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26 **Key Points:**

- 27 • A “three stages hypothesis” was developed to generalize the diverse responses of forest
28 soil CH₄ flux to N input
- 29 • CH₄ uptake by global forest soils was estimated to be 11.2 Tg yr⁻¹, with N deposition
30 suppressing ~3% of this uptake
- 31 • Effective regulation to reduce N deposition would promote CH₄ uptake by N-saturated
32 forests and mitigate global warming

33

34 Abstract

35 Methane (CH₄) is the second most important atmospheric greenhouse gas (GHG) and forest soils
36 are a significant sink for atmospheric CH₄. Uptake of CH₄ by global forest soils is affected by
37 nitrogen (N) deposition; clarifying the effect of N deposition helps to reduce uncertainties of the
38 global CH₄ budget. However, it remains an unsolved puzzle why N input stimulates soil CH₄
39 flux (R_{CH_4}) in some forests while suppressing it in others. Combining previous findings and data
40 from N addition experiments conducted in global forests, we proposed and tested a “stimulating-
41 suppressing-weakening effect” (“three stages”) hypothesis on the changing responses of R_{CH_4} to
42 N input. Specifically, we calculated the response factors (f) of R_{CH_4} to N input for N-limited and
43 N-saturated forests across biomes; the significant changes in f values supported our hypothesis.
44 We also estimated the global forest soil CH₄ uptake budget to be approximately 11.2 Tg yr⁻¹.
45 CH₄ uptake hotspots were located predominantly in temperate forests. Furthermore, we
46 quantified that current level of N deposition reduced global forest soil CH₄ uptake by ~3%. This
47 suppression effect was more pronounced in temperate forests than in tropical or boreal forests,
48 likely due to differences in N status. The proposed “three stages” hypothesis in this study
49 generalizes the diverse effects of N input on R_{CH_4} , which could help improve experimental
50 design. Additionally, our findings imply that by regulating N pollution and reducing N
51 deposition, soil CH₄ uptake can be significantly increased in the N-saturated forests in tropical
52 and temperate biomes.

53

54 Plain Language Summary

55 Methane is an important greenhouse gas. Forest soils can absorb methane from the atmosphere
56 and mitigate its warming effect. Meanwhile, forests suffer from high atmospheric nitrogen

57 deposition, yet the effect of nitrogen on the methane uptake by forest soils remain unclear. Using
58 data from global nitrogen addition experiments, we validated a “stimulating-suppressing-
59 weakening effect” (“three stages”) hypothesis, which could explain the diverse responses of soil
60 methane flux to nitrogen input observed in different forests. On the basis, we quantified that
61 nitrogen deposition decreased global forest soil methane uptake by approximately 3%. Our
62 findings also imply that that by regulating nitrogen pollution, soil methane uptake can be
63 significantly increased in the nitrogen-saturated forests in tropical and temperate biomes,
64 potentially mitigate global warming.

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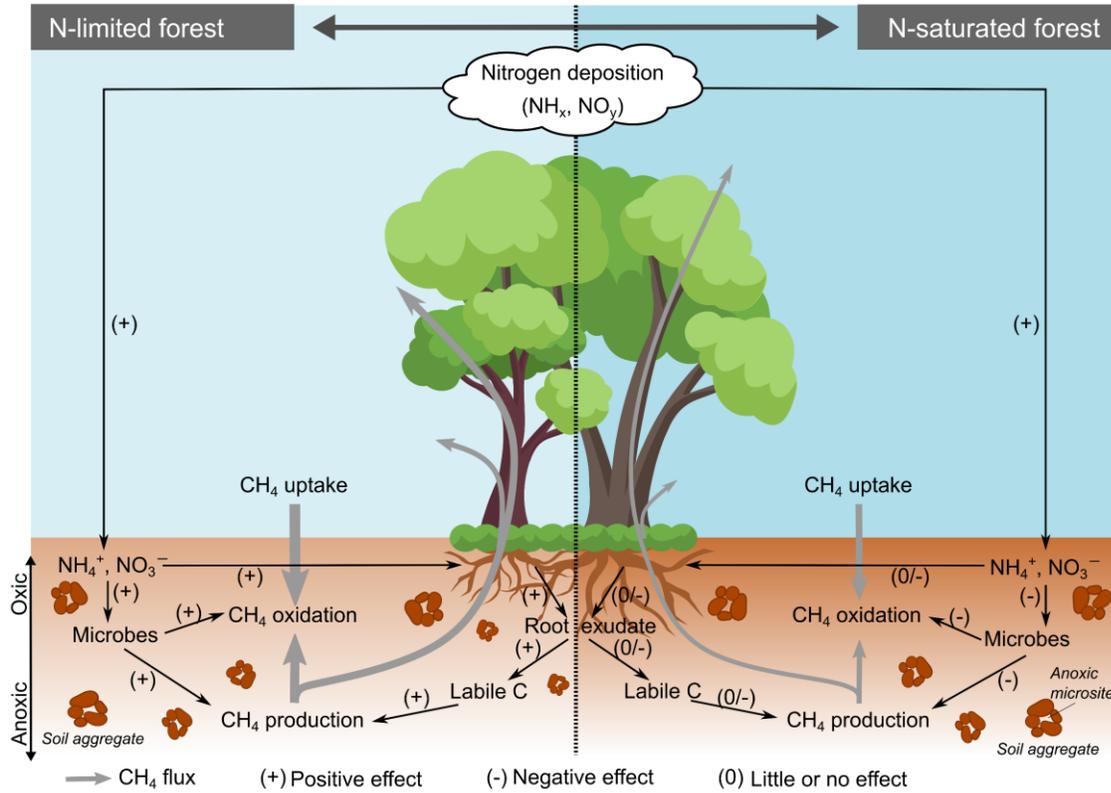
67 **1 Introduction**

