1 Reply to Reviewers:

The manuscript with tracked changes is given below. A number of important changes 2 have been made to the manuscript: (1) we have added a sensitivity study where 3 temperature is increased by 1 °C in the TAN nitrogen reactions; (2) we have added a 4 5 sensitivity study where the actual soil pH is used in the TAN nitrogen reactions instead of 6 a set pH; (3) we have added considerable discussion vis-à-vis agricultural practices. In particular we have discussed at some length animal storage facilities. The global 7 ammonia emissions estimated here are in line with other global estimates. In the new 8 9 version of the paper we more forcefully acknowledge that while important we do not 10 simulate regional ammonia emission variations due to agro-management practices; 11 however, at the same time we stress that we are able to capture the geographical and 12 temporal impact of meteorology on ammonia emissions. The approach taken here is unique in that is suitable for incorporating agricultural nitrogen and meteorological 13 14 variablity into biogeochemical models. This is something that has not been attempted in traditional global approaches in simulating ammonia emissions. In the conclusions we 15 emphasize that a hybrid approach including both a more realistic incorporation of 16 17 agricultural practices and meteorological variability is ultimately necessary.

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For the most part we have already replied to the reviewers comments in our published replies, but a few additional comments are warranted. (1) In particular, reviewer 2 inquired about units in a few of the derivations. All units are given in appendix A as well as in the text where appropriate. In particular the Henry's law coefficient is unitless in equation 11. It is derived from the Henry's Law coefficient given in Sutton et al. (1994)

24	where $NH_3(aq)=NH_3(g)*H$, where $NH_3(g)$ is in atmospheres and the Henry's law
25	coefficient H is in moles/(liter-atmospheres). However, we convert units so that NH ₃ (aq)
26	and $NH_3(g)$ are given in the same units. (2) Reviewer 2 found a misprint in our equation
27	for NH_4^+ (equation 12). This is now corrected. The equation was coded correctly in the
28	computer code. (3). As suggested by Reviewer 2 we looked for additional data in Sogaard
29	et al. (2002) with regard to ammnonia emission factors, but did not find suitable data in
30	that paper nor access to measured emission factors. However, we did track down a
31	subsequent paper by Sintermann et al. (Biogeosciences, 9, 1611-1632, 2012) in which
32	350 measurements of emission factors for ammonia emissions from manure over a recent
33	10 year period were tabulated. We selected the more recent field measurements from this
34	dataset for cattle manure and included them in Figure 2. We note here that there is a
35	discrepancy in the emission factors between these later field scale measurements and
36	early reported measurements from wind tunnels, with the earlier measurements
37	suggesting higher emission factors. A plausibility analysis suggests the earlier emissions
38	are biased high (Sintermann et al., 2012).
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47	Estimate of changes in agricultural terrestrial nitrogen pathways and ammonia
48	emissions from 1850 to present in the Community Earth System Model
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60	Abstract. Nitrogen applied to the surface of the land for agricultural purposes represents
61	a significant source of reactive nitrogen (N_r) that can be emitted as a gaseous N_r species,
62	be denitrified to atmospheric nitrogen (N2), run-off during rain events or form plant
63	useable nitrogen in the soil. To investigate the magnitude, temporal variability and
64	spatial heterogeneity of nitrogen pathways on a global scale from sources of animal
65	manure and synthetic fertilizer, we developed a mechanistic parameterization of these
66	pathways within a global terrestrial model. The, the Community Land Model (CLM).
67	In this initial version the parameterization uses aemphasizes an explicit climate
68	dependent approach whereby the relationshipswhile using highly simplified
69	representations of agricultural practices including manure management and fertilizer
70	application. The climate dependent approach explicitly simulates the relationship
71	between meteorological variables and biogeochemical processes are used to calculate the
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72	volatilization of ammonia (NH ₃), nitrification and run-off of N_r following manure or
72	volatilization of ammonia (NH ₃), nitrification and run-off of N _r following manure or
72 73	volatilization of ammonia (NH ₃), nitrification and run-off of N_r following manure or synthetic fertilizer application. For the year 2000, we approximately 125 Tg N yr ⁻¹ and 62
72 73 74	volatilization of ammonia (NH ₃), nitrification and run-off of N _r following manure or synthetic fertilizer application. For the year 2000, we approximately 125 Tg N yr ⁻¹ and 62 Tg N yr ⁻¹ is applied to the model land surface as manure and synthetic fertilizer,
72 73 74 75	volatilization of ammonia (NH ₃), nitrification and run-off of N _r following manure or synthetic fertilizer application. For the year 2000, we approximately 125 Tg N yr ⁻¹ and 62 Tg N yr ⁻¹ is applied to the model land surface as manure and synthetic fertilizer, respectively. We estimate the resulting global NH ₃ emission and N _r -emissions are 21 Tg
72 73 74 75 76	volatilization of ammonia (NH ₃), nitrification and run-off of N _r following manure or synthetic fertilizer application. For the year 2000, we approximately 125 Tg N yr ⁻¹ and 62 Tg N yr ⁻¹ is applied to the model land surface as manure and synthetic fertilizer, respectively. We estimate the resulting global NH ₃ emission and N _r -emissions are 21 Tg N yr ⁻¹ from manure (17% of manure applied) and 12 Tg N yr ⁻¹ from fertilizer (19% of
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72 73 74 75 76 77 78	volatilization of ammonia (NH ₃), nitrification and run-off of N _r following manure or synthetic fertilizer application. For the year 2000, we approximately 125 Tg N yr ⁻¹ and 62 Tg N yr ⁻¹ is applied to the model land surface as manure and synthetic fertilizer, respectively. We estimate the resulting global NH ₃ emission and N _r -emissions are 21 Tg N yr ⁻¹ from manure (17% of manure applied) and 12 Tg N yr ⁻¹ from fertilizer (19% of fertilizer applied); reactive nitrogen dissolved during rain events from manure at 21 and is calculated as 11 Tg N per year, respectively; for synthetic fertilizer we estimate the
72 73 74 75 76 77 78 79	volatilization of ammonia (NH ₃), nitrification and run-off of N _r following manure or synthetic fertilizer application. For the year 2000, we approximately 125 Tg N yr ⁻¹ and 62 Tg N yr ⁻¹ is applied to the model land surface as manure and synthetic fertilizer, respectively. We estimate the resulting global NH ₃ emission and N _r -emissions are 21 Tg N yr ⁻¹ from manure (17% of manure applied) and 12 Tg N yr ⁻¹ from fertilizer (19% of fertilizer applied); reactive nitrogen dissolved during rain events from manure at 21 and is calculated as 11 Tg N per year, respectively; for synthetic fertilizer we estimate the NH ₃ -emission and N _r -run off during rain events at 12-yr ⁻¹ from manure and 5 Tg N per
72 73 74 75 76 77 78 79 80	volatilization of ammonia (NH ₃), nitrification and run-off of N _r following manure or synthetic fertilizer application. For the year 2000, we approximately 125 Tg N yr ⁻¹ and 62 Tg N yr ⁻¹ is applied to the model land surface as manure and synthetic fertilizer, respectively. We estimate the resulting global NH ₃ emission and N _r -emissions are 21 Tg N yr ⁻¹ from manure (17% of manure applied) and 12 Tg N yr ⁻¹ from fertilizer (19% of fertilizer applied); reactive nitrogen dissolved during rain events from manure at 21 and is calculated as 11 Tg N per year, respectively; for synthetic fertilizer we estimate the NH ₃ -emission and N _r -run-off during rain events at 12 yr ⁻¹ from manure and 5 Tg N per year, respectivelyyr ⁻¹ from fertilizer. The parameterization was implemented in the

83 to the soil nitrogen pools. In a transient simulation which predicted that, even though absolute values of from 1850 to 2000 all nitrogen pathways are increasing with 84 increased increase in magnitude as manure and synthetic fertilizer application, partitioning 85 increase. Partitioning of applied nitrogen in manure to NH₃ emissions from manure is 86 increasing on a percentage basis, increases from 14 % of nitrogen applied (3 Tg NH₃ yr⁻¹) 87 in 1850 to $\frac{1817\%}{12}$ of nitrogen applied in 2000 ($\frac{22 \text{ Tg NH}_3 \text{ yr}^4}{21 \text{ Tg NH}_3 \text{ yr}^{-1}}$). Under 88 current manure and synthetic fertilizer application rates we find a global sensitivity of an 89 additional 1 Tg of NH₃ (approximately 3% of manure and fertilizer) emitted per year °C 90 91 of warming. While the model confirms earlier estimates of nitrogen fluxes made in a 92 range of studies, its key purpose is to provide a theoretical framework that can be employed within a biogeochemical model, that can explicitly respond to climate and that 93 94 can evolve and improve with further observation observations and characterizations of agricultural practices. 95

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97 **1. Introduction**

Nitrogen is needed by all living things for growth. However, it is relatively inert in its most abundant form, diatomic nitrogen (N_2), and needs to be converted to a form of reactive nitrogen (N_r) before it can be used by most plants for growth [Visek, 1984]. Supplying sufficient N_r for maximum crop yield is a major concern in agriculture. In preindustrial times N_r demand was partly solved with the use of animal manure and seabird guano as well as crop rotation and the use of nitrogen fixing crops [Smil, 2000]. However, by the early 20th century the supply of these N_r sources could not match the 105 demands of an increasing population and a process of creating synthetic N_r was 106 developed: the Haber-Bosch process [Galloway et al., 2004].

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The use of N_r to improve crop yield has recently become an environmental concern as N_r 108 109 in synthetic fertilizer and manure cascades through the soil, water and the atmospheric 110 nitrogen cycles. Plants can readily use applied N_r for plant growth; however, N_r washed 111 off fields or volatilized as gas can reduce ecosystem biodiversity through acidification and eutrophication [Sutton et al., 2013]. Increased N_r in the hydrosphere can lead to the 112 113 subsequent degradation of riverine and near shore water quality as the water becomes more acidic and the growth of primary producers blooms [Turner and Rabalais, 1991; 114 115 Howarth et al., 2002], which can alter the local interspecies competition and biodiversity 116 [Sutton et al., 2012]. Reactive nitrogen emissions into the atmosphere impacts air quality through the ozone generation associated with NO emissions [e.g., Hudman et al., 2010] 117 and the contribution of ammonia (NH₃) to aerosol formation [e.g., Gu et al., 2014]. 118 119 Nitrogen cycling also impacts climate through the stimulation of plant growth and 120 associated increased carbon storage; through the associated emissions of N₂O, a strong 121 greenhouse gas; through emissions of nitrogen oxides and the associated ozone production; and through the emissions of ammonia (NH₃) with its potential to cool the 122 123 climate through aerosol formation [e.g., Adams et al., 2001].

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125 As a result of their <u>dependencydependence</u> on environmental conditions, N_r pathways 126 following manure or synthetic fertilizer application are likely to change in the future 127 under climate change scenarios. This study describes a biogeochemically consistent 128 process driven parameterization suitable for incorporation into Earth System Models that simulates N_r flow following the surface addition of N_r as manure or synthetic fertilizer. 129 The parameterization is evaluated on both the local and global scales against local 130 131 measurements and independent global NH₃ flux estimates. The calculated emission estimates for NH₃ and the N_r runoff due to manure and synthetic fertilizer application will 132 133 be used in ensuing studies in both present and future climates to investigate their impact on nitrogen cycling and climate within the earth system. To our knowledge, no Earth 134 System model has yet to explicitly predict changing nitrogen pathways from manure and 135 136 synthetic fertilizer in response to climate. We note at the outset that the representation of 137 agricultural processes is highly simplified in the initial model version described here.

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139 Sources of N_r largely fall into two categories, 'new' sources, created by chemical and biological processes, and those that are 'recycled', such as manure excretion of animals. 140 The largest natural new N_r producers are biological nitrogen fixers, found in the ocean 141 142 and, biological nitrogen fixers on land, and as the by-product of lightning estimated at 140 Tg N yr⁻¹ \pm 50%, 58 Tg N yr⁻¹ \pm 50% and 5 Tg N yr⁻¹ \pm 50%, respectively [Fowler 143 et al., 2013]. The dominant anthropogenic sources of new Nr are Haber-Bosch derived 144 fertilizer (estimated at 120 Tg N yr⁻¹ ±10 %),% in 2005), the burning of fossil fuels, (30 145 Tg N yr⁻¹ \pm 10 $\frac{10}{2000}$, and a further 60 Tg N yr⁻¹ \pm 30 % (circa 2005) estimated 146 from biological nitrogen fixers grown for human consumption, such as legumes [Fowler 147 et al., 2013]. Since pre-industrial times, anthropogenic N_r creation has increased from 15 148 Tg N yr⁻¹ to the present estimate of 210 Tg N yr⁻¹ [Galloway et al., 2004; Fowler et al., 149 150 2013]. Animal manure is used to stimulate plant growth in agriculture. It contains N_r recycled from the soil produced when animals eat plants. A comprehensive increase in livestock population is estimated to have increased global manure production from 21 Tg $N ext{ yr}^{-1}$ in 1850 to the present estimate of 141 Tg N $ext{ yr}^{-1}$ [Holland et al., 2005]. It is suggested that this increase in recycled N_r production speeds up the decay and processing of plant biomass, releasing different N_r products to the atmosphere when compared to natural decay processes [Davidson, 2009].

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Projections of agricultural activity [Bodirsky et al., 2012] suggest continued increases in the application of synthetic fertilizers until the mid-21st century (and possibly beyond) concurrent with likely increases in manure production [Tilman et al., 2001]. In addition to the increased use of organic and synthetic fertilizers <u>in the future</u>, NH₃ emissions are expected to increase because of <u>the impact of changing climate</u> on nitrogen <u>biochemistrybiogeochemistry</u> [Tilman et al., 2001; Skjoth and Geels, 2013; Sutton et al., 2013].

