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Measurement of ammonia emissions from temperate and sub-polar seabird colonies

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1 **Measurement of ammonia emissions from temperate and sub-polar seabird** 2 **colonies**

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14 **Key Words:** Coastal nitrogen; seabirds; penguins; temperate; sub-polar; NH³ 15 emissions; atmospheric dispersion; inverse modelling

16 **Abstract**

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t: S. Riddick, T 17 The chemical breakdown of marine derived reactive nitrogen transported to the land 18 as seabird guano represents a significant source of ammonia (NH3) in areas far from 19 other NH₃ sources. Measurements made at tropical and temperate seabird colonies
20 indicate substantial NH₃ emissions, with emission rates larger than many indicate substantial NH₃ emissions, with emission rates larger than many 21 anthropogenic point sources. However, several studies indicate that thermodynamic 22 processes limit the amount of NH_3 emitted from guano, suggesting that the percentage
23 of guano volatilizing as NH_3 may be considerably lower in colder climates. This study of guano volatilizing as NH_3 may be considerably lower in colder climates. This study 24 undertook high resolution temporal ammonia measurements in the field and coupled 25 results with modelling to estimate NH_3 emissions at a temperate puffin colony and 26 two sub-polar penguin colonies (Signy Island, South Orkney Islands and Bird Island, 27 South Georgia) during the breeding season. These emission rates are then compared 28 with NH3 volatilization rates from other climates. Ammonia emissions were 29 calculated using a Lagrangian atmospheric dispersion model, resulting in mean 30 emissions of 5 μ g m⁻² s⁻¹ at the Isle of May, 12 μ g m⁻² s⁻¹ at Signy Island and 9 μ g m⁻² 31 s⁻¹ at Bird Island. The estimated percentage of total guano nitrogen volatilized was 32 5% on the Isle of May, 3% on Signy and 2% on Bird Island. These values are much 33 smaller than the percentage of guano nitrogen volatilized in tropical contexts (31- 34 65%). The study confirmed temperature, wind speed and water availability have a 35 significant influence on the magnitude of NH₃ emissions, which has implications for 36 reactive nitrogen in both modern remote regions and pre-industrial atmospheric 37 composition and ecosystem interactions.

38 **1. Introduction**

39 Nitrogen is found in all living cells and is necessary for the growth and survival of all 40 living things. However, nitrogen in its most abundant form, diatomic nitrogen (N_2) , is 41 a relatively un-reactive molecule and needs to be 'fixed' to become useable as 42 reactive nitrogen (N_r) compounds. Nr includes all N forms with the exception of N_2 ,
43 including ammonium and nitrate ions, gases such as nitrous oxide (N_2O) , nitrogen including ammonium and nitrate ions, gases such as nitrous oxide (N_2O) , nitrogen 44 oxides (NO_x) and ammonia (NH_3) and organic nitrogen compounds. Human activities,

45 including the Haber-Bosch process, legume cultivation and fossil fuel combustion, are 46 estimated to create 210 Tg of plant-useable N_r annually (Fowler et al., 2013).
47 Reactive nitrogen added to the Earth's surface as fertilizer can wash off into the Reactive nitrogen added to the Earth's surface as fertilizer can wash off into the 48 hydrosphere, volatilize to the atmosphere as $NH₃$ or form organic nitrogen compounds 49 in soils. Further decomposition of oceanic, terrestrial, plant and animal N_r can produce N_2 as well as NO_x and N_2O . produce N_2 as well as NO_x and N_2O .

iems causing acidification and eutrophication, which has been shown to alter
perspects competition and biodiversity (Cape et al., 2009). Stuton et al., 201
Currently, the biogeochemical processes following the addition of 51 Studies suggest the emission of NH_3 gas is likely to negatively impact local ecosystems causing acidification and eutrophication, which has been shown to alter 52 ecosystems causing acidification and eutrophication, which has been shown to alter 53 local interspecies competition and biodiversity (Cape et al., 2009; Sutton et al., 2011, 54 2012). Currently, the biogeochemical processes following the addition of seabird 55 derived N_r to the surface of land are not well understood. However studies have 56 reported NH3 emission from poultry excreta which has similar properties to seabird 57 guano (Elliott and Collins, 1982; Harper et al., 2010) and a study of Adelie penguin 58 colony on the Antarctic continent suggests volatilized NH3 creates a spatial impact 59 zone of up to 300 km^2 surrounding the colony where phosphomonoesterase activity is 60 increased in indigenous organisms (Crittenden et al., 2014). In order to be emitted as 61 NH3, excreted uric acid must first be hydrolysed under microbial decomposition to 62 produce ammonium and bicarbonate ions. Both the processes of uric acid hydrolysis 63 and NH_3 volatilization appear to be affected by environmental conditions, including water availability and temperature (Nemitz et al., 2001; Sutton et al., 2013). Food water availability and temperature (Nemitz et al., 2001; Sutton et al., 2013). Food 65 composition and pH may also play a significant role in $NH₃$ emission (Elliott and 66 Collins, 1982; Harper et al., 2010) where NH_3 emission depends on the ratio between the nitrogen and energy content of the food (Wilson et al., 2004) and the pH affects 67 the nitrogen and energy content of the food (Wilson et al., 2004) and the pH affects 68 the rate at which uric acid is converted to ammonium (Elliott and Collins, 1982).

69 In a theoretical study on seabird N_r excretion by Riddick et al. (2012), the estimated 70 percentage of N_r that volatilizes (P_v) ranged from 9 % in colder temperatures (average 71 temperature during breeding season *c.* 5°C) to 100 % at colonies in higher 72 temperatures ($> 19^{\circ}$ C). Recent measurement-based estimates showed mean P_v values 73 of 31 to 65 % at two tropical seabird colonies estimated (Riddick et al., 2014). 74 Additionally, some variation in *Pv* is expected in relation to habitat, so that birds 75 nesting in vegetation and breeding in burrows (such as puffins), would show a lower 76 percentage emission as NH_3 as compared with birds nesting and breeding on bare
77 rock surfaces (Blackall et al., 2007: Riddick et al., 2012). Similarly, Zhu et al. (2011) 77 rock surfaces (Blackall et al., 2007; Riddick et al., 2012). Similarly, Zhu et al. (2011) 78 suggest temperature is an important driver in the production of NH_3 , however they
79 also suggest temperature may not be the sole climatic variable that affects NH_3 also suggest temperature may not be the sole climatic variable that affects $NH₃$ 80 emission.

81 Seabird colonies are well suited for measuring NH_3 emissions because they are searchly remote from human activity, resulting in near-background NH_3 generally remote from human activity, resulting in near-background $NH₃$ 83 concentrations in the surrounding area. Biogeochemical processes are relatively 84 simple because the majority of seabirds nest on rocky surfaces where excreted guano 85 can: (1) build up on the surface; (2) decompose, converting uric acid to ammoniacal 86 forms which are liable to volatilization, or (3) be washed into the sea. As a model 87 system for studying the effect of climate/environment on NH_3 emissions, seabird 88 colonies also have the advantage that they are generally not influenced by human 89 management practices (other than those which may affect seabird numbers). In 90 addition to this, the penguin species' annual presence in the nitrogen poor regions of 91 the Southern Ocean supplies 858 Gg of N_r per year (\sim 3 kg m⁻²) in the form of guano 92 to the land (Riddick et al., 2012). In agriculture terms, the average penguin colony 93 receives 30,000 kg ha^{-1} compared with 246 kg ha^{-1} for fertilizer consumption on 94 arable land in the UK in 2015 (Worldbank, 2015).