68 Methane (CH₄) is the second most important greenhouse gas (GHG), responsible for
69 approximately 20% of global warming since the industrial revolution (Kirschke et al., 2013;
70 Saunio et al., 2020). Biological CH₄ absorption by soils contributes to 5–7% of total CH₄
71 removal from the atmosphere (Dlugokencky et al., 2011; Saunio et al., 2020). Soils, however,
72 do not always function as net sinks of atmospheric CH₄. The net effect of two biological
73 processes, namely CH₄ production ("methanogenesis", widespread in anoxic microsites and deep
74 soils; Angel et al., 2012; Kotelnikova, 2002; Lacroix et al., 2023) and CH₄ oxidation
75 ("methanotrophy", widespread in oxic surface soils; Le Mer & Roger, 2001), determines whether
76 a soil is a source or sink of CH₄. The delicate, variable balance between soil CH₄ consumption
77 and production depends on various changing environmental factors, which leads to uncertainties
78 in soil-atmosphere CH₄ exchange dynamics and the potential feedback of soil CH₄ uptake to
79 climate change (Bodelier & Steenbergh, 2014; Feng et al., 2020). Approximately 30% of the
80 Earth's land surface are forests, which are significant for regulating global climate (Bonan,
81 2008). Recently, forests received much attention because forestland-based management
82 practices, such as afforestation, are crucial for achieving net-zero emissions by mid-21st century
83 and mitigating global warming (Griscom et al., 2017; IPCC, 2021). Mechanisms underlying
84 forest GHG fluxes are fundamental to assessing and predicting the effectiveness of the practices.
85 Therefore, it is important and urgent to understand global forest soil CH₄ flux variations under
86 environmental changes.

87 Since the 19th century, following an exponential increase in the artificial production and
88 anthropogenic emission of reactive nitrogen compounds (e.g., through fertilizer use, combustion
89 processes), deposited nitrogen (N) to terrestrial ecosystems has increased by more than threefold

90 (Galloway et al., 2004). This exogenous N input impacts the structure and functioning of
91 ecosystems by altering plant and microbial properties (Vitousek et al., 1997). Furthermore,
92 enhanced N deposition has led to widespread "N saturation" of forests (Ågren & Bosatta, 1988),
93 resulting in divergent responses of ecological processes (such as primary production and N
94 mineralization) to N input in N-saturated as compared to N-limited forests (Aber et al., 1998). To
95 quantify the effect of N deposition on forest system functioning, researchers have conducted N
96 addition experiments in forests worldwide during the past half-century. Although the effects of N
97 input on some ecosystem properties have been clarified by the experiments, the relationship
98 between N input and soil CH₄ flux remained an unsolved puzzle. Some experiments revealed
99 stimulating effects of N input on soil CH₄ uptake, while some others showed inhibited soil CH₄
100 uptake by N input (Veldkamp et al., 2013; Zhang et al., 2012). Currently, there is no universally
101 applicable framework to explain the diverse responses of soil CH₄ flux to N input. This lack of
102 understanding hinders the development of quantitative models and assessment of the change in
103 global forest soil CH₄ budget caused by N deposition.

104



105

106 **Fig. 1.** Soil CH₄ fluxes exhibit varying responses to N deposition as forests transition from a N-
 107 limited status to a N-saturated status (or vice versa) due to human activities.