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Current estimates of the direct forcing of nitrate aerosols present as ammonium nitrate 166 encompass the range from -.03 Wm⁻² to -.41 Wm⁻² over thein ACCMIP (Atmospheric 167 Chemistry and Climate Model Intercomparison Project) [Shindell et al., 2013] and 168 169 AeroCom Phase II [Myhre et. al., 2013] simulations. With a future reduction in sulfate 170 emissions the relative importance of nitrate aerosols is expected to dominate the direct 171 aerosol forcing by 2100 with a resulting increase in radiative forcing of up to a factor of 8.6 over what it would have been otherwise [Hauglustaine et al., 2014]. These estimates 172 do not consider the temperature dependence of NH₃ emissions. Skjoth and Geels [2013] 173

predict increases in future NH₃ emissions of up to 60% over Europe by 2100 largely due to increased NH₃ emissions with temperature. Sutton et al. [2013] predicts future temperature increases may enhance global NH₃ emissions by up to approximately 40% assuming a 5° C warming. In addition to <u>future_changes in climate-induced_NH₃</u> volatilization from manure and synthetic fertilizer application, <u>future changes in agro-</u> <u>management practices, soil microbiological processes and nitrogen runoff_may be</u> <u>expected.</u>

Studies calculating NH₃ emission from manure and synthetic fertilizer have broadly 181 182 fallen into two categories: models that use empirically derived agriculturally-based 183 emission factors and more complex process-based models. Global emissions have almost 184 been universally estimated using the former approach with specified emission factors 185 taking into account the. animal feed, the type of animal housing if any and the field application of the synthetic fertilizer or manure [e.g., Bouwman et al., 1997]. Very 186 simplified representations of the effect of climate have been taken into account-by 187 grouping countries into industrial or developing categories [Bouwman et al., 1997]. For 188 example, this type of emission inventory was used in the Atmospheric Chemistry and 189 Climate Model Intercomparison Project (ACCMIP) [Lamarque et al., 2013] for assessing 190 191 historical and future chemistry-climate scenarios. The global impact of nitrogen on the carbon cycle as well as on atmospheric chemistry has traditionally been assessed using 192 these type of inventories of NH₃ emissions. A seasonal emission dependence is not 193 implicit in these bottom-up inventories although sometimes an empirical relationship is 194 applied [e.g., Adams et al., 2001; also see Skjøth et al., 2011]. Emission factors were 195 used by Bouwman et al. [1997] to estimate global NH₃ emissions in 1990 of 54 Tg N yr⁻¹, 196

with the greatest emission of 21.6 Tg N yr⁻¹ from domestic animals [Bouwman et al., 197 1997]. Beusen et al. [2008] also used emission factors to estimate global NH₃ emission 198 from agricultural livestock (21 Tg N yr⁻¹) and synthetic fertilizers (11 Tg N yr⁻¹) in 2000; 199 Bouwman et al. [2013] estimated emissions of 34 Tg NH₃ yr⁻¹ on agricultural land, with 200 10 Tg NH₃ yr⁻¹ from animal housing. A number of more recent global models have 201 included emission factors explicitly as a function of temperature [e.g., Huang et al., 2012; 202 Paulot et al., 2014]. Paulot et al. [2014] estimates global current NH₃ emissions of 9.4 Tg 203 yr⁻¹ for synthetic fertilizer and 24 Tg yr⁻¹ for manure. 204

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Alternatively process-based or mechanistic models have been developed that estimate N_r 206 flows, equilibria and transformations between different nitrogen species as well as 207 208 nitrogen emissions from synthetic fertilizer and manure. Process models have been used 209 on the field to regional scale, but not on the global scale. These models generally do not simulate the run-off of Nr. For example, Génermont and Cellier [1997] model the 210 211 transfer of NH₃(g) to the atmosphere after considering the physical and chemical equilibria and transfer of Nr species ($NH_3(g)$, $NH_3(aq)$, $NH_4^+(aq)$) in the soil. The 212 213 resulting model is used to calculate the NH_3 emissions from synthetic fertilizer over France within the air quality model, Chimere [Hamaoui-Laguel et al., 2014]. Other 214 examples include Pinder et al. [2004], who describes a process model of NH₃ emissions 215 216 from a dairy farm, while Li et al. [2013] describes a farm-scale process model of the 217 decomposition and emission of NH₃ from manure.

219 The overall goal of this paper is to describe and analyze a global model capable of 220 simulating nitrogen pathways from manure and synthetic fertilizer added to the surface of 221 the land under changing climactic conditions to allow a better global quantification of the 222 climate, health and environmental impacts of a changing nitrogen cycle under climate 223 change. The resulting model is of necessity designed for use within an Earth System 224 Model so as to simulate the interactions between the climate and the carbon and nitrogen cycles. Section 2 presents the overall methodology including a detailed description of the 225 226 process model developed here to calculate climate dependent nitrogen pathways. Section 227 3 analyzes this the model and includes: a comparison of simulated versus site level 228 measurements of NH₃ fluxes; an analysis of the globally heterogeneous nitrogen 229 pathways from applied manure and synthetic fertilizer over a range of climatic regimes; 230 model predictions for changes in nitrogen pathways from 1850 to present and the sensitivity of the results to model parameters. Section 4 gives our conclusions. 231

232

233 **2. Methods**

234 In this section we describe a process model designed to predict for the spatial and 235 temporal variations in the evolutionFlows of N_r-Agricultural Nitrogen (FAN) that 236 results simulates NH₃ emissions and other N_r flows from the application of applied manure and synthetic fertilizer applications, including their spatial and temporal variations, 237 238 within the context of an Earth System Model, the Community Earth System Model 1.1 239 (CESM1.1). The FAN process model developed here simulates the loss major pathways of N_r following the application of synthetic fertilizer or manure to the Earth's surface: its 240 241 incorporation of manure and fertilizer N_r into soil organic matter and soil nitrogen pools

242	[Chambers et al., 1999], theits volatilization of NH ₃ to the atmosphere and the direct
243	runoff of N_r from the surface (Figure 1). The model is global in nature, is designed to
244	conserve carbon and nitrogen and responds to changes in climate. It is designed to
245	provide an interface between the application of manure and synthetic fertilizer and the
246	nitrogen cycling developed within the Community Land Model 4.5 (CLM4.5), the land
247	component of the CESM.
248	
249	2.1 Relation between the process model and the CESM1.1
250	
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252	Nitrogen pathways subsequent to the application of manure or synthetic fertilizer depend
253	on the complex interaction between both human and natural processes. In particular they
254	depend on the biology and physics of the applied substrate, agricultural practices and
255	climate. Bottom-up emission inventories with specified emission factors that take into
256	account the animal feed, the type of animal housing if any and the field application of the
257	synthetic fertilizer or manure [e.g., Bouwman et al., 1997] are generally used in global
258	chemistry and chemistry-climate applications. For example, this type of emission
259	inventory [e.g. Lamarque et al., 2010] was used in the Atmospheric Chemistry and
260	Climate Model Intercomparison Project (ACCMIP) [Lamarque et al., 2013a] for
261	assessing historical and future chemistry-climate scenarios as well as in assessing
262	nitrogen deposition [Lamarque et al., 2013b] with implications for impacts on the carbon
263	cycle. However, these inventories include very simplified representations of the effect of
264	climate on emissions, for example, by grouping countries into industrial or developing
265	categories [Bouwman et al., 1997]. A seasonal emission dependence is not implicit in

266 <u>these bottom-up inventories although sometimes an empirical relationship is applied [e.g.,</u>
267 <u>Adams et al., 2001; also see Skjøth et al., 2011].</u>

268

In the first application of the model described here we take the opposite tact. We have
minimized the description of agricultural practices, and instead emphasize a physically
based climate dependent biogeochemistry of manure and synthetic fertilizer
decomposition and the resultant nitrogen pathways. The truth of the matter, of course, lies
somewhere in between: regional and temporal meteorological differences and changes
with climate as well as regional agro-management practices and their possible changes
impact NH₃ emissions.

276

277 We recognize that in this first application we are simplifying many important agro-278 management processes including: (1) we assume all synthetic fertilizer is urea and the pH 279 of soil is given. Different applied synthetic fertilizers have a strong impact on the pH of 280 the soil-fertilizer mixture with the overall emission factor very dependent on the pH as well as the day since application [Whitehead and Raistrick, 1990]. Urea is the most 281 282 commonly used synthetic fertilizer accounting for over 50% of the global nitrogenous synthetic fertilizer usage [Gilbert et al., 2006] and has one of the highest emission factors 283 284 for commonly used synthetic fertilizers [Bouwman et al., 1997]. Emission factors for 285 other types of fertilizers can be significantly smaller. (2) We do not account for manure management practices. Instead we assume all manure is continuously spread onto fields. 286 287 In a global study Beusen et al. [2008] considered four primary pathways for manure 288 nitrogen: (1) manure nitrogen lost from the system (14% of the manure nitrogen, range 5-

289	26%), (2) manure nitrogen excreted in animal houses followed by storage and subsequent
290	spreading onto croplands (35% of manure nitrogen; range 24%-51%), (3) manure
291	nitrogen excreted in animal houses followed by storage and subsequent spreading onto
292	pasture lands (7% of manure nitrogen; range 3%-11%), (4) manure nitrogen excreted by
293	grazing animals onto pastures (44% of manure nitrogen; range 29-59%). Of the 42% of
294	manure nitrogen excreted in housing, 20% (range: 12-28%) is emitted as NH ₃ from
295	housing and storage facilities [Beusen et al., 2008]. An additional 15-23% of the
296	remaining manure nitrogen is emitted as NH ₃ (range: 11-30%) after it is spread onto crop
297	or pasture land. Of the 44% of manure nitrogen excreted by grading animals on pasture
298	land 11-12% (range 6-17%) is emitted as ammonia. Considering these various pathways
299	the overall emission factor for manure nitrogen is estimated as 19% in Beusen et al.
300	[2008] (compare with 17% in this study). (3) We do not account for specific fertilizer
301	application techniques. For example, the soil incorporation of manure leads to a 50%
302	reduction in NH ₃ emissions compared to soil broadcasting (Bowman et al., 2002). We
303	recognize that there are large spreads in all these ranges and that regional practices may
304	alter these numbers, although large errors may be unavoidable due to insufficient
305	characterization of regional agro-management practices.
306	
307	Even though regional differences in agro-management will result in regional differences
308	in NH ₃ emissions, traditional bottom-up NH ₃ emission inventories do not account for

- 309 physically based geographical and meteorological influences, including temperature,
- 310 turbulence and rainfall. However, these are accounted for in the parameterization
- 311 described below. As with regional differences in agro-management practices,

312 meteorological impacts may also induce large regional and inter-annual variations in NH₃
313 emissions. For example, increasing the ground temperature from 290° K to 300°K (at a
314 pH of 7) increases the NH₃ emissions by a factor of 3 (section 2.2).

315

316 In the present application we also simplify the representation of NH₃ fluxes to the 317 atmosphere. The aerodynamic resistances used to compute the flux of NH₃ to the 318 atmosphere are calculated with the CLM4.5, but due to the configuration of the CLM are 319 not calculated at the plant function type (PFT) level. In addition, the canopy capture of 320 the NH_3 flux is calculated as a global number and not at the PFT level. The simulation of 321 dynamic NH₃ emissions, as described below, with NH₃ emissions responding to temperature on the model timestep, and thus allowing for a regionally resolved flux of 322 323 NH₃ dependent on diurnal fluctuations in boundary layer turbulence and boundary layer height is a first step in representing the coupling between terrestrial NH₃ fluxes with the 324 325 atmosphere. Of course high spatial heterogeneity may preclude an accurate local 326 representation of these exchange processes on the approximately 2 x 2 ° grid cell used here, but even on similar coarse resolutions Zhu et al. [2015] show the implementation of 327 328 a bidirectional scheme has significant global and pronounced regional impacts (e.g. 329 approximately a 44% decrease in NH₃ emissions over China in April).