95 As a result of these features, seabird colonies offer a system that is well fitted to 96 address the question of how NH_3 emission rates vary globally through different 97 climatic regimes as well as develop understanding of atmosphere-ecosystem 97 climatic regimes as well as develop understanding of atmosphere-ecosystem 98 interaction in the natural world. The present study contributes to this question by 99 providing data on NH_3 emissions from seabird guano in temperate and sub-polar conditions. for comparison with previous measurements in tropical conditions 100 conditions, for comparison with previous measurements in tropical conditions 101 (Riddick et al., 2014). By bringing these measurements together with other published 102 datasets, we are then able to investigate the global scale variation in $NH₃$ emission 103 rates.

104 **2. Methods and Materials**

105 **2.1 Ammonia measurements**

106 Two methods were applied in this study to make NH_3 concentration measurements:
107 (1) passive sampling and (2) an on-line active sampling NH_3 analysis instrument, as (1) passive sampling and (2) an on-line active sampling $NH₃$ analysis instrument, as 108 summarized below.

109 The passive samplers used (ALPHA samplers, CEH Edinburgh) consist of a 23 mm 110 diameter sampler with a 6 mm diffusion path between a Teflon membrane and an 111 adsorbent sampling surface (filter-paper disc impregnated with citric acid). Further 112 details of ALPHA sampler and its system of pre- and post-sampling protective caps 113 are provided by Tang et al. (2001). In this study, triplicate samplers were used at each 114 sampling location and exposed for periods of 2 to 4 weeks. The samplers were 115 attached by Velcro to an upturned plant saucer (for protection) that was fastened to a 116 pole (The sampling heights above the ground for the different sites are described 117 below, with further details given in Supplementary Material 7). Aluminium strips 118 were mounted on top of each saucer to deter perching birds.

ons, for comparison with previous measurements in tropical condition
set set al., 2014). By bringing these measurements together with other publishes, we are then able to investigate the global scale variation in NH₃ em 119 At all times, except during deployment, the ALPHA samplers were sealed in plastic 120 containers and refrigerated. In the laboratory, the $NH₃$ concentration of the air at the 121 seabird colony was determined using ammonium flow injection analysis, based on 122 selective diffusion of NH3 across a Teflon membrane at high pH (FLORRIA, 123 Mechatronics, NL). Laboratory and field blanks were also analysed to ensure samples 124 were not contaminated. In the present study, the high sensitivity ALPHA samplers 125 were used with a Method Detection Limit $(MDL) = 0.09 \mu g m⁻³$ for two-weekly 126 exposure on Signy Island. A description of how the MDL was calculated is given in 127 Supplementary Material Section 1. ALPHA samplers were also deployed at Bird 128 Island and the Isle of May for comparison with the on-line measurements.

129 The on-line NH3 concentration measurements were made with an AiRRmonia gas 130 analyser (Mechatronics, NL) on Bird Island and a Nitrolux 1000 gas analyser 131 (Pranalytica, USA) on the Isle of May. At each site air was drawn into the instrument 132 through 20 m PTFE tubing, to minimize NH3 sticking the PTFE tubing was heated 133 and insulated a full description of the online active measurement set up is given in 134 Supplementary Material Section 3, with inlet flows of 8 1 min^{-1} .

135 The AiRRmonia analyser (Norman et al., 2009) is based on a similar principle to the 136 FLORRIA. In this case, atmospheric air is passed over a first Teflon membrane with a 137 counterflow of dilute acid to allow gaseous $NH₃$ to transfer to aqueous ammonium in 138 solution. Sodium hydroxide is then added to liberate molecular NH_3 , which then diffuses across a second Teflon membrane into a counter flow of deionized water, diffuses across a second Teflon membrane into a counter flow of deionized water,

140 with reformed ammonium then detected by conductivity. The AiRRmonia has an 141 instrument delay time (the time taken between air sampling and instrument response) 142 of ~ 5 minutes with 15 min averages used to assure quantitative response, with a 143 Limit of Detection (LOD) of $\sim 0.1 \mu g$ m⁻³ and a MDL in this context of 0.07 μg m⁻³. 144 The AiRRmonia measurements were recorded every minute and then the data 145 averaged to 15 minute periods for application in the inverse dispersion model. 146 Calibration of the AiRRmonia was carried out every five days and agreed within 5% 147 over the periods of measurement.

148 The Nitrolux analyser is a photoacoustic instrument that uses absorption of NH_3 molecules from a line-tuneable CO_2 laser to measure concentration. The Nitrolux molecules from a line-tuneable $CO₂$ laser to measure concentration. The Nitrolux 150 1000, as used here, has a detection limit of $\sim 0.1 \mu$ g m⁻³, a MDL in this context of 0.1 151 μ g m⁻³, a range of 0.1 – 2000 μ g m⁻³, and measures concentrations every 45 s. The 152 instrument delay time of the instrument is a function of temperature and relative 153 humidity (typically 4 (3-5) minutes), allowing the data to be averaged up to 15 minute 154 periods for application in the inverse dispersion model. The Nitrolux 1000 requires 155 six-monthly calibrations (Cowen et al., 2004).

156 **2.2 Field Methodology**

157 **Site 1: The Isle of May, Scotland**

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instraction 2.1 and alterction limit of $\neg A$ 158 The Isle of May (56.19 °N, 2.56 °W) is a nesting site for many seabird species, 159 including Common Guillemot (*Uria aalge*), Herring Gull (*Larus argentatus*), Arctic 160 Tern (*Sterna paradisaea*), Black-legged Kittiwake (*Rissa tridactyla*) and Atlantic 161 puffin (*Fratercula arctica*). The island is located at the entrance to the Firth of Forth 162 in eastern Scotland (Figure 1) and has a temperate climate (average temperature of 163 15°C, average humidity of 80% and average wind speed of 4 m s^{-1} during the 164 breeding season). Passive and active measurements of $NH₃$ concentrations and 165 meteorological parameters were made above Atlantic puffin burrows (Figure 1). 166 Atlantic puffins breed on vegetated slopes and amongst rocky outcrops, where they 167 dig and nest in 1-2 m long burrows. Atlantic puffins burrow in most parts of the Isle 168 of May with a colony total of 45,000 pairs during June and July 2009 (Harris et al., 169 2009), with approximately 20,000 burrows in our study area, between the Low Light 170 and Kirk Haven (area shaded dark grey in Figure 1). Measurements were carried out 171 from 01/07/09 to the 06/09/09 during (July) and after (August and September) the 172 period of chick rearing, where large numbers of prospecting juvenile birds are present 173 in addition to breeding birds.

174 Active Sampling Campaign

175 The Nitrolux trace gas analyser measured NH3 concentrations on-line over the 176 Atlantic puffin colony on the Isle of May from 30/06/09 to 23/07/09. The air inlet was 177 positioned 1.26 m above the ground at the measurement site (labelled in Figure 1). 178 Measurements during the Isle of May campaign were limited to daylight hours to 179 reduce disturbance to fledging puffins by the generator. Micrometeorological 180 parameters were measured using a Gill Windmaster Pro sonic anemometer on a mast 181 2.5 m above the ground. Meteorological data were collected by instruments on a mast 182 on the highest point of the island (Figure 1). Data collected included: air temperature, 183 relative humidity, solar radiation (all at 1 m above ground) and ground temperature 184 was using temperature sensors on the surface. The weather station was located away 185 from the colony to avoid interfering with birds' nesting behaviour.

186 <<INSERT FIGURE 1 HERE>>

187 Passive Sampling Campaign

188 Triplicate ALPHA samplers were used to measure NH3 concentrations above the 189 Atlantic puffin colony ("Measurement Site", as labelled in Figure 1), at a height of 1.5 190 m, for 4 periods of 15 days, as described in Supplementary Material Section 7A. 191 Meteorological data were collected by a weather station positioned at the highest 192 point of the island (Figure 1).