108

109 The response of soil CH₄ flux to N deposition is influenced by the rate and persistence of

110 N input and the N availability in forests (Aronson & Helliker, 2010; Chang et al., 2021). In N-

111 limited forests, a low N input rate can stimulate plant and microbial activities. Methanotrophs,

112 which are more active in near-surface soils (Butterbach-Bahl & Papen, 2002), may benefit from

113 the external N supply, with increased abundance and activity (see Fig. 1; Bodelier & Laanbroek,

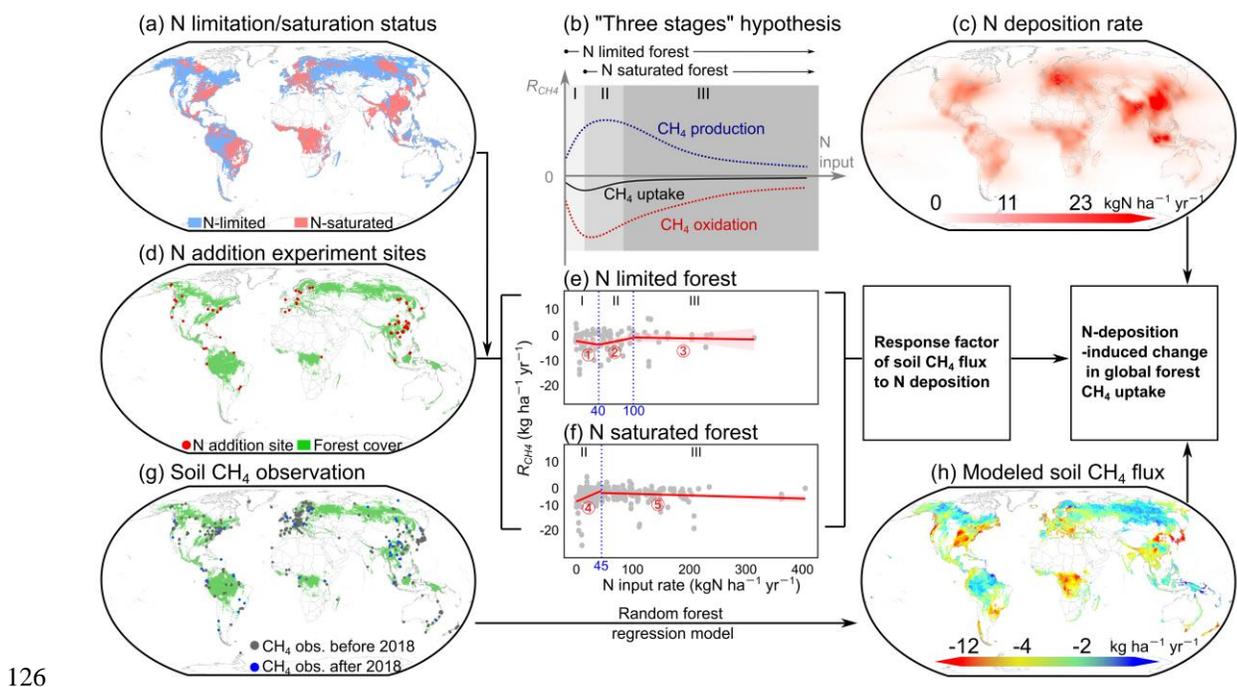
114 2004), causing more CH₄ to be oxidized. However, CH₄ oxidation can be suppressed by high N

115 input, as a result of the inhibitory effect of excessive N on methanotrophs (Agathokleous et al.,

116 2020; Chen et al., 2021; Peng et al., 2019). In N-saturated forests, the N supply surpasses the

117 demands of plants and microbes. Consequently, suppression of soil CH₄ uptake has been

118 observed even under a low N input rate (Mochizuki et al., 2012). Moreover, methanogenesis and
 119 methanotrophy can both be suppressed under high N input rates, resulting in a weak response of
 120 soil CH_4 flux to N input (Keiluweit et al., 2018; Steinkamp et al., 2001). Therefore, there appear
 121 to be distinct stages in the response of soil CH_4 fluxes to N input, with N-limited and N-saturated
 122 forests experiencing different stages under elevated N input rates. In light of these observations,
 123 we have developed a stimulating-suppressing-weakening effect (referred to below as "three
 124 stages") hypothesis (see Fig. 2b) to offer a unified framework that generalizes the response of
 125 soil CH_4 flux to N input.