330

A number of additional requirements are necessary to model NH₃ emissions following
 synthetic fertilizer or manure application within an Earth System Model, specifications
 that are not necessary in more traditional formulations. (1) The model must be global in
 nature to characterize global interactions between applied N_r and climate. (2) The model

335	must conserve nitrogen. In particular the nitrogen associated with manure does not add
336	new nitrogen to the system, but merely represents a recycling of available nitrogen.
337	Artificial sources or sinks of nitrogen may have serious repercussions especially when
338	simulating the global nitrogen cycle on the timescale of centuries. (3) The model must be
339	able to simulate the changing impact of climate on the fate of manure and synthetic
340	fertilizer N _r . In particular, NH ₃ emissions are sensitive to both temperature and to the
341	water content of the soil. In addition the runoff of N _r is likely to change under climate
342	change scenarios. The CESM1.1 simulates atmospheric, ocean, land and sea ice
343	processes, linked together using a coupler, and includes a land and ocean carbon cycle
344	[Hurrell et al., 2013; Lindsay et al., 2014]. The CESM participates in the Climate Model
345	Intercomparison Project (CMIP5), and has been extensively evaluated in the literature
346	[see Hurrell et al., 2013]. The relation between nitrogen cycling within the process model
347	developed here and that within the atmospheric, land and river components of the
348	Community Earth System Model (CESM1.1) is given in Figure 1. In this first study the
349	subsequent fate of N_r within these other components of the CESM1.1 is not further
350	considered. Thus, this first study does not account for the feedbacks between the applied
351	nitrogen in the synthetic fertilizer or manure pools and the carbon cycle. However future
352	studies will consider these effects. In particular, the fate of N _r -incorporated into soil
353	organic matter or the soil nitrogen pools of the Community Land Model (CLM) 4.5, the
354	land component model of the CESM1.1, is not considered (see Figure 1). In addition, the
355	fate of N ₄ -emitted into the atmosphere as NH ₂ -directly from synthetic fertilizer or manure
356	is handled by the atmospheric chemistry component of the CESM (CAM chem) and is
357	not considered here (Figure 1). Note that as a first approximation the model described

358	here does not simulate the direct emission loss of species other than NH2-Atmospheric
359	emission losses of N_2O or N_2 (and potentially $NO_{\underline{x}}$) are simulated in the Community
360	Land Model (CLM) 4.5 [Koven et al., 2013], the land component model of the CESM1.1,
361	'downstream' from the pathways explicitly considered here. The run-off of N_{F} from
362	manure or synthetic fertilizer coupled has been coupled to the river transport model
363	(RTM) [Nevison et al., 2015] (Figure 1).
364	
365	The process model developed here is capable of simulating the physics of changing
366	nitrogen pathways under a changing climate.
367	
368	An ideal model would incorporate a globally more explicit representation of agro-
369	management practices, including manure treatment (housing, storage and spreading) and
370	fertilizer application [e.g., see Sutton et al., 2013]. It would also include an explicit
371	representation of the bidirectional exchange of NH ₃ between the land and atmosphere
372	including the incorporation of PFT dependent canopy deposition and aerodynamic
373	resistances. While the model developed here captures many of the regional and global
374	features seen in models based on emission factors, here we emphasize the importance of
375	regional differences in meteorology.
376	
377 378	2.1 Relation between the FAN process model and the CESM1.1
379	The parameterization developed here acts as the interface between specified manure and
380	synthetic fertilizer application and the CESM1.1. The CESM1.1 simulates atmospheric,
381	ocean, land and sea ice processes, linked together using a coupler, and includes a land

382 and ocean carbon cycle [Hurrell et al., 2013; Lindsay et al., 2014]. The CESM participates in the Climate Model Intercomparison Project (CMIP5), and has been 383 extensively evaluated in the literature [see Hurrell et al., 2013]. described here strongly 384 385 interacts with the The land component of model within the CESM1.1, the CLM 4.5. The 386 CLM 4.5 includes representation of surface energy and water fluxes, hydrology, 387 phenology, and the carbon cycle [Lawrence et al., 2007; Oleson et al., 2008]. The CLM simulations can be forced by meteorology (as done here), or as a part of a coupled-388 carbon-climate model [Lawrence et al., 2007; Oleson et al., 2008]. The current version 389 390 of the carbon model is an improved version of the coupled-carbon-climate model used in 391 Keppel-Aleks et al. [2013], Lindsay et al., [2014] and Thornton et al., [2009]. The carbon 392 model includes a nitrogen limitation on land carbon uptake, described in Thornton et al. 393 [2007, 2009]. Further improvements have been made to the below ground carbon cycle, as well as other elements of the land model in order to improve its [e.g. Koven et al., 394 395 2013; Lawrence et al., 2012]. The impact of increases in nitrogen deposition (NO_v and 396 NH_x from fossil fuels, fires and agriculture [Lamarque et al., 2010]) have been evaluated [Thornton et al., 2007; Thornton et al., 2009] and extensively compared to observations 397 398 [e.g. Thomas et al., 2013]. The CLM4 has been extensively tested and evaluated by many studies at the global [Lawrence et al., 2007; Oleson et al., 2008; Randerson et al., 399 2009] and the site [Stoeckli et al., 2008; Randerson et al., 2009] scale. The CLM4.5 400 401 retains the basic properties of CLM4 but with improvements to better simulate: (1) water 402 and momentum fluxes at the Earth's surface; (2) carbon and nitrogen dynamics within 403 soils and (3) precipitation run-off rates [Koven et al., 2013].

405 As described in Koven et al_{τ}, [2013], the CLM4.5 simulates the basic flows of N_r within soils following the Century N model [Parton et al., 1996, 2001; Grosso et al., 2000] 406 including the processes of nitrification, denitrification, and emissions of Nr and N2 and 407 the loss of N_r from leaching and runoff. The CLM4.5 also simulates the transfer of N_r 408 409 between soils and vegetation, and the loss of Nr from fire. Sources of Nr within the 410 CLM4.5 are from biological nitrogen fixation and from surface deposition. The process model developed here adds an additional source of Nr to the CLM4.5, the addition of 411 synthetic fertilizer. It also adds an additional pathway whereby N_r is recycled: the 412 413 creation and application of manure (Figure 1).

414

415 2.2 Process model for predicting nitrogen pathways from manure or synthetic
416 fertilizer

417

The following specifications are necessary to model the nitrogen cascade following 418 419 synthetic fertilizer or manure application within an Earth System Model. The relation between nitrogen cycling within the FAN process model developed here and that within 420 421 the atmospheric, land and river components of the Community Earth System Model (CESM1.1) is given in Figure 1. In this first study the subsequent fate of Nr from 422 synthetic fertilizer or manure application as is incorporated into the soil organic matter or 423 424 the soil nitrogen pools of the CLM4.5 is not considered here (see Figure 1). As described in more detail below fertilizer and manure is not applied to particular PFTs (e.g., pasture 425 426 or grassland) within the CLM4.5. This is because soil related properties including soil 427 nitrogen are not specified at the PFT level within the CLM4.5, but instead are specified at

428	the column level that includes many PFTs. In practice we expect that the impact of this
429	contamination across PFTs will be small since the major N-application regions (central
430	US, northern India, eastern China) are not PFT-diverse but contain almost exclusively
431	crop and grass PFTs.
432	
433	Note that as a first approximation the model described here does not simulate the direct
434	emission loss of species other than NH ₃ . Atmospheric emission losses of N ₂ O or N ₂ (and
435	potentially NO _X) are simulated in the Community Land Model (CLM) 4.5 [Koven et al.,
436	2013], the land component model of the CESM1.1, 'downstream' from the pathways
437	explicitly considered here. In addition, the fate of N_r emitted into the atmosphere as NH_3
438	directly from synthetic fertilizer or manure is handled by the atmospheric chemistry
439	component of the CESM (CAM-chem) and is not considered here (Figure 1). The run-off
440	of N _r from manure or synthetic fertilizer nitrogen pools has been coupled to the river
441	transport model (RTM) in [Nevison et al., 2016] (Figure 1), but is also not considered
442	here.
443	2.2 FAN Process Model
444	(1) The model must be global in nature to characterize global interactions between
445	applied N _* and elimate. However, as detailed soil types and agricultural practices are not
446	well characterized globally a global picture necessarily sacrifices some of the regional
447	and local details. (2) The model must conserve nitrogen. In particular the nitrogen
448	associated with manure does not add new nitrogen to the system, but merely represents a
449	recycling of available nitrogen. Artificial sources or sinks of nitrogen may have serious
450	repercussions especially when simulating the global nitrogen cycle on the timescale of

451 centuries. (3) The model must be able to simulate the changing impact of climate on the
452 fate of manure and synthetic fertilizer N_r. In particular, NH₃ emissions are sensitive to
453 both temperature and to the water content of the soil. In addition the runoff of N_r is likely
454 to change under climate change scenarios. For this reason the process model developed
455 here is capable of simulating the physics of changing nitrogen pathways under a changing
456 climate.

457

Nitrogen pathways subsequent to the application of manure or synthetic fertilizer depend 458 on the complex interaction between both human and natural processes. In particular they 459 depend on the biology and physics of the applied substrate, agricultural practices and 460 elimate.-Bottom-up inventories with explicit although still incomplete incorporation of 461 462 agricultural practices through the use of emission factors tend to minimize the climate dependence of the emissions. As discussed above this type of model has seen extensive 463 use in the climate and chemical modeling communities. We take the opposite tact here. 464 465 We have minimized the description of agricultural practices, which have not been 466 sufficiently characterized on a global basis, and emphasize the biogeochemistry of manure and synthetic fertilizer decomposition and the resultant nitrogen pathways. As 467 468 shown below, this type of model captures many of the regional and global features seen 469 in models based on emission factors. The truth of the matter, of course, lies somewhere in between. An ideal model would incorporate both emission factors (temperature and wind 470 471 dependent) where appropriate (e.g., from animal housing) as well as a more physically based system simulating the physics of applied manure and synthetic fertilizer 472

473 volatilization and runoff as modified by agricultural practices [e.g., see Sutton et al.,
474 2013].

A schematic of the overall model analyzed here is given in Figure 1. All the equations 475 476 and variables used in the model have been collated and are presented in the appendix. 477 The assumptions used in constructing this model are detailed below where appropriate. 478 Sensitivity to model parameters is given in section 3.4. The nitrogen loss pathways are calculated separately for manure and synthetic fertilizer. While this model assumes that 479 synthetic fertilizer application and manure application can take place in the same 480 approximately $2 \times 2^{\circ}$ grid cell, we also assume that manure and synthetic fertilizer are 481 482 not applied in the exactly the same place. Therefore the NH₃ emissions, the nitrogen incorporation into soil pools, and the nitrogen run-off in rain water are separately 483 calculated for manure and synthetic fertilizer in each column. This means that the Total 484 Ammoniacal Nitrogen (TAN) pools (consisting of $NH_3(g)$, $NH_3(aq)$, NH_4^+) for manure 485 and synthetic fertilizer are discrete and hence the nitrogen pathways are not combined. 486

487

The application rate and geographical distribution used for manure and synthetic fertilizer 488 489 application is taken from the synthetic fertilizer application and manure production datasets developed in Potter et al [2010]. These datasets are valid for circa 2000 for 490 synthetic fertilizer and 2007 for manure [Potter et al., 2010]. Beusen et al. [2008] 491 estimates that 14% of the manure produced is lost from the agricultural system through 492 493 building materials and other uses. In this first study we do not explicitly account for the fate of this lost manure. We further assume that manure is continuously spread onto fields 494 495 by-passing the use of animal houses and storage. While most manure is excreted in 496 housing and storage systems prior to being applied in the field, the emission factors for 497 NH₂ emissions from spreading are not significantly different than from housing and storage: the emission factor for spreading onto grassland is higher and that onto cropland 498 499 is lower. As discussed above we assume that manure is continuously spread onto fields by-passing the use of animal houses and storage and is spread across all PFTs. Future 500 501 model versions will refine these initial assumptions Beusen et al., 2008]. A more sophisticated analysis could take into account differences in manure treatment, although 502 regional differences in animal housing and storage practices would make a global 503 504 analysis quite challenging.

505

To adequately model the conversion timescales of N_r input from animals to TAN, it is 506 507 necessary to separate the manure into different pools depending on the decomposition 508 timescales (sections 2.2.1 and 2.2.2 and Figure 1). A similar strategy was adopted by Li 509 et al. [2013] for manure and is commonly used in simulating litter decomposition. 510 Synthetic fertilizer N_r is added to one pool, where after it decomposes into the TAN pool (Figure 1). Once in the TAN pool N_r (1) washes off during rain events [Brouder et al., 511 512 2005]; (2) volatilizes to the atmosphere as NH₃ [Sutton et al., 1994; Nemitz et al., 2000]; 513 where after it is redeposited onto the canopy (not shown) or enters the atmosphere flow; (3) nitrifies to form nitrate (NO_3^{-}) [Stange and Neue, 2009]; 4) or is incorporated into the 514 515 soil nitrogen pools. Nitrate, in turn, becomes incorporated into the soil (Figure 1). A 516 number of other smaller loss processes are not explicitly simulated. Nitrate, in turn, becomes incorporated into the soil (Figure 1). 517

519 Manure must be added to the model in such a manner as to conserve nitrogen (Figure 1). Here, we assume animals consume carbon and nitrogen from plants and then 520 subsequently excrete this as manure. Within the CLM, carbon and nitrogen in the plant-521 522 leaf pool is thus converted to carbon and nitrogen in manure and urine, conserving 523 overall carbon and nitrogen. The conversion rate from carbon and nitrogen in plants to 524 that in manure and urine is set to equal the rate of manure and urine production. The external dataset of Potter [2010] gives the rate of N_r production from ruminantsanimals, 525 526 and thus allows us to specify the nitrogen flows. The specified C to N ratio in the plant-527 leaf pool determines the associated carbon flows due to ruminant consumption of plant 528 material. The input manure and urine production rate from ruminants animals implicitly 529 includes that produced from transported feed. Thus the subsequent NH_3 emission rate 530 includes the nitrogen contained in transported feed grown elsewhere. Here we make the simplification that the consumption rate of plant matter to balance the manure and urine 531 532 production is local. That is, we do not explicitly consider the lateral transport import of 533 animal feed to match the carbon and nitrogen flows associated with manure and urine 534 production. While this is not entirely consistent, the development of the requisite dataset 535 for feedstock flows from 1850-2000 is outside the scope of this study, although such a dataset could be developed in the future. We do not know of an Earth System Model that 536 does consider the anthropogenic lateral transportimport of nitrogen or carbon. This 537 538 inconsistency could produce cases where there is insufficient local plant material to balance the overall manure and urine production, but this is generally not the case. The 539 540 parameterization also ignores export of N_r in ruminant products such as milk and protein, 541 which could create an additional source of uncertainty.