193 **Site 2: Bird Island, South Georgia**

194 Bird Island is part of South Georgia, 1000 km south-east of the Falkland Islands 195 (Figure 2). Ammonia concentrations were measured at the 'Big Mac' Macaroni 196 penguin (*Eudyptes chrysolophus)* colony at the western end of the island (54.0106 °S, 197 38.0753 °W), where 40,000 breeding pairs were present during the measurement 198 period from 07/11/10 and 26/12/10 (D. Briggs, British Antarctic Survey, pers. 199 comm.). Immediately to the east of the active measurement site, the 'Little Mac' 200 colony is located (450 pairs of Macaroni penguin in a small satellite colony). The 201 average temperature was 3°C, average relative humidity 92 % and average wind 202 speed 5 m s⁻¹ during the measurement period.

203 Active Sampling Campaign

EVALUATION Coordinates were measured at the Balkland Island

is part of South Georgia, 1000 km south-east of the Falkland Island

2). Ammonia concentrations were measured at the 'Big Mac'-Macarot

22). Ammonia concentra 204 On-line NH3 concentrations were measured at Fairy Point to the south of Big Mac 205 (Figure 2). The air inlet for the AiRRmonia analyzer was positioned at 2 m above the 206 ground. All the instruments were housed in a tent to provide protection from the 207 wind, precipitation, sea spray and sun. Micrometeorological parameters were 208 measured using a Gill Windmaster Pro sonic anemometer on a mast 2.5 m above the 209 ground. Meteorological data were collected by instruments on two masts on the 210 highest point at Fairy Point. Data collected included: air temperature, humidity and 211 solar radiation at 1 m above ground, and wind speed at three heights above ground 212 (0.5 m, 1 m, and 2 m). Ground temperature was measured using a Tiny Talk data 213 recorder placed on the ground (Supplementary Material Section 4).

214 <<INSERT FIGURE 2 HERE>>

215 Passive Sampling Campaign

216 Ammonia concentrations were recorded at nine locations on Bird Island using 217 ALPHA samplers mounted at 1 m above ground (Figure 2). These were exposed in

218 seven sampling periods of around 2 weeks from 07/11/2010 to 26/12/2010.

219 **Site 3: Signy Island, South Orkney Islands**

220 Signy Island is a small island in the South Orkney Islands in the Southern Ocean 221 (Figure 3). A relatively flat area on the Gourlay Peninsula was used for passive 222 sampling of NH_3 concentrations from 10/01/2009 to 21/02/2009 at a colony of 10,000 223 pairs of Adélie penguins (*Pygoscelis adeliae)* and 9,000 pairs of Chinstrap penguins 224 (*P. antarcticum)* (60.73° S, 45.59° W). Both species breed in snow free areas and 225 build rudimentary nests of small stones. The climate at this site represents sub-polar 226 conditions with average temperature of 2° C, average relative humidity of 84 % and 227 average wind speed of 5 m s⁻¹ during the breeding season.

228 ALPHA samplers were deployed at five locations (Mast $1 - 5$, Figure 3) over three 229 separate sampling periods of 2 weeks each. Masts 1 and 2 had ALPHA samplers at 1 230 m and 1.5m from the ground. Mast 5 was located as far as possible from any birds to 231 sample background NH3 concentrations, *en route* from the base at Borge Bay to 232 Gourlay (Supplementary Material 3). Representative meteorological data 233 (temperature, wind speed, relative humidity and precipitation) were obtained from the
234 nearest weather station, the Argentinean Orcadas Base on Laurie Island, South nearest weather station, the Argentinean Orcadas Base on Laurie Island, South 235 Orkney Islands (US National Climatic Data Center (NCDC) Integrated Surface 236 Hourly (ISH) database; NCDC, 2011).

237 <<INSERT FIGURE 3 HERE>>

238 **2.3 Estimation of NH3 Emissions**

239 Estimates of NH_3 emissions were calculated using an inverse application of the 240 WindTrax atmospheric dispersion model version 2.0 (Flesch et al., 1995). Given 241 potential temporal covariance between atmospheric NH3 concentrations and 242 dispersion, such calculations should ideally be based on short-term measured 243 concentrations.

244 For input into WindTrax, both the on-line NH_3 concentrations and meteorological 245 data were averaged over 15 minutes to minimise any effects of turbulence while 246 preserving variation caused by environmental or atmospheric change (Laubach et al. 247 $\,$ 2008; Flesch et al. 2009). Fifteen minute averages of wind speed $(u, m s⁻¹)$, wind direction (*WD*, \degree), temperature (*T*, \degree C), NH₃ concentration at 2 m (*X*, ug m⁻³). 249 roughness height (z_0 cm) and the Monin-Obukhov length (L , m) were used as input to 250 WindTrax.

251 For each on-line NH3 concentration dataset, data were removed for calibration 252 periods, any periods when the instrument was not sampling the colony due to wind 253 direction and any periods of high atmospheric stability (wind speed, $u < 0.15$ ms⁻¹, 254 friction velocity, $u^* < 0.1$ ms⁻¹ and Monin-Obukhov Length $|L| < 2$). Each WindTrax 255 simulation used 50,000 particle projections to back-calculate the NH₃ emission.

thand of Terry Emassions were calculated using an inverse application of the
trax atmospheric dispersion model version 2.0 (Flesch et al., 1995). Give
the of NH₃ emissions were calculated version 2.0 (Flesch et al., 199 256 While the first focus of the emission calculations was on applying the on-line $NH₃$ 257 concentration measurements, it is also of interest to assess how the inverse model 258 performs when using time-integrated NH_3 concentrations, since it is not always feasible to deploy on-line NH_3 instrumentation (e.g. as at Signy Island). For this feasible to deploy on-line NH_3 instrumentation (e.g. as at Signy Island). For this 260 reason, we also applied the Windtrax model using two-weekly averaged NH³ 261 concentrations, coupled with the time-resolved estimates of atmospheric turbulence. 262 In principle, this relaxation is expected to contribute significant errors in the resulting 263 flux estimates. However, experience under other conditions indicates that these errors 264 may be small when compared with other sources of error or with the difference in 265 emission rates between sites (Riddick et al., 2014; Theobald et al., 2013). The 266 deployment of both passive and active sampling at the Isle of May and at Bird Island 267 allowed comparison these two approaches, providing a basis to assess confidence in 268 the passive measurements at Signy Island, where only the passive NH_3 concentration 269 data were available.

270 The comparison of estimated NH_3 emissions calculated using the passive and on-line 271 sampling methods can also be used to provide an indicative estimate of the respective 272 sources of error in each approach (Riddick et al., 2014). To do this, the concentrations 273 recorded by the on-line continuous NH_3 detector are first averaged for the same 274 periods as the passive ALPHA sampler data, and then used to estimate NH_3 fluxes 275 using the WindTrax system. The difference in mean flux between the approach using 276 15 minute NH_3 concentrations and the 2-weekly averaged data from the on-line 277 system gives an estimate of the micrometeorological error associated with low-time 278 resolution NH_3 concentration data. By comparison, the difference in mean flux 279 between the 2-weekly averaged data of the on-line system and the 2-weekly estimates 280 from the ALPHA samplers gives an estimate of the chemical sampling error. This 281 chemical sampling error can be mostly associated with the on-line system, because it 282 only samples for part of the time (i.e. semi-continuous), as compared with the passive 283 system, which samples continuously.

284 **2.4 Other Uncertainties**

285 In order to further understand the uncertainties in the emission calculation, the input 286 variables were assessed for both field sites. The uncertainty caused by each variable 287 was estimated using WindTrax to back-calculate the consequent change in estimated 288 NH₃ emission. The total uncertainty was then calculated as the square root of the sum of the squares of the individual uncertainties. Further details are provided in the 289 of the squares of the individual uncertainties. Further details are provided in the 290 Supplementary Material Section 6.