126

127 **Fig. 2.** Workflow for quantifying the impact of N deposition on forest soil CH_4 flux (R_{CH_4}) in
 128 global forests. (a) N limitation or saturation status of global forests, indicated by the sensitivity
 129 of soil N_2O emissions to N deposition (See Supporting Text S1 for details). (b) Proposed "three
 130 stages" hypothesis on the response of R_{CH_4} to N input. (c) Global map of N deposition rates, data
 131 from Ackerman et al. (2019). (d) Forest sites where N addition experiments was conducted and

132 R_{CH_4} was observed (CH₄_exp dataset). (e) Segmented regression models on R_{CH_4} and N input
133 rate, using data from N-limited forests where N addition experiments lasted for no more than 3
134 years. (f) Segmented regression models on R_{CH_4} and N input rate, using data from N-saturated
135 forests and forests with N addition experiments lasting more than 3 years (see Supporting Table
136 S1 for model parameters). (g) Forest sites where no experiment was conducted and R_{CH_4} was
137 observed under natural conditions (CH₄_obs dataset). (h) Estimated R_{CH_4} in global forests.

138

139 In this study, we gathered data from N addition experiments conducted in forests
140 worldwide. We aimed to examine the validity of the "three stages" hypothesis by comparing the
141 alterations in soil CH₄ flux (R_{CH_4}) caused by each unit of N input, known as "response factors",
142 in N-limited and N-saturated forests. Furthermore, we aimed to estimate the global budget for
143 CH₄ uptake by forest soils and determine the specific contribution of N deposition to this budget.

144

145 **2 Methods**

146 2.1 Data source

147 We conducted a systematic compilation of soil CH₄ flux data observed in N addition
148 experiments by searching relevant literature published prior to 1/1/2022 in the Web of Science
149 Core Collection (www.webofscience.com) and the China National Knowledge Infrastructure
150 Theses and Dissertations Database (<https://oversea.cnki.net/kns?dbcode=CDMD>). The search
151 utilized keywords "forest" AND ("greenhouse gas" OR "CH₄" OR "methane"). Subsequently, we
152 manually refined the obtained 8702 papers and theses based on the following criteria: (i) N
153 addition experiments were conducted in forest ecosystems with recorded site locations and N

154 addition doses and (ii) field observations of soil CH₄ flux were measured using gas
155 chromatograph technique (Holland et al., 1999). The resulting compiled dataset, named
156 "CH₄_exp", comprises 465 observations from 85 sites (refer to Fig. 2; Supporting Information
157 Data Set S1).

158 Additionally, we compiled data on soil CH₄ flux observed under natural conditions. Soil
159 CH₄ flux observations before 2018 were obtained from three published datasets by Dutaur and
160 Verchot (2007), L. J. Yu et al. (2017), and Ni and Groffman (2018), while data observed after
161 2018 were gathered from the abovementioned 8702 literature using a different set of criteria: (i)
162 forest soil CH₄ fluxes were observed in the field and measured using gas chromatograph
163 technique (Holland et al., 1999) and (ii) no N or other nutrient addition experiments were
164 conducted at the forest sites. The compiled dataset, referred to as "CH₄_obs", consists of 1946
165 observations from 652 forest sites worldwide (see Fig. 2; Supporting Information Data Set S2).

166 We also collected supplementary information on environmental factors, including
167 climate, N deposition, and soil properties. In cases where not all required information was
168 provided for a particular site, we extracted data from spatial datasets based on the coordinates of
169 the sites. Specifically, temperature and precipitation data were sourced from the Climatic
170 Research Unit, University of East Anglia (https://crudata.uea.ac.uk/cru/data/hrg/cru_ts_4.03/);
171 soil texture data were obtained from the Harmonized World Soil Database
172 ([https://www.fao.org/soils-portal/data-hub/soil-maps-and-databases/harmonized-world-soil-](https://www.fao.org/soils-portal/data-hub/soil-maps-and-databases/harmonized-world-soil-database-v12/en/)
173 [database-v12/en/](https://www.fao.org/soils-portal/data-hub/soil-maps-and-databases/harmonized-world-soil-database-v12/en/)); global N deposition data were derived from a published dataset by Ackerman
174 et al. (2019); forest cover data were obtained from the GLASS-GLC project (Liu et al., 2020);
175 and forest biome boundaries were sourced from the Global Forest Monitoring project (Hansen et
176 al., 2010).