2.2.1 Manure and Urine. Prescribed manure (including urine) is input at a constant 543 annual rate $(\alpha_{applied}(m))$ (g m⁻² s⁻¹) depending on latitude and longitude into the 544 545 manure nitrogen pools. Nitrogen applied to the land as manure (or synthetic fertilizer) is 546 assumed to be spread uniformly on each grid cell irrespective of plant functional type (pft) or surface type. Future development will spread the input into different pfts (e.g., 547 grassland or agricultural land). It is assumed that a fraction ($f_u = 0.5$) of nitrogen excreted 548 549 is urine (urea), with the remaining 50 % excreted as faecal matter [Gusman and Marino, 1999]. The excreted urine is directly added to the TAN pool (g N m⁻²). Faeces are 550 551 composed of matter with varying carbon to nitrogen ratios that take different times to 552 decompose depending on how easily they can be digested by microbes. Excreted faeces are assumed to form three different pools (g m⁻²) depending on their rate of 553 mineralization [e.g., Gusman and Marino, 1999]: (1) we assume a fraction $f_{un} = 5 \%$ is 554 excreted as unavailable nitrogen $(N_{unavailable})$, the lignin component of manure where the 555 nitrogen remains immobilized by bacteria (C:N ratio > 25:1), (2) a fraction f_r =45% goes 556 557 to the resistant pool ($N_{resistant}$) which forms the cellulose component of manure (C:N ratio c. 15:1) resistant to forming which forms TAN relatively slowly; (3) and a fraction $f_a = 50\%$ 558 goes to the available pool $(N_{available})$ that is readily available to form TAN $(N_{available})$. In 559 560 reality the fractions within each of these broadly defined pools will be dependent on the type of animal and the type of feed. 561 The equations governing the three manure pools (see Figure 1) are: 562

563 $dN_{available}/dt = f_a x \alpha_{applied}(m) - K_a \cdot N_{available} - k_m \cdot N_{available}$ (1)

564 $dN_{resistant}/dt = f_r x \alpha_{applied}(m) - K_r \cdot N_{resistant} - k_m \cdot N_{resistant}$ (2)

565 $dN_{unavailable}/dt = f_{un} x \alpha_{applied}(m) - k_m \cdot N_{unavailable}$ (3)

where $\alpha_{applied}(m)$ is the amount of manure applied (g m⁻² s⁻¹); f_a , f_r and f_{un} are the 566 fractions of manure applied to each pool; K_a and K_r (s⁻¹) are temperature dependent 567 mineralization rates and $k_m k_m (s^{-1})$ is the mechanical loss rate of nitrogen out of these 568 manure pools and into soil nitrogen pools. The decay constants, K_a and K_r are measured 569 570 as the fast and slow decomposition rates for biosolids added to various soils and incubated at 25° C [Gilmour et al., 2003], where a two-component decay model 571 accurately fit approximately 73% of the samples incubated. The decay timescales for 572 manure are 48 days and 667 days at 25 °C. The temperature dependence of the decay 573 574 constants is derived from a fit of temperature dependent mineralization rates (see appendix) [Vigil and Kissel, 1995] corresponding to a Q10 value of 3.66. To prevent the 575 576 manure pools from building up over long-timescales we assume that manure is 577 incorporated into soils with a time constant of 365 days with a mechanical rate constant k_m . This timescale is consistent with the base bioturbation rate of 1 cm² year⁻¹ assumed 578 579 in Koven et al. [2013] and a typical length scale of 1 cm. The sensitivity of the 580 subsequent nitrogen pathways to this timescale is small (section 3.4). Note, that nitrogen 581 in the $N_{unavailable}$ pool does not mineralize and is thus only incorporated into soil organic matter on the timescale determined by k_m . We assume nitrogen prior to conversion to 582 TAN comprises a range of insoluble organic compounds that do not wash away or 583 otherwise volatilize. 584

585

586

2.2.2 Synthetic fertilizer. Synthetic fertilizer nitrogen is added to the N_{fertilizer} pool

(g N m⁻²) (Figure 1) at a rate $(\alpha_{applied}(t)(f))$ (g N m⁻² s⁻¹) that depends on geography and time. The amount of nitrogen within the synthetic fertilizer pool is subsequently released into the TAN pool with the rate $k_{f}k_{f}$ (s⁻¹):

(4)

590 $dN_{fertilizer}/dt = \alpha_{applied}(f) - k_f \cdot N_{fertilizer}$

591 Here we assume all synthetic fertilizer is urea. Urea is the most commonly used synthetic 592 fertilizer accounting for over 50% of the global nitrogenous synthetic fertilizer usage [Gilbert et al., 2006]. Many other fertilizer types have significantly lower emission 593 594 factors depending largely on changes in soil pH due to interactions between the soil and 595 the fertilizer (Whitehead and Raistrick, 1990). We do not simulate this interaction here, 596 but it should be accounted for in future model development. Thus the estimates here for 597 fertilizer NH₃ emissions may be considered as an upper estimate. We set the decay timescale of urea fertilizer to be 2.4 days consistent with the decay rate measured in 598 Agehara and Warncke [2005] for temperatures from 15 to 20 ^{-o}C. In a series of 599 experiments Agehara and Warncke [2005] show that 75% of the urea hydrolyzes in a 600 week at temperatures from 10 to 25 °C without a significant dependence on temperature 601 especially for temperatures above 15 to 20 °C. 602

603

The timing of the for synthetic fertilizer application in the is determined internally within the CLM4.5 crop model coincides with as the spring planting date. This for corn. We use corn since the CLM4.5 crop model only specifically includes corn, soybean and temperate cereals and the planting date for corn lies between the earlier planting date for temperate cereal crops and the later planting of soy. The date for fertilizer application is determined for each grid point location using the surface temperature-based criteria 610 developed by Levis et al. [2012] for simulating the planting date of corn. In Levis et al. [2012]: the ten-day running mean temperature, ten-day running mean daily minimum 611 612 temperature and growing degree days must all surpass fixed threshold values (283.15K, 279.15K and 50 days, respectively, for corn) before planting can take place. We do not 613 614 use the Levis et al. [2012] crop model in this study but use these criteria to determine a planting date for each grid point and assume synthetic fertilizer is applied on this date. 615 Future applications may assume a more complete algorithm for fertilizing the spectrum of 616 global cropsFertilizer application dates can have a large influence on the seasonality of 617 618 the emissions (e.g., see Paulot et al., 2014) and the nitrogen loss pathways following 619 fertilization (section 3.4). Future applications will assume more complete algorithms for fertilizing the spectrum of crops, as well as multiple fertilizer applications and double 620 621 cropping. A global accounting of fertilization practices and application techniques (e.g., 622 fertilizer injection) nevertheless remains a considerable source of uncertainty in global 623 modeling of the NH₃ emissions from agriculture.

624

625 2.2.3 Total Ammonical Nitrogen (TAN). We consider two TAN pools (g N m⁻²), one for 626 the nitrogen produced from synthetic fertilizer $N_{TAN}(f)$ the other for nitrogen from manure 627 $N_{TAN}(m)$. The budget for the manure and synthetic fertilizer TAN pools respectively is 628 given by:

629

630
$$N_{TAN}(m)/dt = f_u x \, \alpha_{applied}(m) + K_r \cdot N_{resistant} + K_a \cdot N_{available}$$

$$631 - F_{run}(m) - K_D^{NH4} \cdot N_{TAN}(m) - F_{NH3}(m) - F_{NO3}(m)$$
(5)

632

 $N_{TAN}(f)/dt = k_f \cdot N_{fertilizer}$

633
$$-F_{run}(f) - K_D^{NH4} \cdot N_{TAN}(f) - F_{NH3}(f) - F_{NO3}(f)$$
 (6)

634

Here $F_{run}(m/f)$ (g N m⁻² s⁻¹) is the loss of nitrogen by runoff from the manure or 635 synthetic fertilizer pool, K_D^{NH4} (s⁻¹) the loss rate of nitrogen to the soil nitrogen pools, 636 $F_{NH3}(m)$ and $F_{NH3}(f)$ (g N m⁻² s⁻¹) the NH₃ emissions from the TAN pool to the 637 atmosphere from the soil manure and synthetic fertilizer pools, respectively, and $F_{NO3}(m)$ 638 and $F_{NO3}(f)$ (g N m⁻² s⁻¹) the loss of nitrogen through nitrification from the manure and 639 synthetic fertilizer pools respectively. The formulation of each of these terms is given 640 below. Inputs into $N_{TAN}(m)$ pool are from the fraction (f_u) of applied manure as urine 641 $(\alpha_{applied}(m))$, and from the decomposition of the nitrogen within the available and 642 resistant manure pools. Input into the $N_{TAN}(f)$ pool is through decomposition of 643 644 nitrogen within the synthetic fertilizer pool.

645 2.2.4 Runoff of nitrogen to rivers. The immediate runoff of fertilizer and manure 646 nitrogen to rivers is derived from the runoff rate of water (R) (m s⁻¹) in the CLM 647 multiplied by concentration of nitrogen in the TAN water pool:

648
$$F_{run}(m/f) = R \cdot \frac{N_{TAN}(m/f)}{N_{water}(m/f)}$$
(7)

The value of *R* is calculated within the CLM and is a function of precipitation, evaporation, drainage and soil saturation. The amount of water within the TAN pool $(N_{water}(m/f))(m)$ is needed to convert N_{TAN} (g N m⁻²) to a concentration (g N m⁻³). An expression for $N_{water}(m/f)$ is given in 2.2.9. Section 2.2.9. It should be emphasized that this is the immediate runoff of manure and synthetic fertilizer nitrogen from the TAN pools. Subsequent loss of nitrogen derived from manure and synthetic fertilizer application occurs following the nitrogen transfer to the soil pools, but is not tracked in
these simulations. Additional hydrological losses will also occur following NH₃
volatilization to the atmosphere, the subsequent deposition and loss through runoff or
leaching. These losses are also not tracked in the current simulation.

Initially, we attempted to use the runoff parameterization based on the global Nutrient Export from Watersheds 2 (NEWS 2) Model [Mayorga et al., 2010] where runoff is also parameterized in terms of *R*. However, the amount of nitrogen that runs off in NEWS 2 is represented in terms of the annual nitrogen initially applied to the land and thus is not directly related to the amount of nitrogen in the TAN pool.

2.2.5 Diffusion through soil. Nitrogen is assumed to diffuse from the TAN pool to the soil 664 665 pools. Génermont and Cellier [1997] represent the diffusion coefficient of ammonium through soils as dependent on soil water content, soil porosity, temperature and an 666 empirical diffusion coefficient of ammonium in free water (see appendix). For example, 667 assuming a temperature of 21° C, a soil porosity of 0.5 and a soil water content of 0.2 the 668 resulting diffusion coefficient is approximately 0.03 cm² day⁻¹, in reasonable agreement 669 670 with measurements in Canter et al. [1997]. Here we assume a typical length scale of 1.0 671 cm to convert the diffusion rate to a timescale. The resulting diffusion of ammonical nitrogen is added to pre-existing nitrogen pools in the CLM4.5- and is not further tracked. 672

673 2.2.6 Flux of Ammonia to the Atmosphere. The flux of NH₃ (F_{NH3} , g m⁻² s⁻¹) to the 674 atmosphere –is calculated from the difference between the NH₃ concentration at the 675 surface ($NH_3(g)$, g m⁻³) of the TAN pool and the free atmosphere NH₃ concentration 676 $\left(\frac{NH_{3}(a), \chi_{a_{\perp}}}{(R_{a}), \chi_{a_{\perp}}} g m^{-3}\right)$ divided by the aerodynamic (R_{a}) and boundary layer (R_{b}) resistances 677 (Equation 8) [Nemitz et al., 2000; Loubet et al., 2009, Sutton et al., 2013].

678
$$F_{NH3} = \frac{NH_3(g) - \chi_a}{R_a(z) + R_b}$$
 (8)

679

680 The calculation of $NH_3(g)$ is given below. For compatibility with the NH₃ emission model we compute average values of R_a and R_b for<u>over</u> each CLM soil column, which 681 may contain several PFTs. Continental NH3 concentrations between 0.1 and 10 $\mu g\ m^{-3}$ 682 have been reported by Zbieranowski and Aherne [2012] and Heald et al. [2012]. A 683 background atmospheric NH₃ concentration ($\chi_{a} = We$ specify χ_{a} to be 0.3 µg m⁻³ in 684 685 Equation 8) is specified, representative of aconcentrations over low activity agricultural sitesites [Zbieranowski and Aherne, 2012].-.. This concentration is intermediate between 686 concentrations at low to moderate pollution sites as diagnosed in GEOS-chem [Warner et 687 688 al., 2015]. The sensitivity to this parameter is small as NH₃(g) is usually very large-(section 3.4). While equation (8) allows for negative emissions ($NH_3(g) < \chi_a$) or 689 deposition of atmospheric NH₃ onto the soil we currently disallow negative emissions in 690 691 the current simulations. In future studies the atmospheric concentration of NH₃ will be calculated interactively when by coupling the NH₂-emission FAN model is coupled with to 692 693 the atmospheric chemistry componenent of the CESM (CAM-chem), thus allowing the dynamics of the NH₃ exchange between the soil, the atmosphere and vegetation to be 694 captured [e.g., Sutton et al., 2013]. 695

A large fraction of the NH₃ emitted to the atmosphere is assumed captured by vegetation.
The amount emitted to the atmosphere is given by:

699	$F_{NH3_{atm}}(m/f) = (1 - f_{capture}) \times F_{NH3}(m/f) (9)$
700	where <i>f_{capture}</i> is set to 0.6
701	
702	
703	where $f_{capture}$ is set to 0.6, where this accounts for the capture of the emitted NH ₃ by
704	plants or even onto the soil surface. Plant recapture of emitted NH ₃ is often reported to be
705	as high as 75 % (Harper et al., 2000; Nemitz et al., 2000; Walker et al. 2006; Denmead et
706	al., 2008; Bash et al., 2010). Using seabird nitrogen on different substrates (rock, sand,
707	soil and vegetation) inside a chamber Riddick (2012) found NH ₃ recapture to be 0% on
708	rock, 32% on sand, 59% on soil and 73% on vegetation. We set $f_{capture}$ to 0.6 in-line
709	with the findings of Wilson et al. [2004] as a mid-way between the value for soil (when
710	the crops are planted) to when they are fully grown. Bouwman et al. [1997] also used
711	canopy capture to estimate emissions with the captured fraction ranging from 0.8 in
712	tropical rain forests to 0.5 in other forests to 0.2 for all other vegetation types including
713	grasslands and shrubs. Bouwman et al. [1997] omitted canopy capture over arable lands
714	and intensively used grasslands. Overall, the deposition of NH_3 onto the canopy (or even
715	the soil surface) is poorly constrained [e.g., see Erisman and Draaijers, 1995] and often
716	ignored in model simulations. In reality canopy capture is not constant but depends on
717	surface characteristics and boundary layer meteorology. Variations in canopy capture will
718	induce temporal and regional variations in NH ₃ emissions. Explicitly including the
719	canopy capture fraction allows us to explicitly differentiate between different
720	biogeochemical pathways in future studies. In the future when the model is fully coupled
721	with the atmospheric NH ₃ cycle a compensation point approach would be desirable for

723724It is assumed that the nitrogen in the TAN pool is in equilibrium between $g \cdot Nm^3 gN$ 725 $m^3 N H_g(g), N H_g(aq) and N H_4^+(aq) \cdot NH_3(g) (g m^{-3}), NH_3(aq) (g N m^{-3}) and NH_4^+(aq) (g$ 726N m^{-3}). The equilibrium that governs the speciation of these species is determined by the727Henry's Law coefficient (K_H), where K_H is a measure of the solubility of NH₃ in water,728and the disassociation constant of NH₄⁺ in water (K_{NH_4}) (moles l⁻¹) [e.g., Sutton et al.,7291994]

calculating the net NH₃ emissions, but we feel it is outside the scope of the present study.

