48	0.0035	0.36	1.01	0.89
Hot	Step	0.10		0.24	1.41	0.86
Hot	Gradient	0.54	0.0023	0.24	1.55	0.89
Hot	External gradient	0.47	0.0073	0.39	1.52	1.21

Table 3: Retrieved CO profile parameters assuming the nominal, hot, and cold temperature profiles in Figure 2. The external gradient profile has zero abundance for pressures greater than p_1 . χ^2/N is the reduced χ^2 misfit and should be ~1 for a model fitting the data to within error – all profiles except the hot external gradient profile provide adequate fits to the data. The nominal temperature profile with a step CO abundance profile (shown in bold) provides the best fit to the observations (Figure 5 and 6).

Temperature	Optimal	PH_3 band centre	Pressure sensitivity	3- σ UL	3- σ UL
profile	spectrometer	(cm^{-1})	(bar)	(ppb)	(x solar P/H^{\dagger})
Cold	SLW	17.81	0.8 - 1.2	$<\!\!5.52$	$< 1.13 \times 10^{-2}$
Nominal	SSW	44.46	0.4 - 0.8	<1.10	$<\!\!2.25{ imes}10^{-3}$
Hot	SSW	44.46	0.1 – 0.7	< 0.192	$< 3.93 \times 10^{-4}$

Table 4: PH₃ 3- σ upper limits for the three temperature profiles. In each case the optimal spectrometer is the one giving the lowest upper limit. Pressure sensitivity is the full-width half-maximum of the contribution functions shown in Figure 4 and in the absence of photochemistry is determined by the saturation vapour pressure curve. † Equivalent enrichment relative to solar P/H using the abundances from Lodders (2010), i.e. solar P/H=0.281×10⁻⁶, implying a solar composition volume mixing ratio PH₃/(H₂+He)=0.489×10⁻⁶ (assuming that He/H₂=0.15 and all phosphorous is the form of PH₃).



Figure 1: Neptune SPIRE SLW and SSW spectra. (a,b) Individual spectra from the 17 observations in Table 2. (c,d) Weighted mean spectra compared with synthetic baseline spectra (no CO or PH₃) for nominal, hot, and cold temperature profiles. (e,f) Ratio of synthetic spectra to observation in continuum regions for nominal temperature case. The smooth cubic b-spline curve is used to correct the observation for large scale continuum ripples. (g,h) Corrected spectrum for nominal temperature case and (i,j) standard error uncertainties. CO features are clearly visible in the data (labelled).



Figure 2: Neptune temperature profiles. The hot and cold profiles are ± 5 K from the nominal case, which is based on Lindal (1992) and Fletcher et al. (2010). Our analysis is performed using all three profiles to include the current uncertainty in Neptune's temperature structure.



Figure 3: Contribution functions for CO microwave bands. Calculations assume the SPIRE spectral resolution of 0.07373 cm^{-1} , the nominal temperature profile in Figure 2, and 0.1 ppm uniform CO abundance throughout the atmosphere. The upper tropospheric CO abundance gives wide absorption (negative/blue) features that are most prominent at lower frequencies, whereas the stratospheric abundance gives a narrow emission peak (positive/red) that is most prominent at higher frequencies. (a-c) Lowest frequency CO bands studied previously (e.g. by Luszcz-Cook and de Pater, 2013). (d-m) CO bands covered SPIRE, with * indicating the bands analysed here. (n) Total normalised contribution function summed over all analysed wavenumbers in (e-m) for 1 ppm CO (red) and 0.1 ppm CO (blue), appropriate for the stratosphere and troposphere respectively. Solid/dashed line indicates where these abundances are appropriate/not appropriate. Information content is restricted to the 0.001–0.01 and 0.1^{-44} bar pressure regions for SPIRE. The lower frequency lines (a–c) probe slightly deeper to ~ 3 bar, assuming a typical 5–10 GHz bandwidth (Luszcz-Cook and de Pater, 2013). The transition pressure used to define the step profile in previous studies typically falls in the 0.01-0.1 bar region, where there is little sensitivity.



Figure 4: Phosphine contribution functions for the 17.81 cm⁻¹ (a–d) and 44.46 cm⁻¹ (e–h) features under different temperature profile assumptions. For the nominal and cold cases saturation limits stratospheric abundances to negligible amounts. For the hot case some PH₃ can enter the stratosphere giving a small emission feature. (d,h) Total normalised contribution function summed over all wavenumbers in each band. Information is limited to the upper troposphere (0.1-1.2 bar).



Figure 5: Fitted CO features using the nominal temperature profile and a step VMR profile (bold profile highlighted in Table 3, and plotted in Figure 6). All nine CO bands are well fitted. The other profiles in Table 3 provide comparable fit qualities.



Figure 6: Fitted CO VMR profiles and PH₃ upper limits. (a) CO profiles have 0.80– 1.55 ppm in the stratosphere and 0.21–0.39 ppm in the upper troposphere. These data do not require significant deep CO abundance. (b) Phosphine upper limits are consistent with our simple photochemical model profiles, which predict significant PH₃ removal by photolysis for pressures less than \sim 2 bar. Upper limit profiles are limited by condensation, which occurs at pressures less than \sim 1 bar. Note that the hot profiles require more stratospheric CO due to the rescaling of the continuum, which increases the contrast of the emission peak (the opposite is true for the cold profile).



Figure 7: Phosphine upper limits. (a,d,g) variation of χ^2 as a function of deep phosphine abundance for the two spectral bands and all three temperature profiles. No significant PH₃ is detected and upper limits of 0.192–5.52 ppb are inferred. Observations and synthetics with zero PH₃ abundance, 3- σ PH₃ abundance, and an enhanced abundance to show the PH₃ feature shape more clearly are shown in (b,c) for the cold profile, (e,f) for the nominal profile, and (h,j) for the hot profile.