177

178 2.2 Calculating response factor of soil CH₄ flux to N input

179 Considering the distinct responses of R_{CH_4} to N input in N-limited and N-saturated
180 forests, we partitioned the CH₄_exp dataset into two subsets based on whether the experimental
181 sites were N-limited or N-saturated. Since the N limitation or saturation status was largely
182 unknown at most experimental sites, we initially predicted the N limitation or saturation status of
183 global forests, using the sensitivity of soil N₂O emissions to N deposition as an indicator; the
184 accuracy of prediction exceeded 70% on both regional and global levels (see Supporting Text S1
185 for details about determining the N status of global forests). On the basis, we determined the N
186 limitation or saturation status of forest sites in CH₄_exp dataset; the dataset was then divided into
187 two sub-datasets, CH₄_exp_NL and CH₄_exp_NS, consisting of data from N-limited and N-
188 saturated forests, respectively (Supporting Fig. S1). Moreover, recognizing that long-term N
189 addition can lead to the transition of N-limited forests into N-saturated forests, we also included
190 data from experimental sites where N addition had been implemented for more than three years
191 in the CH₄_exp_NS sub-dataset.

192 Using data from the CH₄_exp_NL sub-dataset, we constructed segmented linear
193 regression models to account for the changing relationship between soil CH₄ flux and N input
194 (see Fig. 2e,f, Supporting Table S1). In accordance with the segmented regression models, we
195 further divided the sub-dataset into several groups based on N input levels (low, medium, and
196 high). Using data from each group, we computed the change in soil CH₄ flux per unit of N input
197 on site level (Eq. 1), which we referred to as the response factor of soil CH₄ flux to N input (f).
198 Similarly, we calculated f for N-saturated forests using data from the CH₄_exp_NS sub-dataset.

199
$$f = \frac{R_2 - R_1}{N_2 - N_1} \quad (\text{Eq. 1})$$

200 where f is the response factor of the soil CH₄ flux to N input, N_1 and N_2 are the two different N
 201 input rates (kg N ha⁻¹ yr⁻¹), R_1 and R_2 are the corresponding soil CH₄ fluxes (kgCH₄ ha⁻¹ yr⁻¹).

202

203 2.3 Estimating global forest soil CH₄ flux using random forest regression method

204 Using soil methane flux (R_{CH_4}) observed under natural conditions (CH₄_obs dataset), we
 205 predicted R_{CH_4} of global forests on grid level with random forest regression models (Breiman,
 206 2001). In practice, R_{CH_4} observed at the same site were aggregated by taking the mean value.

207 After excluding one outlier that is approximately two times lower than the second lowest R_{CH_4}
 208 value, we randomly sampled 20% of the 872 entries of data to form a testing dataset (n = 175).

209 75% of the remaining data (i.e., 60% of all data) were randomly chosen to form a training dataset
 210 (n = 523), and the rest data were to allow for the variation of training dataset (n = 174). Climate,
 211 N deposition, soil texture, and soil N status variables were used as predictors (Supporting Table
 212 S2).

213 Because the constructed models can vary depending on which data were used to train the
 214 models, the random sampling of training data was repeated for 1000 times, which derived 1000
 215 models. When estimating R_{CH_4} on grid level, each grid had 1000 predicted R_{CH_4} values from the
 216 1000 models. The mean R_{CH_4} of the 1000 values were used as the estimated R_{CH_4} of the grid, and
 217 the standard error of the estimation was also calculated from the 1000 values.

218 Estimated R_{CH_4} for grids in the test dataset (which were never used in model
 219 construction) were then compared with observed values to measure the accuracy of prediction.

220 Also, we randomly sampled a different 80% of data to form different training datasets, repeated
 221 the above processes, and checked the robustness of our prediction on grid level.

222

223 2.4 Quantifying the contribution of N deposition to global forest soil CH₄ budget

224 By summarizing the grid-level R_{CH_4} data (Eq. 2), we obtained soil CH₄ budgets for
 225 forests in various regions. Combining the N deposition rate with the previously quantified
 226 response factor (f), we determined the N-deposition-induced changes in the soil CH₄ budget.
 227 This allows us to quantify the contribution rate of N deposition to the global forest soil CH₄
 228 budget (Eq. 3).

$$229 \quad \text{Budget} = \sum_i (R_{CH_4,i} \times A_i) \quad (\text{Eq. 2})$$

$$230 \quad \text{Contribution rate} = \frac{\sum_i (N_{depo,i} \times f_i)}{\text{Budget}} \times 100\% \quad (\text{Eq. 3})$$