722

730
$$NH_3(g) + H_2O \stackrel{\kappa_H}{\leftrightarrow} NH_3(aq)$$
 (10)

731
$$NH_4^+(aq) \xrightarrow{K_{NH_4}} NH_3(aq) + H^+$$
 (11)

Combining these two expressions $NH_3(g)$ can be expressed as a function of the total TAN (e.g., Pinder et al. [2004], although note their different units for K_H and K_{NH4})

$$NH_{3}(g)(m/f) = \frac{N_{TAN}(m/f)/N_{water}(m/f)}{1+K_{H}+K_{H}[H^{+}]/K_{NH4}}$$
734 (11)
735

736

where $[H^+]$ is the hydrogen ion concentration in moles/liter. Both $K_{\rm H}K_{\rm H}$ and $K_{\rm NH4}K_{\rm NH4}$ 737 are temperature dependent. As temperature and pH increase the concentration of NH₃(g) 738 739 increases. The pH of the solution depends on the type of soil, the exposure of the manure 740 to air and may change with the aging of the manure or synthetic fertilizer TAN pool. In 741 Eghball et al. [2000] the majority of the reported measurements of pH for beef cattle 742 feedlot manure are between 7 and 8, although in one case a pH of 8.8 was measured. The 743 recommended pH for various crops ranges from approximately 5.8 to 7.0 depending on the crop (e.g., http://onondaga.cce.cornell.edu/resources/soil-ph-for-field-crops). 744 For now we simply set the pH of the solution to 7 for both the synthetic fertilizer and manureTAN pools. Sensitivity to pH is explored in section 3.4.

747

2.2.7 Conversion of TAN to NO_3^- . The flux from the TAN pool to NO_3^- by nitrification 748 $(N_{NO3}F_{NO3-}, g m^{-2} s^{-1})$ was adapted from that derived by Stange & Neue [2009] to describe 749 the gross nitrification rates in response to fertilization of a surface with manure or 750 synthetic fertilizer. In particular Stange & Neue [2009] fit measured gross nitrification 751 rates to an expression using a maximal nitrification rate r_{max} , (µg N kg⁻¹ h⁻¹) modified by 752 a soil temperature response function (f(T)) and a soil moisture response function (f(M))753 [Stange and Neue, 2009] (see appendix). However, since r_{max} is fit from their 754 experimental data the dependence of the nitrification rate on the ammonium 755 756 concentration is not explicitly included in the formulation of Stange & Neue [2009]. We have remedied this by setting the maximum nitrification rate (r_{max}) in the formulation of 757 [Stange and Neue, 2009] to $1.16 \ 10^{-6} \ s^{-1}$ consistent with the formulation in Parton et al. 758 [2001]: 759

760
$$F_{NO3}(m/f) = \frac{2 r_{max} N_{water}(m/f) N H_3(g)(m/f) K_H[H^+] / K_{NH4}}{\frac{1}{f(T)} + \frac{1}{f(M)}}$$
(12)

761

where f(T) and f(M) are functions of soil temperature and moisture and the ammonium concentration is assumed to be in equilibrium with the other forms of ammoniacal nitrogen and is thus expressed in terms of pH, K_H and K_{NH4} and N_{TAN} (NH₃(g) (m/f).).

$$dN_{NO3}(m/f)/dt = F_{NO3}(m/f) - K_D^{NO3}N_{NO3}(m/f)$$
(13)

The source of nitrate ions is nitrification from the TAN pool (see Eq. 13). Nitrate is lost 766 767 to the soil nitrate pool through diffusion. Nitrate leaching is not explicitly taken into 768 account in the current model as the diffusion of nitrate into the soil pools occurs very rapidly. The loss of nitrate through runoff and leaching can, however, occur within the 769 CLM. NO₃, but is not tracked in the current simulations. Nitrate ions diffuse significantly 770 faster than the NH₄⁺ ions because they are not subject to immobilization by negatively 771 charged soil particles [Mitsch and Gosselink, 2007]. Diffusion rates used in this study 772 773 are derived from the same formulation as assumed for the diffusion of ammonium [e.g., see Jury et al., 1983] with a different base diffusion rate. The summary of measurements 774 775 given in Canter et al. [1997], where both the diffusion of ammonium and nitrate were measured in the same soil types and wetness, suggest the base diffusion rate of NO_3^{-1} is 13 776 times faster than that of ammonium. 777

778

2.2.9 TAN and Manure Water pools. The evolution of the TAN manure and synthetic
fertilizer water pools depends on the water added during manure or synthetic fertilizer
application and the subsequent evolution of the water in the pools. The equations for the
manure and synthetic fertilizer water are:

$$dN_{water}(m)/dt = s_w(m) \times \alpha_{applied}(m) - k_{relax} \times (N_{water}(m) - M_{water})$$
(12)

$$dN_{water}(f)/dt = S_w(f) \times \alpha_{applied}(f) - k_{relax} \times (N_{water}(f) - M_{water})$$
(13)

783 These equations include a source of water $(s_w(m) \text{ or } S_w(f))$ added as a fraction of the synthetic fertilizer or manure applied and a relaxation term (k_{relax}, s^{-1}) to the soil water 784 (M_{water}, m) calculated in the CLM for the top 5 cm of soil. The value for M_{water} 785 explicitly takes into account the modification of the water pool due to rainfall, 786 787 evaporation and the diffusion of water into deeper soil layers. We assume the TAN pool equilibrates with water within the top 5 cm of the soil with a rate of 3 days⁻¹. The solution 788 is insensitive to this parameter within the ranges examined of 1 to 10 days⁻¹ (section 3.5). 789 790 The water content of manure applied to fields depends on the animal, its feedstock and on agricultural practices. Here we assume cattle manure is added as a slurry with a dry 791 fraction of 74.23 g kg⁻¹ and a nitrogen content of 1.63 g kg⁻¹, resulting in 5.67 10^{-4} m 792 793 water applied per gram of manure nitrogen applied [Sommer and Hutchings, 2001]. In 794 the case of synthetic fertilizer we assume urea is added as a liquid spread, where water 795 added is calculated from the temperature dependent solubility of urea in water [UNIDO 796 and FIDC, 1998].

797

798 **2.3 Model spin up and forcing**

Two different type of model simulations were conducted using the CLM4.5: a present day control simulation (1990-2004) and a historical simulation (1850-2000). The resolution used in these simulations is: 1.9 degrees latitude by 2.5 degrees longitude.

802

2.3.1 Present day control simulation. This simulation uses the manure and synthetic
fertilizer input as given in Potter et al. [2010]. Forcing at the atmospheric boundary is set
to the Qian et al. [2006] reanalysis for solar input, precipitation, temperature, wind and

specific humidity. The simulation is run for fifteen model years (1990-2004) with the last ten years of the simulation used for analysis. The spinup period allows for the more decomposition resistant N pools to approach a steady state with respect to the loss from mechanical incorporation into the soil.

810

811 2.3.2 Historical simulation. The historical simulation uses transient forcing conditions (accounting for changes in atmospheric CO₂, nitrogen deposition, aerosol deposition and 812 813 land use change forcings) and the Qian et al. [2006] atmospheric forcing dataset. Quality 814 meteorological 6-hourly meteorological datasets for the period prior to 1948 do not exist. 815 Therefore from 1850 to 1973 the CLM4.5 is driven by recycled meteorological data, using meteorological data from the 1948-1973 time period. During this time there is little 816 817 increase in temperature: the statistically significant changes in temperature (outside of 818 natural variability) occur after 1973. After 1973 the meteorological data is not recycled 819 but is valid for the year applied.

820

821 The temporal distribution of manure and synthetic fertilizer application from 1850-2000 822 is specified by applying the temporal distribution of Holland et al. [2005] to the base 823 values as calculated in Potter et al. [2010]. For lack of detailed information on the geography of historical manure and synthetic fertilizer we use the scaled spatial 824 distribution from Potter et al. [2010]. We assume manure production has changed from 825 26.3 Tg N yr⁻¹ in 1860 to 138.4125 Tg N yr⁻¹ in 2000 [Holland et al., 2005; Potter et al., 826 2010]-, but acknowledge these temporal changes are uncertain. Synthetic fertilizer was 827 first used in the 1920s -with use increasing to $\frac{8662}{10}$ Tg N yr⁻¹ in 2000. 828

829

830 **3. Results**

831 3.1 Model evaluation

832 To evaluate model output, measurements of the percentage of applied nitrogen that was 833 emitted as NH₃ (P_{ν}) from literature were compared against corresponding model 834 predictions. The model predictions are obtained from the present day control simulation. 835 The percent-volatilized ammoniaNH₃ was used as a metric because it can be compared across time irrespective of the absolute amount of nitrogen applied to the surface. To be 836 837 able to compare emissions to published measurements we require field studies with 838 published data on: nitrogen excretion rates, NH₃ emissions, ground temperature, location, 839 and date of measurement. Given all of these requirements we found that only a small 840 selection of publications had enough data.

841

842 For the manure emissions, 35 measurements in a range of climates (temperatures from 843 1.4 °C to 28 °C) and a range of livestock management methods (commercial beef cattle 844 feedyard, dairy cow grazing on ryegrass, beef cattle grazing on ryegrass and dairy cattle 845 grazing on pasture land) were used (Supplementary Table 1). Each P_{ν} reported by the measurement campaign was compared against the P_{ν} at the corresponding grid cell in the 846 847 model. For the synthetic fertilizer scenario, 10 measurements in a range of latitudes 848 (43 °S to 50 °N) over a range of land use surfaces (pasture, sown crops, turf and forest) 849 were used (Supplementary Table 2). Each total annual P_{ν} reported by the measurement 850 campaign was compared against the annual P_{ν} of the corresponding grid cell.

852 3.1.1 Nitrogen volatilized as NH_3 from manure. There is a general increase in the percentage of applied manure lost as NH₂ (P_{ν}) with temperature, in both the model and 853 854 measurements (Figure 2). However, temperature is not the only factor in determining 855 NH₃ emissions where wind speed, water availability and below ground soil properties can also effect NH₃ emission. This is particularly demonstrated by the measurements of 856 Todd et al. [2007] at temperatures less than 5° C where the measured emissions are 857 higher than those predicted at higher temperatures [e.g., Bussink, 1992]. It is also worth 858 noting that the model predicts the emissions of Todd et al [2007] at lower temperatures 859 860 with relative success.

861

The agreement between measured and modeled P_{ν} from manure appears reasonable, with 862 an R^2 of 0.78 that is significant at the 99.9% confidence level (p-value - 1.87 x 10⁻¹⁶). On 863 closer inspection, the model appears to agree best with measurements made on grassland 864 and differs considerably with measurements made by both campaigns for beef cattle 865 866 feedlots in Texas, where beef cattle feedlots are commercial operations to prepare livestock for slaughter and comprise of thousands of animals contained in a pen [US EPA, 867 2010]. This is perhaps not surprising, as the parameterization developed here explicitly 868 represents emissions from manure spreading as opposed to the more managed conditions 869 in feedlots. 870

871

872 *3.1.2 Nitrogen volatilized as NH*₃ *from synthetic fertilizer*. The comparison between 873 measured and modeled annual average P_v from synthetic fertilizer applied to a range of 874 land use types appears weak with an R² of 0.2 that is significant at the 90% confidence level (p-value - 0.15) (Figure 3). The lowest emissions in the model and measurements tend to be associated with the higher latitudes of both hemispheres. There does not appear to be any noticeable bias with land use type where the model estimates are both higher and lower than measured values of P_v for surfaces covered in turf, pasture land and crops. The fact that the R² for the synthetic fertilizer measurements is lower than the R² of the manure measurements is potentially caused by the single application date applied in the model, where actual farming practices may differ from model assumptions.