291 **3. Results**

292 **3.1 Isle of May**

293 Active measurements and meteorological data

294 Measured NH₃ concentrations ranged from 0 to 105 μ g m⁻³ and were found to be 295 lower during the morning and evening than during the day (Figure 4). Ground 296 temperature ranged from 12 to 27 \degree C and peaked during the early afternoon. The 297 roughness length estimated using the ultrasonic anemometer on the Isle of May 298 ranged from 0.1 to 13.8 cm, i.e., within the useable range of WindTrax. Ammonia 299 emissions generally followed a diurnal pattern with low emission early in the morning 300 $\left($ <5 μ g m⁻² s⁻¹), building to a peak in the early afternoon (10 to 25 μ g m⁻² s⁻¹), before 301 dropping back to low values (<5 μ g m⁻² s⁻¹) in the evening (Figure 4). Overall, for the 302 active measurements the average emission rate was 5 μ g m⁻² s⁻¹.

To trained using WindTrax to be an
example and the constantine of the sense are severe assessed for both field sites. The uncertainty caused by each variably
inision. The total uncertainty was then calculated as the squar 303 The uncertainty in background NH_3 concentration for the southern North Sea (0.03 -304 1.49 μ g m⁻³) resulted in an emission uncertainty of 6%. The uncertainty in the size of 305 the NH₃ emission area (range of $0.2 - 0.3 \text{ km}^2$), caused by puffins moving around 306 near their burrows during the day, resulted in an uncertainty in NH₃ emission of 10 % 307 (Supplementary Material Section 6). Considering only these components, the overall 308 uncertainty in the modelling of the emission estimate on the Isle of May is estimated 309 at 12 %. A major source of uncertainty is the representatively of the NH₃ sampling, 310 given that measurements were only made for part of the time, with the generator 311 having to be switched off during the hours of darkness. This is addressed further in 312 section 3.4.

- 313 <<INSERT FIGURE 4 HERE>>
- 314 Passive measurements

315 Ammonia concentrations decreased from a maximum of 36.1 μ g m⁻³ during the first 316 period to a minimum of 0.9 μ g m⁻³ during the fourth measurement period, due to 317 measurements being made towards the end of the breeding season. The NH₃ emission 318 was highest during Period 1 (01/07/09 - 15/07/09), estimated at 5.1 µg m⁻² s⁻¹. By mid-319 July, most puffins had fledged and had left the nesting site. As a consequence, $NH₃$ 320 emission decreased to 1.9, 0.4, 0.1 μ g m⁻² s⁻¹ during measurement periods 2, 3 and 4, 321 respectively (for more details see Supplementary Material Section 7A). Temperatures 322 were broadly similar through the four sampling periods (Supplementary Material 323 Section 7A).

 324 The uncertainty in the estimated emission caused by the roughness length, NH₃ 325 background and emission area were 12, 8 and 10 %, respectively (See Supplementary 326 Material Section 6). The largest estimated uncertainty was the Monin-Obukhov length 327 at 28%. Overall, these factors contributed a combined uncertainty of \pm 38 % to the 328 model results from the passive campaign on the Isle of May. However, this does not 329 include the micrometeorological uncertainty associated with long-averaging periods, 330 which is considered separately in Section 3.4.

331 **3.2 Bird Island, South Georgia**

332 Active measurements and meteorological data

d Island, South Georgia

measurements and meteorological data

H₃ concentrations measured by the AiRRmonia trace gas analyser wer

H₃ concentrations measured by the AiRRmonia trace gas analyser wer

n 0 and 60 µ m⁻ 333 The NH₃ concentrations measured by the AiRRmonia trace gas analyser were 334 between 0 and 60 μ g m⁻³, with higher concentrations recorded during the daytime 335 (Figure 5). Ground temperature ranged from 1 to 12 °C, with maximum values 336 during the early afternoon (Figure 5). The roughness length estimated from the ultra-337 sonic anemometer on Bird Island ranged from 6 to 12.5 cm and was within the 338 useable range of WindTrax. Gras (1983) estimated open water background NH³ 339 concentration for Antarctica, a location representative of this area, at 0.15 μ g m⁻³. 340 which was used as the background concentration in WindTrax. The minimum and 341 maximum NH3 emissions from the Big Mac penguin colony during the measurement 342 period were 0.6 μ g m⁻² s⁻¹ and 52.6 μ g m⁻² s⁻¹, respectively (Figure 5). The largest 343 emissions occurred during the daytime, associated with higher wind speeds (Figure 344 5), with smaller emissions at night.

345 The emission uncertainty caused by the uncertainty in the size of the excretion area, 346 again caused by penguins moving around the edge of the nesting site, and NH³ 347 background were estimated at 27 % and 4 %, respectively (Supplementary Material 348 Section 6). The combined uncertainty calculated for the modelled emission 349 estimate from the Big Mac penguin colony was at \pm 28 %. The additional uncertainty 350 associated with the semi-continuous nature of the NH3 measurements is examined in 351 Section 3.4.

352 <<INSERT FIGURE 5 HERE>>

353 Passive measurements

354 Ammonia concentrations nearest the colony (3 m from the edge of Big Mac) 355 decreased from a maximum of 34.2 μ g m⁻³ during the third period (21/11/2010 to 356 28/11/2010) to a minimum of 11.3 μ g m⁻³ during the fifth measurement period 357 (06/12/2010 to 12/12/2010; NH₃ concentration data is presented in Supplementary 358 Material Section 7B, full transect data to be published elsewhere (Tang et al. in prep.). 359 The NH3 emission, calculated with WindTrax, was highest during Period 2 (Table 1), 360 estimated at 11.2 μ g m⁻² s⁻¹ and lowest during the fifth measurement period at 3.2 μ g $361 \text{ m}^2 \text{ s}^{-1}.$

 362 The uncertainty in the estimated emissions caused by the roughness length, NH₃ 363 background and emission area were 15, 12 and 12%, respectively (Supplementary 364 Material Section 6). The largest estimated uncertainty was associated with 365 micrometeorology at 35%. Overall, these amount to a combined uncertainty for the 366 passive campaign on Bird Island of \pm 42%.

367 **3.3 Signy Island**

368 On Signy Island the ALPHA samplers were exposed for three two-week periods 369 (Supplementary Material Section 7C). The NH₃ concentrations at Masts 1 and 2,

370 measured at a height of 1 m above the ground in the middle of the colony, were the 1371 highest (maximum 483 μg m⁻³) of the different sampling locations at Signy. NH₃ concentration decreased with distance from the penguin colony to a minimum at Mast 372 concentration decreased with distance from the penguin colony to a minimum at Mast $5(0.9 \text{ to } 2.1 \text{ µg m}^{-3})$. The ALPHA samplers lower to the ground (1 m height) 374 measured larger NH_3 concentration, as expected (see Supplementary Material Section 375 \degree 7C for details). The atmospheric conditions averaged over the measurement period 375 7C for details). The atmospheric conditions averaged over the measurement period 376 were estimated as neutral, (i.e. $(L = |\infty|)$ because of low ground heating and relatively 377 high wind (Seinfeld and Pandis, 2006). The most obvious sources of aerodynamic 378 roughness in the otherwise very flat area were the penguins (average height 60 cm) 379 and any larger rocks (maximum height estimated at 1 m). Therefore, a roughness 380 height of 10 cm, corresponding to an object height of 1 m (Seinfeld and Pandis, 381 2006), was used for modelling. The NH3 source area was assumed to be the observed 382 nesting area, which was 2.7×10^3 m².

383 The calculated NH_3 emission fluxes for the penguin colony on Signy Island were 18, 384 8 and 9 μ g m⁻² s⁻¹ for periods1, 2 and 3, respectively. The wind was almost constantly 385 from the north-west, which suggests that the footprint of the source sampled by each 386 ALPHA sampler was not a very significant source of variation. The 387 micrometeorological conditions on Signy Island could only be estimated from 388 available data on Laurie Island, South Orkney Islands, and therefore a larger 389 uncertainty is associated with meteorological data needed to estimate NH_3 emissions.