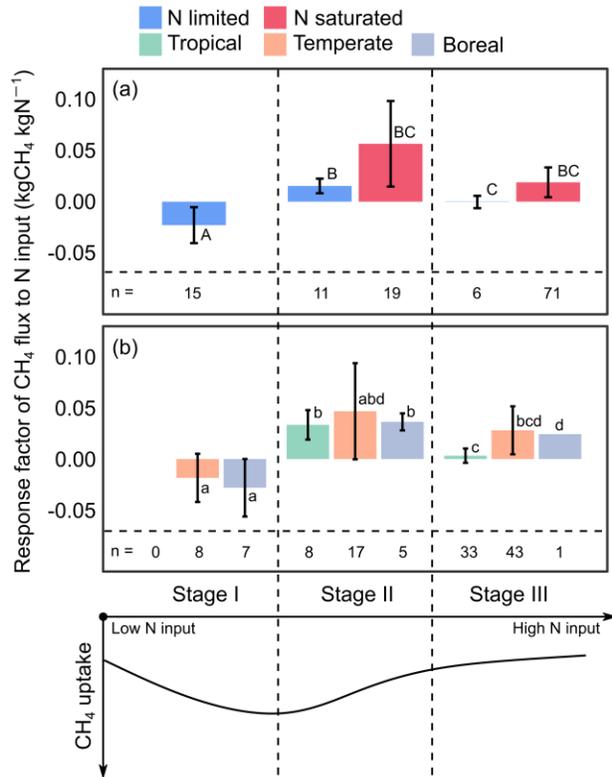
231 where $R_{CH_4,i}$ is the soil CH₄ flux in grid i (kgCH₄ ha⁻¹ yr⁻¹), A_i is the forest area in grid i (ha),
 232 $N_{depo,i}$ denotes the N deposition rate in grid i (kgN ha⁻¹ yr⁻¹), and f_i is the response factor
 233 determined based on the N deposition rate and the N limitation/saturation status of the forests in
 234 grid i (kgCH₄ kgN⁻¹).

235 Additionally, we employed the bootstrap method (Davison & Hinkley, 1997) to compare
 236 the mean R_{CH_4} values among forests in different biomes. Furthermore, we conducted an analysis
 237 to determine the relative importance of environmental factors in explaining the spatial variation
 238 in R_{CH_4} (Grömping, 2006). Data analyses were performed using R software (R Core Team,
 239 2020), with a significance level set at $p < 0.05$. The production of maps was accomplished using
 240 ArcGIS software (ESRI, 2011).

241

242 **3 Results**243 3.1 Response of forest soil CH₄ fluxes to N input

244 Locally weighed regression models showed different patterns in the responses of soil CH₄
245 flux (R_{CH4}) to N input in N-limited and N-saturated forests (Supporting Fig. S1; see Supporting
246 Text S1 for determination of N limitation or saturation status of global forests), and the responses
247 changed with N input level. Segmented linear regression models were separately fitted to data
248 from N-limited and N-saturated forests to detect the thresholds in the phased responses of R_{CH4} to
249 N input. In accordance with the detected N input thresholds (40 and 100 kgN ha⁻¹ yr⁻¹ for N-
250 limited forests, and 45 kgN ha⁻¹ yr⁻¹ for N-saturated forests; refer to Fig. 2e,f), data obtained in
251 N-limited forests were then divided into three groups based on N input levels (low, medium, and
252 high), whereas data from N-saturated forests were divided into two groups (low and high N
253 inputs). The quantified response factor (f) for N-limited forests under low, medium, and high N
254 inputs represented the response of R_{CH4} to N input in Stages I, II, and III, respectively (Fig. 3).
255 Similarly, the quantified f for N-saturated forests under low and high N inputs represented the
256 response of R_{CH4} to N input in Stages II and III. The observed changes in f values across different
257 stages and different biomes provide support for our "three stages" hypothesis (Fig. 3).



258

259 **Fig. 3.** Changes in the response factors (f) of soil CH_4 flux to N input at three stages. (a) Mean
 260 response factors of N-limited and N-saturated forests; (b) Mean response factors of forests in
 261 different biomes. The error bars represent the standard errors of the mean values. Different letters
 262 beside each column indicate that the mean values of f were significantly different ($p < 0.05$).
 263 Numbers below each column are the number of f values derived from CH_4 _exp dataset.