882

883 3.1.3 Nitrogen run-off. Here we simulate the direct nitrogen runoff from the manure and 884 synthetic fertilzer TAN pools, but do not track the resulting nitrogen flows. These flows are tracked, however, in Nevison et al. [2015] routes2016] where the nitrogen runoff 885 886 from manure and synthetic fertilizer usingpools is routed into the River Transport Model (RTM) [Dai and Trenberth, 2001; Branstetter and Erickson, 2003] within the CESM. 887 Nevison et al. [20152016] assumes denitrification occurs within the simulated rivers at a 888 889 rate inversely proportional to the river depth (amounting to approximately 30% of the 890 nitrogen inputs on average) and compares the simulated dissolved inorganic nitrogen 891 (DIN) export at the river mouths against the measured nitrogen export measurements 892 [Van Drecht et al., 2003] partitioned into the proportion that is DIN (Dissolved Inorganic 893 Nitrogen) following Global NEWS [Mayorga et al., 2010]. The simulated nitrogenDIN 894 export is nearly unbiased for six identified rivers with high human impact: the Columbia, Danube, Mississippi, Rhine, Saint Lawrence and Uruguay. Explicit comparisons against 895 the Mississippi River show that the amplitude and seasonality of the simulated Nr runoff 896 897 is in reasonable agreement with the measurements. While the comparison in Nevison et al

898 (2015). [2016] gives confidence the runoff is reasonably simulated, the complications in
899 simulating river runoff preclude tight model constraints.

900

901 **3.2 Global Nitrogen Pathways: Present Day**

902 3.2.1 Geography of Nitrogen Inputs. Global maps of nitrogen input from synthetic 903 fertilizer and manure application during the present-day simulation are given in Potter et al. [2010] and are not repeated here. Heavy synthetic fertilizer use generally occurs in the 904 upper Midwest of the U.S. (mostly east of 100° W and north of 40° N), Western Europe 905 (mostly west of 20° E and north of 40° N), the Northern part of India and much of 906 907 Northeastern and North Central China. High manure usage coincides with the areas of heavy synthetic fertilizer use but is more widespread extending across much of Eastern 908 South America from 20-40° S and across Africa at approximately 10° N. 909

910

3.2.2 Geography of Nitrogen Losses. There are strong geographical differences in the loss 911 912 pathways of nitrogen following manure or synthetic fertilizer application. The importance 913 of the various loss pathways from the TAN pool (the amount nitrogen volatilized as NH₃, 914 runoff, nitrified or diffused directly into the soil, Figures 4-8) is dependent on temperature, precipitation and soil moisture. In hot, arid climates, the percentage 915 916 volatilized is high (Figures 4 and 5). For example, regions of high NH₃ volatilization of 917 applied manure N_r approach 50% across the southwest U.S. and Mexico, Eastern South America, central and southern Africa, parts of Australia, and across southern Asia from 918 919 India to Turkey (Figure 5). The absolute highest emissions of NH_3 from applied synthetic fertilizer and from applied manure approach 20 kg N ha⁻¹ yr⁻¹ over hot regions with high 920

921 applications, e.g. the Indian peninsula and parts of China (Figure 4 and 5). Ammonia emissions from manure are more broadly distributed globally than those of synthetic 922 923 fertilizer with high NH₃ emissions not only over the synthetic fertilizer hotspots, 924 characterized by heavy application of both synthetic fertilizer and manure, but also over 925 southeastern South America and central Africa. For the most part, the largest synthetic 926 fertilizer NH₃ emissions occur during April-June reflecting the single fertilization date used in this study as calculated in the CLM for corn. While Paulot et al. [2014] also show 927 the maximum synthetic fertilizer emissions generally occur from April-June they obtain 928 929 relatively higher emissions than simulated here during the other seasons. This is likely 930 due to differences in the assumed timing of applied synthetic fertilizer: Paulot et al. [2014] consider three different synthetic fertilizer applications for each crop as well as a wide 931 932 variety of crops. The seasonal emission distribution of NH₃ emissions from manure is broader than that of synthetic fertilizer but with maximum emissions usually occurring in 933 934 April-June or July-Sept. The simulated geographical and seasonal NH₃ emission 935 distribution from manure is in broad agreement with Paulot et al. [2014].

936

937Runoff of N_r from applied synthetic fertilizer and manure applications
TAN pools as well938as nitrification and diffusion into the soil depend on precipitation and soil moisture-(see939appendix).939High manure and synthetic fertilizer N_r run off from the TAN pools (see940Figure 6-7) occur particularly across parts of China, Europe (particularly the Northern941parts) and the East central U.S. The global hotspot for simulated N_r runoff from the TAN942pools is China where runoff approaches 20 kg N ha⁻¹ yr⁻¹ for nitrogen applied as either in943manure andor synthetic fertilizer. However, we do find otherIn general the importance of

951

<u>The percent of the TAN pool nitrified or diffused directly into the soil (see Figs 7 and 8)-</u>.)
<u>also tends to be largest in wetter and cooler regions.</u> The amount of nitrogen nitrified has
an optimal temperature of 28° C and tends to occur more rapidly under moist conditions;
the diffusion of nitrogen into the soil is also promoted under wet conditions—(see
appendix)...

957

958 3.2.3 Regional and Global accounting of nitrogen losses. As nitrogen cascades through the environment it can be emitted as NH₃ or runoff or leached at many different stages. 959 960 Here we only examine the losses directly from manure or fertilizer application. Globally, the direct loss of applied nitrogen to the atmosphere as NH₃ is similar for manure and 961 synthetic fertilizer (17% for manure, 2019% for synthetic fertilizer; see Figure 9). Our 962 963 global estimates of the percent of manure and synthetic fertilizer volatilized as NH₃ are 964 similar to Bouwman et al. [2002] and Beusen et al. [2008], although our estimate for synthetic fertilizer volatilization as NH₃ is somewhat high. Bouwman et al. [2002] 965 966 estimates 19-29% of applied manure and 10-19% of applied synthetic fertilizer volatilizes as NH₃; Beusen et al. [2008] concludes 15-23% of applied manure is lost as NH₃
(including losses from housing and storage, grazing and spreading) and 10-18% of
applied synthetic fertilizer is lost.

970

We calculate the global <u>direct</u> run-off <u>from manure or fertilizer TAN pools</u> as 8% for manure N_r and 9% for synthetic fertilizer. Bouwman et al [20112013] find that 23% of deposited N_r (comprised of synthetic fertilizer, manure and <u>atmospheric</u> nitrogen deposition) runs off, higher than our estimate. However, our estimate only includes the direct runoff from the TAN pool; further loss of nitrogen due to runoff <u>and leaching</u> may also occur from the soil nitrogen pools or downstream following NH₃ emission and redeposition.

978

979 Our simulations assume a large fraction of emitted nitrogen is captured by the canopy, 980 where canopy capture accounts for 25.5% of manure losses and 30% of synthetic 981 fertilizer losses. The nitrogen captured by the canopy may have a number of fates. First, 982 Sparks [2008] posits that since foliar nitrogen uptake is a direct addition of N to plant 983 metabolism it could more readily influence plant growth than uptake from soils. As such it would decrease plant demand on soil uptake and thus conserve the soil nitrogen 984 reservoirs. Secondly, nitrogen uptake by the plants, even if not directly used in plant 985 986 metabolism, may redeposit onto the surface with litter fall. Finally, it may be emitted 987 back to the atmosphere from plants. The latter process can be represented through a 988 compensation point model between the atmosphere, the ground and stomata [e.g., Massad 989 et al., 2010]. A full accounting of this requires the simulation to be run in afully coupled 990

mode with the atmosphere and <u>soil chemistry and biogeochemistry which</u> is beyond the scope of the present study.

992

991

993 In the case of synthetic fertilizer the direct diffusion of TAN N_r into the soil pool (22%) 994 is larger than nitrification (17%); for manure it is just the opposite: the nitrification (29%)995 is larger than the direct diffusion (14%) (Figure 9). In practice, as simulated here, this makes little difference as the diffusion of nitrate into the soil pool occurs very rapidly, an 996 order of magnitude faster than the diffusion of nitrogen from the TAN pool. Thus NO₃⁻ is 997 998 directly incorporated into the soil nitrate pool without any subsequent loss. Recall, also, a 999 small percentage of manure is mechanically stirred into the soil organic nitrogen pools. 1000 Accounting for the N_r diffused from the TAN pool into the soil pools, and assuming the 1001 NH_3 emissions captured by the canopy, as well as the ammonium nitrified to NO_3 also 1002 end up in the soil pools we find that globally 75% of TAN-manure nitrogen and 71% of TAN synthetic fertilizer nitrogen ends up in the soil nitrogen or soil organic nitrogen 1003 1004 pools. Of course, once in these soil pools there may be subsequent losses of nitrogen due to runoff and leaching or emissions, but these are not calculated in this initial study. 1005

1006

The <u>global</u> percentages <u>given above</u> change appreciably when examined over subsets of countries (Figure 10). For example, over all developed countries the percentage of emissions of manure and synthetic fertilizer TAN as NH₃ [13%] is substantially smaller than for developing countries [21%]. These differences can be largely explained by the fact that developing countries tend to be located in warmer climates than developed countries. Bouwman<u>et al.</u> [2002] took these differences into account when developing emission factors for developing and industrialized countries. Bouwman [2002] calculated
NH₃ emission factors for manure of 21% and 26% for developed and industrialized
countries, respectively and for synthetic fertilizer of 7% and 18%, respectively. The

1016

1017 In our simulations 16% and 9% of applied agricultural nitrogen is emitted as NH_3 in the US and the European Union have N_r emission percentages of 16% and 9%, respectively 1018 and. The direct runoff percentages of nitrogen accounts for 9% and 14%%, of the losses 1019 of agricultural nitrogen in the US and the European Union, respectively, within a factor 1020 1021 of two although nitrogen. Nitrogen runoff is favored in the cooler moister climate of 1022 Europe. However, note the large contrast between India and China, where for India NH₃ emissions are 27% of the applied Nr with very little runoff, whereas for China the runoff 1023 1024 and emissions are approximately equal (13% and 10%, respectively).

1025

3.2.4 Comparison to other emissions inventories. Figure 11 gives a comparison of 1026 1027 manure and synthetic fertilizer NH₃ emissions from our the FAN process oriented model for 2000 and various bottom-up emission inventories, as collated by Paulot et al. [2014]. 1028 1029 The bottom-up inventories rely on emission factors depending on animal husbandry, types of synthetic fertilizer usage and other details of agricultural practices. Only the NH₃ 1030 1031 emission inventory of Huang et al. [2012] for China and Paulot et al. [2014] explicitly 1032 account for temperature to modify their emission factors; the inventory of Paulot et al. [2014] also uses wind speed to modify the emission factors. The inventories of Paulot et 1033 1034 al. [2014] for 2005-2008, Beusen et al. [2008] for 2000, and EDGAR v4.2 for 2005-2008 1035 are global inventories. We supplement these estimates over North America with The

1036 EDGAR inventory does not strictly separate the Goebes et al. [2003] estimate for 1995 1037 for NH₃ emissions into those from manure and synthetic fertilizer NH₂ emissions and the 1038 US EPA [2006] estimate of so we simply show the overall NH₃ emissions. Over the US 1039 we also give an estimate for synthetic fertilizer NH₃ from 1995 [Goebes et al., 2003] and for NH₃ emissions from animal agricultural operations- [US EPA, 2006]. Over China the 1040 1041 global NH₃ emission estimates are supplemented by Huang et al. [2012] for 2006 and Streets et al. [2003] for 2000. Over Europe results using the Greenhouse Gas and Air 1042 1043 Pollution Interactions and Synergies [GAINS] model are given [Klimont and Brink, 2004] as reported in Paulot et al. [2014]. In this study synthetic fertilizer application dataset is 1044 valid circa 2000 and the manure application dataset is valid circa 2007 [Potter et al., 1045 2010]. 1046

1047

1048 Globally all inventories give approximately the same overall NH₃ emissions of 30-35 Tg N yr⁻¹. The global apportionment of emissions between manure and synthetic fertilizer in 1049 1050 this study in approximately the ratio of 2:1, roughly consistent with that of Paulot et al. 1051 [2014] and Beusen et al. [2008]. The apportionment of manure to synthetic fertilizer emissions in the EDGAR inventory (approximately in the ratio 1:3, respectively) is not 1052 consistent with the other three inventories presented. The European and Chinese NH₃ 1053 emissions estimated here are on the low side of the other inventories, while the U.S. 1054 1055 emissions are on the high side. In Europe the current parameterization underestimates the 1056 manure emissions compared to the other estimates, while the synthetic fertilizer emissions fall between are on the Paulot et al. (2014) and GAINS emission inventories 1057 1058 and that oflow side. The EDGAR- emissions are somewhat higher than the other 1059 estimates over Europe, although may depend on exactly what is assumed for the
1060 European boundary.

1061

1062 In the U.S. the manure NH₃ emissions are close to the estimate of all theother inventories 1063 except that of EDGAR while the synthetic fertilizer emissions are high compared to all 1064 inventories, although the synthetic fertilizer emissions are close to that of EDGAR. In 1065 China our synthetic fertilizer emissions are similar to those of Huang et al. [2012], but underestimate the manure NH₃ emissions of all the other inventories except EDGAR. Of 1066 1067 the three regions examined all inventories suggest the Chinese emissions are highest. 1068 Note, however, there is considerable variation amongst the Chinese inventories for both 1069 synthetic fertilizer and manure. Our results appear to match those of Huang et al. [2012] 1070 the best.