- 390 The difference in the NH3 emission rates between the first and second/third 391 measurement periods may be explained by the birds' behaviour, with colony 392 attendance during the first measurement period being high for both Adélie and 393 Chinstrap penguins. The lower emissions during the second and third periods may be 394 associated with the departure of the Adélie penguins around late January.
- not (Semmen data random and particular and particular and particular of the control of the meaning (semper rocks (maximum height estimated at 1 m). Therefore, a roughness of a reduced by larger rocks (maximum height stima 395 Together, the uncertainty in roughness length and stability resulted in an uncertainty 396 in emission of 26 % (Supplementary Material Section 6). The uncertainty associated 397 with background concentration from Gras (1983) was 7 % and the associated 398 uncertainty in area was estimated at \pm 6 %. The combined uncertainty in modelling 399 NH₃ emissions for Signy Island was estimated at \pm 37 %, although this does not 400 include uncertainty related to application of the time-integrated ALPHA sampling, 401 which is addressed in Section 3.4.

402 **3.4 Comparison of Active and Passive Sampling methods**

- 403 A summary of the measurements made at the different colonies of this study is 404 provided in Table 1. For the Isle of May, the mean fluxes from the passive and active 405 sampling campaigns were 5.1 and 5.3, μ g m⁻² s⁻¹, respectively. The estimate of the 406 flux from the active sampling averaged for the same period as the ALPHA 407 measurements was 6.0 μ g m⁻² s⁻¹. The difference between the first and third of these 408 fluxes represents the Uncertainty in Sampling Period (USP), at -1.0 μ g m⁻² s⁻¹, while 409 the difference between the second and third of these represents the Uncertainty in 410 chemical Sampling Method (USM), at -1.0 μ g m⁻² s⁻¹. In both cases the USP and 411 USM amount to around +/-20% of the mean flux at Isle of May.
- 412 <<INSERT TABLE 1 HERE>>

413 A similar comparison of active and passive sampling at Bird Island gave a mean flux 414 during the first period from the passive and active sampling campaigns of 11.2 and 415 10.3 μ g m⁻² s⁻¹, respectively. The mean fluxes during the second period from the 416 passive and active sampling campaigns were 8.9 and 10.5 μ g m⁻² s⁻¹, respectively. 417 The estimate of the flux from the active sampling averaged for the first and second 418 periods as the ALPHA measurements was 10.6 and 10.7 μ g m⁻² s⁻¹, respectively. The 419 estimate of the flux from the active sampling averaged for the average of the two 420 periods of the ALPHA measurements was 10.7 μ g m⁻² s⁻¹. In this case the USP 421 amounts to around 3% of the mean measured fluxes, whereas the USM was 6% for 422 the first period and 17% for the second period (Table 1).

423 In the case of Signy, only passive estimates of the flux were available, where the 424 overall mean of the three runs was 12 μ g m⁻² s⁻¹. Although active sampling was not 425 possible at this site, the performance comparison distinguishing USP and USM at Isle 426 of May and Bird Island may be taken as an indication of the scale of uncertainty 427 associated with the long sampling periods on Signy.

428

429 **4. Discussion**

430 **4.1 Variation in NH3 emissions from seabird colonies**

431 The largest weekly average NH₃ emission measured by this study was 18 μ g m⁻² s⁻¹ 432 on Signy Island, South Orkney Islands. Higher rates of NH₃ emission (22 μ g m⁻² s⁻¹) 433 were observed above the Brown noddy colony on Michaelmas Cay, Great Barrier 434 Reef, Australia (Riddick et al., 2014), while Blackall et al. (2007) reported even larger 435 emission rates equivalent to 240 μ g m⁻² s⁻¹ from Atlantic gannets on the Bass Rock, 436 Scotland. These results illustrate how NH₃ emissions from seabird colonies are 437 considerable discrete NH3 sources in a wide range of climates.

438 However, such figures tend to mask the climatic dependence of NH3 emission, since 439 they are also a function of nesting density, and for total colony emissions, of bird 440 numbers, types and colony attendance, etc. It is therefore helpful to normalize the 441 emission rates per g of bird biomass. In this case, it can be seen that NH_3 emission is 442 much higher at the tropical colony (7.5 \pm 2.6 mg NH₃-N g⁻¹ bird yr⁻¹; Michaelmas Cay) than at the sub-polar Bird Island colony reported here $(0.05 \pm 0.01 \text{ mg NH}_3\text{-N g}$ 443 444 ¹ bird yr⁻¹).

mean of the three runs was 12 µg m² s⁻¹. Although active sampling was no
e at this site, the performance comparison distinguishing USP and USM at 1sl
at distinguishing USP and USM at 1sl
at and Bird Island may be take 445 Another way to normalize the NH_3 emission data is to calculate the percentage of 446 excreted nitrogen that volatilizes as NH_3 (P_v , %), as described in Supplementary 447 Material Section 8. An excretion rate (Furness et al., 1991; Wilson et al., 2004), 448 calculated from the adult/chick mass, nitrogen content of the food, energy content of 449 the food, assimilation efficiency of ingested food and proportion of time spent at the 450 colony during the breeding season has been used instead of direct measurements of 451 guano depth up at the colony to reduce disturbance to breeding birds and minimize the 452 risk of egg/chick abandonment. For the measurements reported here, a P_v value of 453 4.7 \pm 0.5 % was calculated for the Atlantic puffin colony on the Isle of May, 4.7 ± 0.5 % was calculated for the Atlantic puffin colony on the Isle of May, 454 compared with 1.6 ± 0.4 % for Bird Island and 3.1 ± 1.1 % for Signy Island, 455 respectively (percentage error in measurement and modelling; Table 1).

456 In Table 2 the values from the present study are compared with emission rates and 457 estimates of P_v from other published studies. This shows the largest values of P_v at 458 tropical colonies, such as the Brown noddy colony on Michaelmas Cay, where P_ν was estimated at 65 \pm 22 % (Riddick et al., 2014), and the smallest values in sub-polar estimated at 65 ± 22 % (Riddick et al., 2014), and the smallest values in sub-polar 460 conditions, with comparable values for Bird Island and Signy Island (2%, 3%, 461 respectively) and Cape Hallet on mainland Antarctica (2%, Theobald et al., 2013). 462 These observations are in agreement with Zhu et al. (2011) who also found that $NH₃$ 463 emissions are larger under increased temperature. However, moisture limitation can 464 also be important at high temperatures.

465 As Riddick et al. (2014) showed for the two tropical islands, the higher value for 466 Michaelmas Island (67%) than for Ascension Island (32%) reflected a moisture 467 limitation at the latter site. In this instance, of two sites with similar temperatures, it 468 appears that the limited water availability at Ascension Island resulted in a lower rate 469 of uric acid hydrolysis, thereby leading to lower NH_3 emissions. By contrast, the 470 overall increase in observed P_y with increasing temperature across the sites (Table 2) overall increase in observed P_v with increasing temperature across the sites (Table 2) 471 may be a consequence of both increasing volatility of $NH₃$ and increasing rates of uric 472 acid hydrolysis, where sufficient moisture is available, although it is not possible to 473 distinguish these component effects from our measurements. In order to examine 474 these drivers separately, specific process modelling is needed (Riddick, 2012; Riddick 475 et al. in prep).

476 <<INSERT TABLE 2 HERE>>

increase in observed P_v with increasing temperature across the sites (Table 2 consequence of both increasing volatility of NH₃ and increasing rates of university and the sconfigured in M₃ and increasing rates of uni 477 It is worth noting that the measured P_v for the Atlantic puffin colony on the Isle of 478 May (5%) is much lower than the estimate by Riddick et al. (2012) and the 479 measurements made in similar conditions on the rocky cliffs of the Isle of May 480 (Guillemot) and Bass Rock (Northern gannet) by Blackall et al. (2004; 2007) (16- 481 36%). The much lower emission rate for Atlantic puffins, compared with Northern 482 gannets and Guillemot under the same climate, may be attributed to their habitat 483 preference as burrow nesters in grassland. This illustrates how climatic conditions are 484 not the only factors to affect NH_3 emission. In the case of the puffins on the Isle of 485 May case, the comparison suggests that emissions rates are about 14-31% of what May case, the comparison suggests that emissions rates are about 14-31% of what 486 would be emitted by bare-rock breeding birds under the similar temperate climatic 487 conditions.