264

265 At Stage I, mean value of f in N-limited forests was significantly lower than 0 ($p <$
 266 0.005), indicating that low N input stimulated soil CH_4 uptake (or suppressed soil CH_4
 267 emissions). At Stage II, the mean f values for both N-limited and N-saturated forests were
 268 significantly higher than 0 ($p < 0.005$), signifying that medium N input suppressed soil CH_4
 269 uptake in N-limited forests, while low N input had a suppressing effect on R_{CH_4} in N-saturated

270 forests. At Stage III, the mean f value for N-limited forests approached zero, and the mean f
271 value for N-saturated forests at this stage was lower than the mean f value for N-saturated forests
272 in Stage II. These findings suggested that the response of R_{CH_4} generally diminished under high
273 N input in both N-limited and N-saturated forests. Furthermore, we observed a decrease in the
274 standard deviation of R_{CH_4} under high N input (Supporting Fig. S2), indicating the weakening of
275 at least one process underlying soil CH_4 flux (methanogenesis or methanotrophy). This in
276 combination with the relatively stable mean R_{CH_4} values under high N input (refer to Supporting
277 Fig. S1), suggested that both CH_4 production and oxidation rates declined (see Supporting Text
278 S2 for inference processes), which agreed with our hypothesis (Fig. 2b).

279 On biome level, the significant changes in f values at different stages were as well
280 consistent with our hypothesis (Fig. 3).

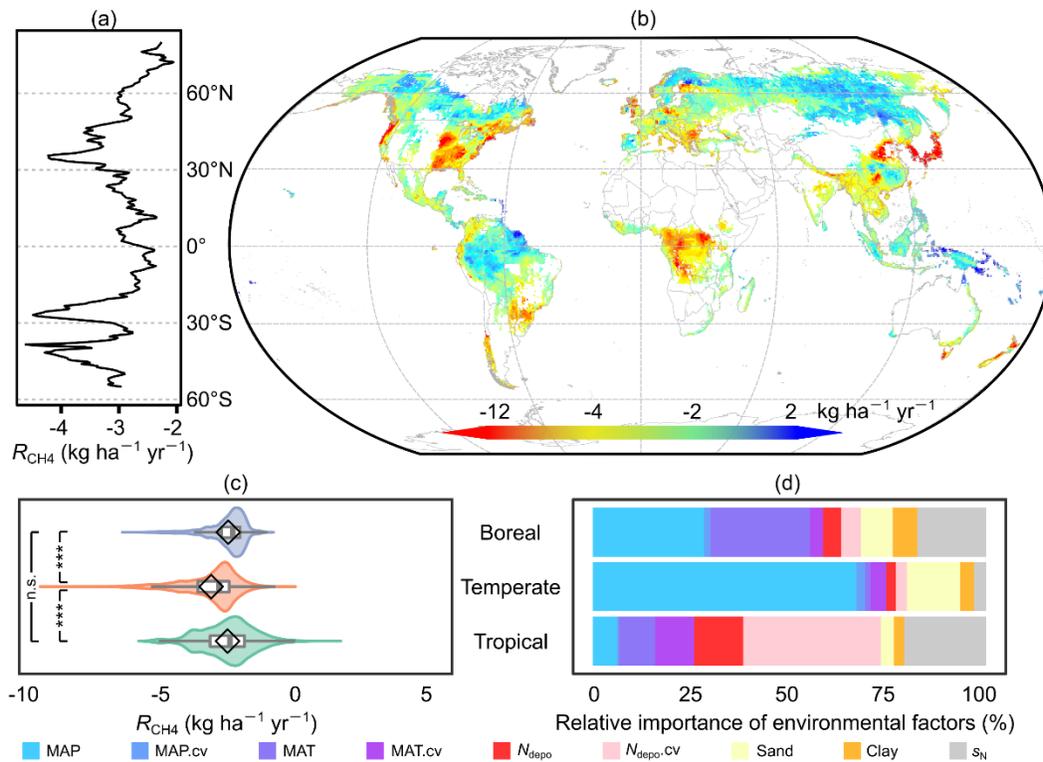
281

282 3.2 Global forest soil CH_4 fluxes, and the underlying environmental variables

283 We built random forest regression models using natural R_{CH_4} observations ($R_{CH_4_obs}$
284 dataset; Supporting Table S2), and predicted R_{CH_4} in global forests on grid level. Estimated R_{CH_4}
285 was compared with observations from the testing dataset not used in model construction. The
286 correlation coefficient of 0.6 proved the reliability of this method (Supporting Fig. S3). Also, we
287 randomly sampled data to form different training datasets and built a different set of random
288 forest models. Estimated R_{CH_4} were in good agreement with our initial estimations ($r = 0.91$;
289 Supporting Fig. S4), showing the robustness of grid-level estimations of R_{CH_4} .