1071

3.2.5 Site specific simulated pathways. The hourly time series of the fate of applied 1072 1073 nitrogen from manure and synthetic fertilizer at a single site better illustrates the 1074 relationship between the different pathways and the local meteorology (Fig. 12). This The 1075 large fluctuations in the NH₃ emissions and the resultant implications for atmospheric chemistry also demonstrate the desirability of inventories that respond on hourly 1076 1077 timescales to meteorological conditions. The site shown in Fig. 12 is near the Texas 1078 panhandle. It experiences several large rain events and surface temperatures ranging from 1079 0 to 18 degrees Celsius^oC over a period of about two months during the spring season. 1080 The response of the NH₃ emissions to the diurnal temperature range is clearly evident. 1081 The nitrogen losses of manure TAN due to NH₃ volatilization is initially small at the 1082 beginning of the examined period, on par with the diffusive loss and somewhat less than the loss due to nitrification. The loss by nitrification and diffusion from the TAN manure 1083 pool remain roughly constant through the period examined although both processes show 1084 1085 some response to precipitation, particularly. Note in particular the diffusion 1086 which diffusive loss reaches a maximum near May 21 presumably due to the increased 1087 water content in the soil by the prior rain event. With the rise in temperatures towards the end of the period, the emission loss of manure TAN becomes the dominant loss pathway 1088 and the TAN manure pool decreases. Closer inspection suggests, however, that the large 1089 1090 increase in the NH₃ emissions towards the end of the period cannot solely be attributed to 1091 temperature, but must also be attributed to decreased water in the TAN pool as the soil 1092 dries. The latter process increases the concentration of nitrogen species within the TAN 1093 pool. The TAN manure pool is punctuated by sharp decline events, associated with precipitation and increased runoff (Fig. 12c). Synthetic fertilizer TAN responds similarly 1094 during these events but the different temporal distribution of N application for synthetic 1095 1096 fertilizer is clearly evident in these plotsfigures. The decrease in the synthetic fertilizer TAN pool occurs on a timescale of approximately a week, consistent with the timescale 1097 1098 used in the MASAGE NH3 model (Paulot et al., 2014).

1099

1100 **3.3 Global Nitrogen Pathways: Historical**

Historical nitrogen pathways are accessed since 1850 in a simulation with changing
climate and changing application amounts. These simulations do not include changing
agricultural practices such as changes in animal housing and storage, changes in animal

1104 <u>diet and explicit changes in land use, all of which may substantially alter the nitrogen</u>
1105 pathways. Thus the results must be treated with caution.

1106

1107	The nitrogen applied produced as manure increases in the historical simulation from 21
1108	Tg N yr ⁻¹ in 1850 to 125 Tg N yr ⁻¹ in 2000 (Figure 13). In 1900 we estimate that 37 Tg N
1109	yr^{1} of manure is produced, similar to the Bouwman et al (2011) estimate of 35 Tg N yr^{-1} .
1110	Emissions of NH_3 from applied manure increase from approximately 3 Tg N yr ⁻¹ in 1850
1111	(14.3% of the manure produced) to 22 Tg N yr ⁻¹ in 2000. Bouwman et al (2011)
1112	estimates that 35 Tg N yr ⁴ is produced as manure in 1900 similar to our estimate of 37
1113	Tg N yr ⁻¹ . The (17.6% of the applied manure). On the other hand the percentage of
1114	manure nitrogen applied as manure that volatilize to NH3 increases by 4% since the
1115	preindustrial while the percentage of manure TANis nitrified decreases from 33 to 27%%
1116	since the preindustrial. Note that the year 2000 emissions in the historical simulation differ
1117	slightly from the results of the present day control for which we report the 1995-2004
1118	average emissions for the year 2000.
1119	
1120	Synthetic fertilizer nitrogen application has increased dramatically since 1960 from
1121	essentially zero tothe 1960s with an estimated 62 Tg N yr ⁻¹ applied as synthetic fertilizer
1122	in 2000. Accompanying this increase, We estimate the volatilization of synthetic fertilizer
1123	$\frac{\text{reaches}as \text{ NH}_3 \text{ is}}{12 \text{ Tg N yr}^{-1} \text{ in } 2000}$

1124

1125 For synthetic fertilizer there is an increase of emissions to the atmosphere and a decrease
1126 in nitrogen runoff. Since 1920 the (19% of that applied). The percent of synthetic
1127 fertilizer nitrogen volatized to the atmosphere as NH₃ increases from 8% to 20%, while

1128	the runoff has in 1920 was 8%. On the other hand, the percentage of synthetic fertilizer
1129	that is lost through runoff decreased since the preindustrial by 8%. It is evident that much
1130	of this changethese percentage changes can be explained by the fact the runoff of
1131	synthetic fertilizer runoff canacts to completely drain the TAN synthetic fertilizer pool
1132	atin when the application rate is small.
1133	
1134	In part the historic emission increases in NH ₃ can also be explained by changes in climate.
1135	The globally average has warmed by approximately 1° C since the preindustrial. In a
1136	sensitivity experiment the temperature was artificially increased by 1° C in the rate
1137	equations governing the nitrogen pathways following manure and synthetic fertilizer
1138	application rate prior to 1960 Under current manure and synthetic fertilizer application
1139	rates we find a global sensitivity of an additional 1 Tg NH ₃ is emitted from the manure
1140	and synthetic fertilizer pools per degree of warming. The resulting manure emissions
1141	increase by 4% and the fertilizer emissions by 3%.
1142	
1143	
1144	
1145	3.4 Sensitivity Tests
1146	We have conducted a large number of sensitivity tests to evaluate the effect of changes in
1147	individual model parameters on NH3 emissions. The various parameters may co-vary, of
1148	course, with non-linear impacts on the NH ₃ emissions; however, we have not attempted
1149	to evaluate these effects. The sensitivity tests for manure are given in Table 1, those for
1150	synthetic fertilizer in Table 2. The sensitivitiessensitivity tests are labeled with a number

1151 denoting the sensitivity parameter perturbed and a letter denoting whether the test is with 1152 respect to manure emissions (m) or synthetic fertilizer emissions (f). In each case we give 1153 the percent change in NH_3 emissions due to the parameter change and the relative 1154 emission change with respect to the relative parameter change (the sensitivity). Rationale 1155 for the assumed parameter bounds is given in the supplement.

1156

1157 Except for changes in the canopy capture parameter ($\frac{EX7mEX8m}{f}, \frac{EX8mEX9m}{f}$) and changes in the timing or composition of manure or synthetic fertilizer inputs (EX17m, 1158 1159 EX18fEX18m, EX19f, EX20f, EX21f), changes in the sensitivity parameters directly 1160 change the nitrogen cycling within the TAN pool (as described below). For the most part 1161 the synthetic fertilizer and manure TAN pools respond similarly to the parameter changes. 1162 Note also, that except for EX17EX18, where the amount of nitrogen input into the TAN pools is reduced, the total input and loss of nitrogen from the TAN pools remain the same 1163 1164 for all sensitivity experiments. In general, the sensitivity of NH_3 emissions to the 1165 imposed parameter changes are within the range of \pm 20% with many processes within 1166 the range of $\pm 10\%$. The sensitivity to the mechanical mixing of manure (EX1m, EX2m), 1167 the adjustment timescale for the water pool (EX3, EX4), the diffusion rate into the soil 1168 (EX13, EX14, EX15), the assumed depth of the water pool (EX11, EX12, EX13) and the 1169 maximum nitrification rate (EX15, EX16, EX17) all impact NH₃ emissions by less than 1170 20%. The sensitivity to the assumed background NH₃ concentration is also low (EX9, 1171 EX10). The high NH₃ concentration in equilibrium with the TAN pool renders the emissions rather insensitive to the background concentration.EX10, EX11). 1172

1174	The NH ₃ emissions are most sensitive to changes in pH (EX5m/f, EX6m/f <u>, EX7m/f</u>). The
1175	NH_3 emissions increasedecrease by a factor of 3-4approximately 60% when the pH is
1176	changed from 6 to 8. Increased pH pushes the solution towards $NH_3(aq)$ and away from
1177	$NH_4^+(aq)$ (equations 10 and 11). As $NH_g(aq)$ is in equilibrium with $NH_g(g)$ -increased
1178	pH increases from 7 to 8 and increase by 50 to 70% (for manure and synthetic fertilizer,
1179	respectively) when the pH is decreased from 7 to 6. We also test the concentration of
1180	$NH_{\mathcal{F}}(g)$ and consequently sensitivity of the emissions to the spatially explicit pH from
1181	ISRIC-WISE dataset [Batjes, 2005], with a global pH average of 6.55. The spatially
1182	explicit pH changed the manure NH ₃ emissions. Decreased pH has the opposite effect. by
1183	<u>23% and the synthetic fertilizer NH_3 emissions by 14%</u> . Changes in pH also have a large
1184	impact on the nitrification rate. Increased pH reduces $NH_4^+(aq)NH_4^+(aq)$ and thus the
1185	rate of conversion of $\frac{NH_4^+(aq)}{NH_4^+(aq)}$ to NO ₃ . The effect of pH on the <u>rate constant</u>
1186	for nitrification rate constant-is not included in the current parameterization. Parton et al.
1187	(2001) suggests this effect is small _{a} between a pH of 6 and 8, varying only on the order of
1188	15%. Changes in pH also results result in marked changes in the runoff and soil diffusion
1189	due to the large changes in emissions and nitrification: low pH's act to increase the flux
1190	of nitrogen through these loss pathways, high pH's act to decrease them.

1191

Emissions are also highly sensitive to changes in canopy capture (i.e., the parameter fcapture) as shown in EX7m/f, EX8m/f, EX9m/f. Decreasing the fraction captured by the canopy by a factor of 2 increases the emissions by approximately a factor of 3. Changes in this fraction modify the fixed ratio between the amount of nitrogen captured by the canopy and that emitted to the atmosphere, but do not impact nitrogen cycling within the 1197 TAN pools. Of course, the nitrogen captured in the canopy impacts the overall soil
1198 nitrogen budget, but this impact is not simulated here.

1199

1200 The NH_3 emissions are somewhat sensitive to the depth of the assumed water pool (EX11m/f, EX12m/f), where the water budget is calculated over depth of the water pool. 1201 1202 EX13m/f). Smaller depths (less water) give higher concentrations of all the constituents within the TAN pool resulting in largerhigher NH₃ emissions (equations 7 and 11) and 1203 larger nitrogen runoff (section 2.4.1). Larger depths (more water) have the opposite effect. 1204 1205 The diffusion of nitrogen into the soil is somewhat sensitive to changes in the assumed 1206 water depth as the coefficient of diffusion is proportional to the water content to the 10/3power (see appendix). Increased diffusion at higher depths likely reflects changes in the 1207 1208 water content of the soil with depth.

1209

We conducted various sensitivities to synthetic fertilizer applications. Early synthetic 1210 1211 fertilizer applications decrease NH₃ emissions due to their strong temperature dependence and increase the susceptibility of the TAN pool to washout. An early fertilization date 1212 1213 (set to March 15) decreases the NH_3 emissions by 23% and increases the nitrogen runoffrun off from the TAN pool by 62% (EX18fEX19f). To investigate the sensitivity to 1214 the application rate of synthetic fertilizer, synthetic fertilizer was applied over 20 days as 1215 1216 opposed to the single day application assumed in the default version (EX19fEX20f). This 1217 did not have a significant impact on the emissions. The assumed synthetic fertilizer type 1218 in the default version of the model (urea) was replaced with ammonium nitrate fertilizer 1219 in EX20fEX21f. Whereas urea is converted to NH₃ rather slowly, the conversion of 1220 ammonium nitrate is rapid (in the sensitivity test it is assumed to be instantaneously released into the TAN pool-, and result in no changes in pH). However, the emissions are 1221 not particularly sensitive to this change. This is in contrast to differences in volatilization 1222 1223 rates of different synthetic fertilizers given in Bouwman (2002). Whitehead and 1224 Raistrick (1990) show that one of the primary differences between the addition of urea 1225 versus ammonium nitrate as fertilizer is in the effect of the fertilizer on the soil pH, an effect that we do not consider in this first study. In particular urea increases the soil pH 1226 and thus the NH₃ emissions. 1227

1228

1229 Finally we test the impact of manure composition on the NH₃ emissions (EX17fEX18f). The composition of manure nitrogen excreted by animals depends in part on the 1230 digestibility of the feed, which can vary in both time and space. To investigate this 1231 uncertainty we varied the composition of the manure assumed in the default model 1232 version (50% urine, 25% available, 22.5 % resistant and 2.5% unavailable) to the less 1233 1234 soluble N excreta from dairy cattle in sensitivity simulation EX17mEX18m (41% urine, 21% available, 25% unavailable and 13% resistant [Smith, 1973]). This decreased the 1235 1236 NH₃ emissions by 21 percent.

1237

1238 It is important to emphasize that these sensitivity simulations only test the parameter 1239 sensitivity within the imposed model. In particular, the sensitivities to various farming 1240 practices are generally extraneous to the model assumptions with some exceptions. The 1241 sensitivities to synthetic fertilizer or manure input assumptions are tested in simulations 1242 <u>EX17m, EX18fEX18m, EX19f, EX20f, EX21f</u>; sensitivities to the water depth which

ςς

may crudely represent some of the impacts of plowing manure or synthetic fertilizer into the soil are examined in EX11EX12 and EX12EX13; finally modifications to soil pH are tested in EX5, EX6 and EX6EX7.