488 Excretory behaviour of Atlantic puffins varies between individual birds and can lead 489 to variation in NH_3 emissions. The entrance chambers of most puffin burrows are free 490 from guano, with chicks deeper in the nest excreting inside the burrow, but adults do 491 not excrete in the burrow (M. Newell, pers. comm.). A significant fraction of the NH³ 492 emitted from subterranean excreta can therefore be expected to be absorbed by 493 overlying soil and vegetation. The amount of puffin excretion on the land surface 494 changes during the day as well as between days, puffins can be observed in large 495 numbers across the colony, often at dusk and less so at dawn (Harris & Wanless, 496 2011).

497 In earlier modelling estimates, the presence of substantial amounts of vegetation has 498 been estimated to reduce NH_3 by a multiplier of 0.2 (Wilson et al., 2004), while NH_3
499 emissions from excretion inside burrows was estimated to be 0.1 of that on bare rock. emissions from excretion inside burrows was estimated to be 0.1 of that on bare rock. 500 Based on the *Pv* values presented in Table 2, the present measurements in the Firth of 501 Forth indicate 0.14 or 0.31 times lower emissions for Puffins (grass and burrows) 502 compared with Northern Gannets or Guillemots, respectively (which are both bare 503 rock breeders) which are broadly consistent with the prior model estimates.

504 **4.2 NH3 Emissions and environmental conditions**

 505 The NH₃ emission estimates from the on-line measurements offer the possibility to 506 compare and interpret emission rates with environmental parameters during the course 507 of the measurement campaigns. This is illustrated for the Isle of May and Bird Island 508 in the present study and for Ascension Island (Riddick et al., 2014), based on a 509 comparison of hourly emission estimates to each environmental variable (ground

510 temperature, relative humidity, wind speed and precipitation) at each site 511 (Supplementary Material Section 10).

512 The results show ground temperature is positively correlated to measured NH₃ 513 emission at each site, representing tropical, temperate and sub-polar climates. The 514 strongest correlation with temperature was found at the Isle of May $(R=0.7; P<0.001)$. 515 Conversely, the weakest correlation between ground temperature and NH₃ emissions 516 was found for Ascension Island ($R=0.2$; P<0.001), which appears to have been due to 517 the overriding importance of moisture-limitation on the temporal pattern of emissions 518 at this site (Riddick et al., 2014). This is illustrated by a higher correlation between 519 NH₃ emission and relative humidity $(R = 0.4; P<0.001)$ and NH₃ emission and 520 precipitation events $(R = 0.3; P<0.001)$ at Ascension Island. In fact, Ascension is the 521 only field site where there is a positive correlation between NH₃ emission and both 522 relative humidity and precipitation, whereas relative humidity is inversely correlated 523 to emission at the Isle of May and Bird Island. This indicates that, where there is 524 sufficient water availability for uric acid hydrolysis (as at Bird Island and the Isle of 525 May), excess water tends to suppress the measured $NH₃$ emission.

526 Wind speed has a positive correlation with emission at all sites, with this correlation being strongest in the sub-polar conditions of Bird Island ($R = 0.9$; P<0.001) and being strongest in the sub-polar conditions of Bird Island ($R = 0.9$; P<0.001) and 528 weakest in the tropical conditions of Ascension Island $(R = 0.1; P = 0.09)$. This may 529 reflect the fact that Bird Island is the windiest site $(2 - 18 \text{ m s}^{-1})$ with the smallest 530 moisture limitation and temperature variation, so that turbulence is the major 531 controller of hourly variation in NH_3 emissions. By contrast, wind speeds were lower 532 at Ascension Island, so that the effect of varying moisture limitation largely masked 533 the effect of wind speed.

princing importance of moisture-limitation on the temporal pattern of emissions is
it (Riddick et al., 2014). This is illustrated by a higher correlation between
mission and relative humidity (R = 0.4; P-0.001) and NH₃ 534 It was assumed that the pH at each site remained constant throughout. No direct 535 measurements of pH were taken because of access restrictions to the breeding sites 536 and changes in pH of the guano may explain some of the variance in results. 537 Supplementary Material Section 11 shows there is some correlation between soil pH 538 and P_v ($R^2 = 0.40$, number of points = 11, p-value 0.04). Supplementary Material 539 Section 11 also shows that there is also a negative correlation between seabirds' food 540 energy to nitrogen ratio ($R^2 = 0.61$, number of points = 11, p-value 0.004). The 541 energy to nitrogen ratio is significantly correlated to P_v , but that the response is very 542 weak as the ratio only goes from 167 to 189, ie around 10% variation, so cannot 543 propagate much to other estimates, and may simply reflect input uncertainty in the 544 dataset. The sample size of species and diet is very small and further investigation is 545 required to ensure this is not correlated solely with temperature.

546 **4.3 Comparison of Active and Passive sampling methods**

547 The comparison summarized in Table 1 shows that the approach of calculating time-548 averaged NH3 fluxes from ALPHA samplers provided surprisingly similar estimates 549 to those calculated from on-line sampling with 15 minute averaging. This finding is 550 consistent with a similar comparison by Riddick et al. (2014) for tropical colonies, 551 and by Theobald et al. (2013) for measurements on mainland Antarctica. In principle, 552 while co-variance between NH₃ concentrations and varying atmospheric turbulence is 553 expected to lead to significant errors, these comparisons show that the errors 554 associated with this can be relatively modest in practice. While this finding may be a 555 surprise to micrometeorologists, it appears to result from the fact that non-linearities 556 associated with averaging over periods of changing atmospheric stability are

557 relatively modest when compared with other sources of uncertainty, especially for 558 such sites at relatively windy locations.

559 By calculating the flux using the on-line NH3 sampling, but with the time resolution 560 of the ALPHA samplers, we can also compare the chemical and meteorological 561 sources of uncertainty. In this way, Table 1 shows that the Uncertainty associated 562 with the Sampling Period (USP) is of comparable magnitude to the Uncertainty 563 associated with the chemical Sampling Method (USM). This study therefore further 564 provides support for the utility of low-cost passive sampling measurements at remote 565 locations where it is often logistically much harder to deploy expensive active 566 sampling methods. While such passive NH₃ flux measurements cannot replace 567 continuous measurements for the examination of detailed (e.g. hourly) temporal 568 controls on emissions (Supplementary Material Section 10), they may serve a useful 569 role in gathering data over longer periods (e.g. 2-weekly measurements over several 570 years) for comparison of seabird colonies in different climates.

571 **5. Conclusions**

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sins where it is often logistically much harder to deploy expensive acity
on measurements control ephacy much harder to deploy expensive acity
 572 The analysis shows that each of the environmental variables investigated have an 573 influence on NH₃ emission (ground temperature, relative humidity, precipitation, 574 wind speed). Increases in NH₃ emission caused by increases in relative humidity and 575 rain events were only observed at the arid Ascension Island field site, where lack of 576 moisture appeared to limit rates of uric acid hydrolysis. At other sites in colder 577 climates, increases in precipitation result in decreased NH₃ emission, because rain 578 events dilute available ammonium pools, while having the potential to wash uric acid 579 and NH₃ from the surface. Ammonia emission was found to increase with wind speed 580 especially at the cooler sites, reflecting a reduction in both aerodynamic and boundary 581 layer resistances at higher wind speeds. Overall, the most consistent relationship is 582 the increase in NH₃ emission with increasing ground temperature.

583 Future work will examine these mechanisms more explicitly using a mechanistic 584 model (Blackall, 2004; Riddick, 2012), allowing the observed relationships between 585 environmental conditions and NH3 emission to be better understood, as well as 586 providing a basis for simulating the effect of future climate change scenarios on 587 global NH₃ emissions from seabird colonies.

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597 **References**

- 598 Blackall, T.D. (2004) The emissions of ammonia from seabird colonies. PhD thesis, 599 University of Leeds.
- 600 Blackall, T.D., Theobald, M.R., Milford, C., Hargreaves, K.J., Nemitz, E., Wilson, 601 L.J., Bull, J., Bacon, P.J., Hamer, K.C., Wanless, S. and Sutton, M.A. (2004)
- 602 Application of tracer ratio and inverse dispersion methods with boat-based plume
- 603 measurements to estimate ammonia emissions from seabird colonies. Water, Air, & 604 Soil Pollution: Focus, 4, 279-285.
- 605 Blackall, T.D., Wilson, L.J., Theobald, M.R., Milford, C., Nemitz, E., Bull, J., Bacon,
- 606 P.J., Hamer, K.C., Wanless, S. and Sutton, M.A. (2007) Ammonia emissions from 607 seabird colonies. Geophysical Research Letters, 34, 5-17.

608 Cape, J.N., van der Eerden, L.J., Sheppard, L.J., Leith, I.D. and Sutton, M.A. (2009) 609 Evidence for changing the Critical Level for ammonia. Environmental Pollution, 157, 610 1033-1037.

- 611 Cowen, K., Summer, A. L., Dindal, A., Riggs, K., Willienberg, Z., Hatfield, J. L.,
- 612 Pfeffer, R. and Scoggin, K. (2004) Environmental Technology Verification Report.
- 613 Pranalytica, Inc. Nitrolux 1000 Ambient NH3 Analyser. ETV, pp.57.
- 614 Crittenden, P.D., Scrimgeour, C.S., Minnullina, G., Sutton, M.A., Tang Y.S. and
- 615 Theobald, M.R. (2015) Lichen response to ammonia deposition defines the footprint
- 616 of a penguin rookery Biogeochemistry. 122(2), 295-311.
- 617 Elliott, H. A. and Collins, N. E. (1982) Factors affecting ammonia release in broiler 618 houses. Transactions of the Asae, 25, 413-418.
- 619 Flesch, T.K., Harper, L.A., Powell, J.M., and Wilson, J.D. (2009) Inverse dispersion 620 calculation of ammonia emissions from Wisconsin dairy farms. Transactions of the
- 621 American Society of Agricultural and Biological Engineers. 52:253-265
- 622 Flesch, T.K., Wilson, J.D. and Yee, E. (1995) Backward-time Lagrangian stochastic 623 dispersion models, and their application to estimate gaseous emissions. Journal of 624 Applied Meteorology, 34, 1320-1332.
- 625 Fowler, D., Coyle, M., Skiba, U., Sutton, M. A., Cape, J. N., Reis, S., Sheppard, L. J.,
- 626 Jenkins, A., Grizzetti, B. and J. N. Galloway (2013) The global nitrogen cycle in the
- 627 twenty-first century. Philos. Trans. R. Soc. Lond. B. Biol. Sci., 368:20130164.
- Coverage and Color and Externor animomal. Environmental Contactor, 1973

(1877), K., Summer, A. L., Dindal, A., Riggs, K., Willienberg, Z., Hatfield, J. L., R. and Scoggin, K. (2004) Environmental Technology Verfication Re 628 Furness, R. W. (1991) The occurrence of burrow-nesting among birds and its 629 influence on soil fertility and stability. Symposia of the Zoological Society of London, 630 63, 53-67.
- 631 Gras, J. L. (1983) Ammonia and Ammonium Concentrations in the Antarctic 632 Atmosphere. Atmospheric Environment, 17, 815-818.
- 633 Harris, M.P., Newell, M., Leitch, A., Bruce, B. and Hunt, J. (2009) Dramatic decline 634 in numbers of Atlantic Puffins in the Firth of Forth. Scottish Birds 29: 132-134.
- 635 Harris, M.P. & Wanless, S. (2011) The Puffin. Poyser. pp 133-138.
- 636 Harper, L. A., Flesch, T. K. and Wilson, J. D. (2010) Ammonia emissions from 637 broiler production in the San Joaquin Valley. Poultry Science, 889, 1802-1814.
- 638 Laubach, J., Kelliher, F. M., Knight, T. W., Clark, H., Molano, G. and Cavanagh, A.
- 639 (2008) Methane emissions from beef cattle a comparison of paddock-and animal-640 scale measurements. Australian Journal of Experimental Agriculture, 48, 132-137.
- 641 Nemitz, E., Milford, C. and Sutton, M.A. (2001) A two-layer canopy compensation 642 point model for describing bi-directional biosphere-atmosphere exchange of 643 ammonia. Quarterly Journal of the Royal Meteorological Society, 127, 815-833.
- 644 Norman, M., Spirig, C., Wolff, V., Trebs, I., Flechard, C., Wisthaler, A., 645 Schnitzhofer, R., Hansel, A. and Neftel, A. (2009) Intercomparison of ammonia
- 646 measurement techniques at an intensively managed grassland site (Oensingen, 647 Switzerland). Atmospheric Chemistry, 9, 2635-2645.
- 648 Riddick (2012) Global ammonia emissions from seabird colonies. PhD thesis, King's 649 College, London.
- 650 Riddick S., Dragosits U., Blackall T., Daunt F., Wanless S. and Sutton M.A. (2012) 651 The global distribution of ammonia emissions from seabird colonies. Atmospheric
- 652 Environment, 55, 312-327.
- 653 Riddick S.N., Dragosits U., Blackall T.D., Daunt F., Braban, C.F., Tang, Y.S., 654 MacFarlane, W., Taylor, S., Wanless S. and Sutton M.A. (2014) Measurement of 655 ammonia emissions from tropical seabird colonies. Atmospheric Environment, 89. 35- 656 42. 10.1016/j.atmosenv.2014.02.012
- 657 Riddick S.N., Blackall T.D., Dragosits U., Daunt F., Wanless S. Hamer, K.C. and 658 Sutton M.A. (in prep) High temporal resolution modelling of climate-dependent 659 seabird ammonia emissions: The GUANO Model
- 660 Seinfeld, J. H. and Pandis, S. N. (2006) Atmospheric Chemistry and Physics: From 661 Air Pollution to Climate Change London, John Wiley & Sons.
- 662 Sommer, S. G., Olesen, J. E. and Christensen, B. T. (1991) Effects of temperature, 663 wind-speed and air humidity on ammonia volatilization from surface applied cattle 664 slurry. Journal of Agricultural Science, 117, 91-100.
- 665 Sutton, M.A., Howard, C.M., Erisman, J.W., Billen G., Bleeker A., Grennfelt P., van 666 Grinsven H. and Grizzetti B. (Eds.) (2011) The European Nitrogen Assessment: 667 Sources, Effects and Policy Perspectives, Cambridge University Press.
- 668 Sutton, M.A., Reis, S., Billen, G., Cellier, P., Erisman, J.W., Mosier, A.R., Nemitz, 669 E., Sprent, J., van Grinsven, H., Voss, M., Beier, C. and Skiba, U. (2012) Nitrogen & 670 Global Change: Preface, Biogeosciences, 9(5), 1691-1693, doi:10.5194/bg-9-1691- 671 2012.
- mment, ao, Trasposito U. Blackall T.D., Daunt F., Braban, C.F., Tang, Y.S