1246

1247 **4. Discussion and Conclusions**

1248 In this paper we develop a process-oriented model that predicts the climate dependent reactive nitrogen pathways from synthetic fertilizer and manure application to the surface 1249 of the land. Continued population growth will likely result in an increased application of 1250 1251 synthetic fertilizers with concurrent increases in manure production in the future 1252 (Davidson, 2012). Climate is an important determinant in the ultimate fate of this applied nitrogen, important in determining the resulting emissions of NH₃ and other 1253 1254 reactive nitrogen gases, in the runoff of the applied nitrogen, its nitrification and its incorporation into the soil organic and inorganic pools. The fate of the resultant applied 1255 1256 nitrogen may act to acerbate climate change through the formation of N_2O_1 , or perhaps 1257 mitigate climate change through increased carbon fertilization and the increased 1258 formation of aerosols. On the flip side the impact of a changing climate on agriculture 1259 and the resultant pathways for N_r is likely to be significant.

1260

Agricultural NH₃ emissions are an unusual emission source in that both natural and anthropogenic processes control their emissions. Previous global NH₃ emission inventories have exclusively used bottom up emission factors mainly governed by agricultural practices. In many cases the emission factors only implicitly include temperature dependence by using different emission factors for industrial and developing

1266 countries [e.g., Bouwman et al. 1997], although recently some inventories have included 1267 empirical emission factors that vary with temperature [Paulot et al., 2014; Huang et al., 2012]. Here, however, we take the opposite tact by constructing a model where the N_r 1268 1269 pathways and in particular the NH₃ emissions are explicitly driven by climate but where 1270 the explicit representation of most agricultural practices are minimized. We find the 1271 global emissions of NH₃ due to manure and fertilizer nitrogen sources are similar to other recent inventories, with 21 Tg N vr⁻¹ emitted from manure nitrogen and 12 Tg N vr⁻¹ 1272 emitted from synthetic fertilizer-nitrogen. Strong regional differences in emissions 1273 1274 captured by the bottom up inventories are also simulated. Moreover, we are able to 1275 simulate the inter-annual, seasonal and diurnal changes in NH₃ emissions critical for air pollution applications ([e.g., see De Meij et al., 2006].]. Most previous inventories have 1276 included no seasonal dependence of the emissions, although in some cases a seasonal 1277 dependence is empirically introduced. It is perhaps important to note that the impact of 1278 nitrogen emissions on the global carbon budget has generally made use of these previous 1279 1280 inventories without explicit seasonal or diurnal dependence of NH₃ emissions and with a 1281 rather minimal representation of the geographic meteorological dependence.

1282

The model developed here uses a process level approach to estimate nitrogen pathways from fertilizer and manure application. It is suitable for use within an Earth System model to estimate the resulting NH_3 emissions, nitrogen run-off, and the incorporation of the nitrogen into soil organic and inorganic matter. The modeled N_r pathways dynamically respond to climatic variation: (1) the breakdown timescale of manure and fertilizer into TAN depends on temperature; (2) the formation of NH_3 gas from the TAN 1289 pool is highly temperature sensitive with the rate of formation described by the 1290 temperature dependence of the thermodynamic Henry and dissociation equilibria for NH₃ 1291 [Nemitz et al., 2000]; (3) the rate of nitrification of NH₃ within the TAN pool, determined 1292 by the rate at which ammonium ions are oxidized by nitrifying bacteria to form nitrate 1293 ions [Abbasi and Adams, 1998] is controlled by environmental factors such as soil 1294 temperature and soil moisture; (4) the runoff of N_r is determined by the precipitation. 1295 Predictions for direct nitrogen runoff from fertilizer and manure nitrogen pools and the incorporation of nitrogen into soil pools from applied fertilizer and manure nitrogen are 1296 1297 some of the first made by a global process-level model. Measurements of nitrogen runoff 1298 from rivers heavily impacted by anthropogenic nitrogen input compare favorably with 1299 simulated results using the River Transport Model within the CESM [Nevison et al., 20152016]. 1300

1301

Manure is not a new nitrogen source, but contains recycled Nr from soil nitrogen 1302 1303 produced when animals eat plants. Therefore to conserve nitrogen within an earth system 1304 model, the application of manure determines the consumption of plant matter by model 1305 ruminants.animals. Specifically, the model calculates the amount of nitrogen and carbon needed for a given manure application and subtracts it from the plant leaf pools within 1306 the CLM. The manure production acts to speed up the decay and processing of plant 1307 1308 biomass, releasing different N_r products to the atmosphere than natural decay [Davidson, 1309 2009].

1311 The climate dependency incorporated into the model suggests that the pathways of nitrogen added to the land are highly spatially and temporally heterogeneous. An 1312 examination of nitrogen loss pathways at a point over Texas shows the variation of the 1313 1314 nitrogen pathways on a variety of timescales with changes in temperature, precipitation 1315 and soil moisture. Spatially, values for the percentage of manure nitrogen volatilized to 1316 NH_3 in this study show a large range in both developing countries (average of 20%) (maximum: 36 %)) and industrialized countries (average of 12% (maximum: 39 %)). The 1317 model also predicts spatial and temporal variability in the amount of NH₃ volatilized as 1318 1319 manure from agricultural fertilizers ranging from 14% [maximum 40 %] in industrialized 1320 countries to 22 % [maximum 40 %] in developing countries. As a result of temperature dependency, NH₃ volatilization is highest in the tropics with largest emissions in India 1321 1322 and China where application of fertilizer and manure is high. In comparison, the EDGAR database uses the emission factors based on Bouwman et al. (2002), where 21 % 1323 and 26 % of manure is converted into NH₃ in industrialized and developing countries, 1324 1325 respectively. The respective emission factors for fertilizer application calculated here are 7 % in industrialized countries and 18 % in developing countries. Nitrogen run-off from 1326 1327 the manure and synthetic fertilizer TAN pools is highest in areas of high N_r application and high rainfall, such as China, North America and Europe. Despite high nitrogen input 1328 rates we simulate low nitrogen runoff in India and Spain, for example. We also simulate 1329 1330 climate dependent pathways for the diffusion of N_r into the soil inorganic nitrogen pools 1331 and the nitrification of ammonium to nitrate.

1333 Historically we predict emissions of NH₃ from applied manure to have increased from approximately 3 Tg N yr⁻¹ in 1850 to 22 Tg N yr⁻¹ in 2000 while the volatilization of 1334 fertilizer reaches 12 Tg N yr⁻¹ in 2000. The NH₃ emissions increase by approximately 4% 1335 1336 for manure applications and 5% for fertilizer applications over this historical period (1930 to 2000 for fertilizer). However similar increasesUnder current manure and 1337 synthetic fertilizer application rates we find a global sensitivity of an additional 1 Tg NH₃ 1338 is emitted from the manure and synthetic fertilizer pools per degree of warming. The 1339 resulting manure emissions increase by 4% and the fertilizer emissions by 3%. Increases 1340 1341 are not evident in the runoff of nitrogen. Note, however, we do not include runoff and leaching from the mineral nitrogen pools within the CLM in these calculations. The latter 1342 may be impacted by plant nitrogen demand such that excess fertilization would act to 1343 increase the nitrogen runoff. 1344

1345

The NH₃ emissions appear reasonable when compared to other inventories on the global 1346 1347 scale, but also when compared to the local scale measurements of manure and synthetic fertilizer (Figure 2 and 3), although these latter comparisons highlight the difficulty in 1348 1349 making global scale assumptions about surface parameters and farming methodology. The biggest disagreement with the manure emission measurements is from beef cattle 1350 feedlots in Texas. On the whole the model performs best when estimating NH₃ manure 1351 1352 emissions from cows on grassland. Despite the issues described above, this model gives reasonable NH₃ emission predictions given the limited global information available on 1353 1354 the grazing land of agricultural animals.

1356	The model described here is capable of predicting global to regional impacts of climate
1357	on applied synthetic fertilizer and manure nitrogen. However, given the nature of global
1358	modeling described here and simplifying modeling assumptions there are numerous
1359	sources of error associated with our model predictions. Parameter sensitivity studies show
1360	the largest sensitivity to the assumed pH, consistent with other studies [e.g., Fletcher et
1361	al., 2013], and to the canopy deposition. The actual pH likely depends on a complex
1362	interaction of soil types, and agricultural and animal husbandry practices. Canopy
1363	capture depends on bidirectional exchange models that involve resistances between the
1364	plant canopy, the ground and ground emissions [see, e.g., Massad et al., 2010]. In the
1365	future these processes will be simulated when the CLM is coupled with a chemistry
1366	model, although the conservation of nitrogen in a biogeochemical context may present
1367	peculiar challenges. More accurate specification of the NH3 emissions can be made
1368	within an Earth System model by better accounting of synthetic fertilizer and manure
1369	application within specific PFTs or explicit incorporation into an agricultural model.
1370	
1371	The approach taken here has been rather different from an approach using emission
1372	factors to model NH ₃ emissions. Perhaps, then, the greatest source of uncertainty in this
1373	study is associated with simplifying farming methods. This It is also capable of taking into
1374	account the resulting biogeochemical cycling of nitrogen. Previous estimates of NH ₃
1375	emissions have relied on detailed information on animal type, animal housing if any and
1376	the field application of synthetic fertilizer or manure [e.g., Bouwman et al., 1997] but
1377	have minimized the representation of meteorological processes. These estimates have
1378	also not allowed for an explicit representation of the biogeochemical nitrogen cycling and

1379 loss pathways. Here we take the opposite tact by taking into account the importance of 1380 meteorological variability in accounting for regional and temporal differences in NH₃ emissions and nitrogen cycling. However, we have greatly simplified agricultural 1381 1382 management practices. - model uses a single date for synthetic fertilizer application, considers only urea fertilizer, and does not take into account manure storage methods, 1383 such as slurry pools or different types of animal manures. It also assumes a fixed depth of 1384 manure and synthetic fertilizer application. The use of simplified farming practices may 1385 be acceptable in many locations as more complex farming methods are rarely employed 1386 1387 in the developing world. The Food and Agriculture Organization [FAO, 2005] suggests over 75 % of the global agricultural land uses traditional farming methods. Nevertheless, 1388 one of the largest sources of uncertainty in this study is associated with the simplification 1389 1390 of agricultural practices. This FAN model uses a single date for synthetic fertilizer application, considers only urea fertilizer, and does not take into account manure storage 1391 methods, such as slurry pools or different types of animal manures. It also assumes a 1392 1393 fixed depth of manure and synthetic fertilizer application. Still, adapting a hybrid approach as outlined in Sutton et al. [2013] using both emission factors governing animal 1394 stockyards and the approach outlined here for manure applied to fields may be the most 1395 reasonable. The depth of synthetic fertilizer and manure mixing and a more exact 1396 representation of soil water through the vertical discretization of the soil nitrogen pools 1397 would also help account for additional agricultural practices The truth of course lies 1398 1399 somewhere between: both meteorological variability and a detailed accounting of 1400 management practices is necessary to fully account for nitrogen cycling from agricultural 1401 practices and the resulting NH₃ emissions.

1402	
1403	A number of future model improvements are necessary in the next generation model. (1)
1404	More realistic representation of manure mangement practices. Globally, somewhat over
1405	40% of manure is excreted in animal houses and stored prior to being spread onto fields.
1406	While there is a wide range of variation in animal housing and storage practices, the
1407	unique set of emission factors entailed in animal housing and storage should be
1408	incorporated in the next model generation. (2) A better representation of nitrogen
1409	transport throughout the soil column and the resulting NH ₃ generation. This would allow
1410	a differentiation between NH ₃ emissions resulting from grazing, where urine is rapidly
1411	incorporated into the soil column, versus emissions resulting from the spreading of
1412	manure slurry. It would also allow a representation of fertilizer injection or mixing into
1413	the soil column and the transport of nitrogen into the soil column in association with
1414	water transport. (3) Representation of NH ₃ emissions from different synthetic fertilizer
1415	formulations. Different types of synthetic fertilizer have rather different emission factors.
1416	As shown by Whitehead and Raistrick [1990] many of these differences can be
1417	represented by the impact of the fertilizer on soil pH. (4) A full biogeochemical coupling
1418	of the FAN process model to the overall biogeochemistry within the CLM. This would
1419	allow the nitrogen introduced through agricultural practices to impact the overall model
1420	biogeochemistry and allow a more thorough investigation of the flows of agricultural
1421	nitrogen. Here the fertilizer nitrogen would be added explicitly to the CLM crop model
1422	where appropriate. (5) A full coupling between the NH_3 emissions represented by the
1423	FAN process model and the atmospheric chemistry model through a PFT-dependent

1424 <u>compensation point approach. In this approach the atmospheric model would directly</u>
1425 <u>provide the nitrogen deposition fields to the land model.</u>

1426

1427 The increased use of synthetic fertilizer and growing livestock populations has increased 1428 N_r emission to both the atmosphere and oceans to unprecedented levels with a marked 1429 effect on the environment. We have provided a first estimate of globally distributed temporal changes in nitrogen pathways from manure and synthetic fertilizer inputs in 1430 response to climate. This is relevant to current studies investigating the ecosystem effects 1431 1432 of N_r, and in particular, how adding synthetic fertilizer to farmland affects the ocean, the 1433 atmosphere and impacts climate. The model predicts vastly different nitrogen pathways depending on the region the inputs are applied. Scenarios predicting future synthetic 1434 1435 fertilizer use and livestock populations suggest large increases in nitrogen added to the land surface from both sources [Tilman et al., 2001; Skjoth and Geels, 2013]. The climate 1436 dependence of the nitrogen pathways suggests these pathways will be sensitive to climate 1437 1438 change. The interaction of these changes with climate is not yet clear. The volatilization 1439 of NH₃ increases exponentially with temperature suggesting future increases are likely. 1440 However, increases in temperature may surpass the optimal temperature at which certain biological processes occur, slowing the process. Washout pathways are also likely to 1441 change, not only with climate, but with increases in nitrogen loading. Future applications 1442 1443 of this model will investigate the tight coupling between nitrogen, agriculture and climate. 1444

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