rahae, W., Taylor, S., Wanless S. and Sutton M.A. (2014) Measurement can

intermissions from tropical seabird colonies. Atmospheric Environment, 89 672 Sutton, M.A., Reis, S., Riddick, S.N., Dragosits, U., Nemitz, E., Theobald, M.R., 673 Tang, Y.S., Braban, C.F., Vieno, M., Dore, A.J., Mitchell, R.F., Wanless, S., Daunt, 674 F., Fowler, D., Blackall, T.D., Milford, C., Flechard, C.R., Loubet, B., Massad, R., 675 Cellier, P., Personne, E., Coheur, P.F., Clarisse, L., Van Damme, M., Ngadi, Y., 676 Clerbaux, C., Skjoth, C.A., Geels, C., Hertel, O., Kruit, R.J.W., Pinder, R.W., Bash, 677 J.O., Walker, J.T., Simpson, D., Horvath, L., Misselbrook, T.H., Bleeker, A., 678 Dentener, F., de Vries, W., 2013. Towards a climate-dependent paradigm of ammonia 679 emission and deposition. Philosophical Transactions of the Royal Society B-680 Biological Sciences 368, 20130166.
- 681 Tang, Y.S., Cape, J.N. and Sutton, M.A. (2001) Development and types of passive 682 samplers for NH_3 and NO_x . In Proceedings of the International Symposium on 683 Passive Sampling of Gaseous Pollutants in Ecological Research. The Scientific 684 World, 1, 513-529.
- 685 Tang, Y.S., Schmale , J., Drewer, J., Riddick, S.N., Anderson, M., Coyle, M., Helfter, 686 C., Skiba, U., Blackall, T.D., Sutton, M.A., Cape, J.N., Poskitt, J., Phillips, G., James,
- 687 P., Daunt F., Wanless, S., Leeson, S., Braban C.F. Trathan, P. & Dragosits, U. (in
- 688 prep) Bird Island: Case study of Sub-Antarctic marine and biogenic influences on
- 689 ecosystem nitrogen
- 690

- 691 Theobald, M.R., Crittenden, P.D., Tang, Y.S. and Sutton, M.A. (2013) The 692 application of inverse-dispersion and gradient methods to estimate ammonia 693 emissions from antarctic penguins. Atmospheric Environment 81, 320-329. 694 http://dx.doi.org/10.1016/j.atmosenv.2013.09.009
- 695 Tratt, D.M., Buckland, K.N., Young, S.J., Johnson, P.D., Riesz, K.A., Molina, K.C. 696 (2014) Remote sensing visualization and quantification of ammonia emission from an 697 inland seabird colony. Journal of Applied Remote Sensing, 7, 073475 – 073475. 698 doi:10.1117/1.JRS.7.073475.
- 699 Wilson, L.J., Bacon, P.J., Bull, J., Dragosits, U., Blackall, T.D., Dunn, T. E., Hamer, 700 K.C., Sutton, M.A. and Wanless, S. (2004) Modelling the spatial distribution of 701 ammonia emissions from seabirds in the UK. Environmental Pollution, 131, 173-185.
- 702 Zhu, R., Sun, J., Liu, Y., Gong, Z. and Sun, L. (2011) Potential ammonia emissions
- 703 from penguin guano, ornithogenic soils and seal colony soils in coastal Antarctica:
- 704 effects of freezing-thawing cycles and selected environmental variables. Antarctic

705 Science 23:78-92

1117/LJRS.7.073475.

1, LJ., Bacon, P.J., Bull, J., Dragosits, U., Blackall, T.D., Dunn, T. E., Hame

Sutton, M.A. and Wanless, S. (2004) Modelling the spatial distribution c

itia emissions from seabirds in the UK. Enviro

Figure 1 Left pane: Location of the Isle of May off the coast of Scotland, UK (56.19 °N, 2.56 °W). Right pane: Details of the Isle of May showing the Atlantic Puffin study colony, meteorological station and the site for on-line campaign measurements of ammonia concentration.

Figure 2 Top left pane: Location of measurement site on South Georgia (54.01 °S, 38.08 °W). Bottom left pane: Location of Bird Island in relation to South Georgia. Right pane: North western Bird Island indicating locations of Big Mac Macaroni penguin colony being studied, location of passive samplers and the site of the active ammonia concentration measurements, at Fairy Point.

n colony being studied, location of passive samplers and the site of the active contation measurements, at Fairy Point.

3 Top left pane: Location of measurement stie on South Orkney Island (60.7

59 °W). Bottom Left pane: Figure 3 Top left pane: Location of measurement site on South Orkney Island (60.73 °S, 45.59 °W). Bottom Left pane: Location of Signy Island relative to the South Orkney Islands. Right pane: Details of south-eastern Signy Island showing the ammonia sampling locations (ALPHA masts) in relation to the studied nesting area of Adélie and Chinstrap penguin nests on the Gourlay Peninsula of Signy Island.

Figure 4. Time-course of measured ammonia concentrations (top), calculated NH₃ emissions (bottom) for the active sampling campaign on the Isle of May, Scotland July 2009.

Figure 5 Time-course of measured ammonia concentrations (top), calculated NH₃ emissions (bottom) for the active sampling campaign on Bird Island, South Georgia, November & December 2010.

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Table 1 Comparison of active and passive sampling. Summary of seabird colony NH₃ emissions estimated from temperate and sub-polar measurement campaigns. P_ν is the percentage of excreted nitrogen that volatilizes as $NH₃$, Ground T is the ground temperature, USP represents the uncertainty in the flux attributable to the choice of sample averaging period and USM represents the uncertainty in the flux caused by the choice of sampling method (see notes below). Colony M indicates Isle of May, colony B indicates Big Mac on Bird Island and colony S indicates Signy Island.

2 Summary of seabird colony NH₃ emissions estimated from measurement
gas at the field sites in this study as compared with other recent measurement
 P_v describes the percentage of seabird excreted nitrogen that volatili Table 2 Summary of seabird colony $NH₃$ emissions estimated from measurement campaigns at the field sites in this study as compared with other recent measurements. Column P_v describes the percentage of seabird excreted nitrogen that volatilizes as $NH₃$.

¹ Ammonia concentrations measured in the middle of the colony (Passive Measurement site, Isle of May) and 1.5 m from the ground

 2^2 Ammonia concentrations measured at 3 m from the edge of the colony (Mast 1, Bird Island) and 1 m from the ground

³ Ammonia concentrations measured in the middle of the colony (Mast 1, Signy Island) and 1 m from the ground

 4 Ammonia concentrations measured in the middle of the colony (Active Measurement site, Isle of May) and 1.26 m from the ground

⁵ Ammonia concentrations measured at 300 m from the edge of the colony (Active Measurement site, Bird Island) and 2 m from the ground

Notes:

Flux a. Flux calculated as the mean $(+/-$ uncertainty) of hourly flux estimates based on hourly meteorology and time-integrated NH₃ concentrations from passive sampling

- Flux b. Flux calculated as the mean $(+/-$ uncertainty) of available hourly flux estimates derived from application of the on-line hourly $NH₃$ measurements with hourly meteorology.
- Flux c. Flux calculated as the mean $(+/-$ uncertainty) of flux estimates calculated from the on-line NH_3 measurements based on block averaging the NH_3 concentrations to the same extended sampling periods as used for the passive sampling.

USP is calculated as flux b minus flux c, and estimates the uncertainty in flux a and c due to using time-integrated NH₃ sampling instead of continuous hourly NH₃ concentrations. USM is calculated as flux a minus flux c, and estimates the uncertainty in flux b and c due to incomplete sampling when using the on-line measurement system.

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 $^{\text{a}}$ Estimates based on data in Tratt et al. (2014) and data from Riddick et al. (2012). # mean of the estimates from active and passive sampling (Table 1).

>The effect of meteorology on NH₃ fluxes from temperate and sub-polar seabird colonies is measured. >The percentage of excreted nitrogen that volatilized was 3% at sub-polar penguin colonies. > The percentage of guano nitrogen volatilized in temperate and sub-polar environments is much smaller than in tropical contexts. > Confirms that temperature has a significant influence on the magnitude of NH³ emissions.

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