290 The average R_{CH_4} for global forests was estimated to be $-2.95 \text{ kg ha}^{-1} \text{ yr}^{-1}$. Mean R_{CH_4} of
 291 temperate forests was significantly more negative than those of tropical and boreal forests ($p <$
 292 0.001 ; refer to Fig. 4).

293 Environmental factors influencing the spatial variation in R_{CH_4} differed across biomes. In
 294 tropical forests, approximately 50% of the explainable variation could be attributed to N
 295 deposition and its annual fluctuations. In temperate forests, precipitation played a dominant role
 296 in explaining the spatial variation in R_{CH_4} . Both precipitation and temperature emerged as the
 297 main factors influencing R_{CH_4} in boreal forests.



298

299 **Fig. 4.** Estimation of soil CH₄ fluxes (R_{CH_4}) in global forests using the random forest method. (a)
 300 Latitudinal gradient in soil CH₄ flux. The black line represents the average R_{CH_4} values across
 301 latitudes. (b) Global map illustrating soil CH₄ fluxes in forests. Negative values indicate net CH₄
 302 uptake. (c) Violin plots and boxplots displaying the statistical distribution of R_{CH_4} values in

303 different biomes. (d) Assessment of the relative importance of environmental factors in the
304 spatial variation of R_{CH_4} across different biomes. The factors include mean annual precipitation
305 (MAP), mean annual temperature (MAT), atmospheric N deposition rate (N_{depo}), soil sand
306 content (Sand), soil clay content (Clay), and sensitivity of soil N_2O emission to N deposition
307 (s_N), which serves as an indicator of the N limitation/saturation status of forests.

308

309 3.3 Contribution of N deposition to global forest soil CH_4 budget

310 By summarizing the grid-level R_{CH_4} data, the CH_4 uptake by global forest soils was
311 estimated to be approximately $11.2 \text{ TgCH}_4 \text{ yr}^{-1}$. Currently, N deposition reduced global forest
312 soil CH_4 uptake by $0.29 \text{ TgCH}_4 \text{ yr}^{-1}$, representing a global suppression of 2.6%. The overall
313 effect of N deposition on forest soil CH_4 uptake varied among different biomes (see Table 1). N
314 deposition suppressed soil CH_4 uptake by 3–6% in tropical and temperate forests, whereas it
315 stimulated boreal forest soil CH_4 uptake by 1.1%.

316 **Table 1.** Contribution of N deposition to soil CH₄ budget in global forests.

	Area (10 ⁸ ha)	N deposition rate (kgN ha ⁻¹ yr ⁻¹)	Response factor (kgCH ₄ kgN ⁻¹)	N deposition induced change in CH ₄ emission (TgCH ₄ yr ⁻¹)	CH ₄ emission rate (kgCH ₄ ha ⁻¹ yr ⁻¹)	CH ₄ emission budget (TgCH ₄ yr ⁻¹)	Contribution rate of N deposition to CH ₄ budget (%)
Tropical forest							
N limited	8.4	3.87	-0.019 (0.024)*	-0.06 (0.08)	-2.33 (0.011)	-1.77 (0.01)	3.4
N saturated	9.6	7.23	0.034 (0.015)	0.24 (0.11)	-3.20 (0.012)	-3.07 (0.01)	-7.8
Subtotal	18.0	5.66		0.18 (0.13)	-2.81 (0.012)	-4.84 (0.02)	-3.7
Temperate forest							
N limited	3.6	5.40	-0.019 (0.024)	-0.04 (0.05)	-3.41 (0.011)	-1.34 (0.004)	3.0
N saturated	3.8	10.50	0.047 (0.047)	0.19 (0.19)	-3.48 (0.012)	-1.34 (0.004)	-14.2
Subtotal	7.4	7.99		0.15 (0.20)	-3.44 (0.012)	-2.68 (0.01)	-5.6
Boreal forest							
N limited	10.0	2.08	-0.028 (0.028)	-0.07 (0.07)	-2.65 (0.012)	-2.70 (0.01)	2.6
N saturated	3.0	2.53	0.037 (0.008)	0.03 (0.01)	-3.32 (0.016)	-1.00 (0.01)	-3.0
Subtotal	13.0	2.18		-0.04 (0.07)	-2.79 (0.013)	-3.70 (0.02)	1.1
Total	38.4	4.64		0.29 (0.25)	-2.95 (0.012)	-11.22 (0.05)	-2.6

317 * No observations were available for N-limited tropical forests; hence, the mean response factor of N-limited temperate forests was used instead.

318 Values in parentheses represent standard errors of the mean.

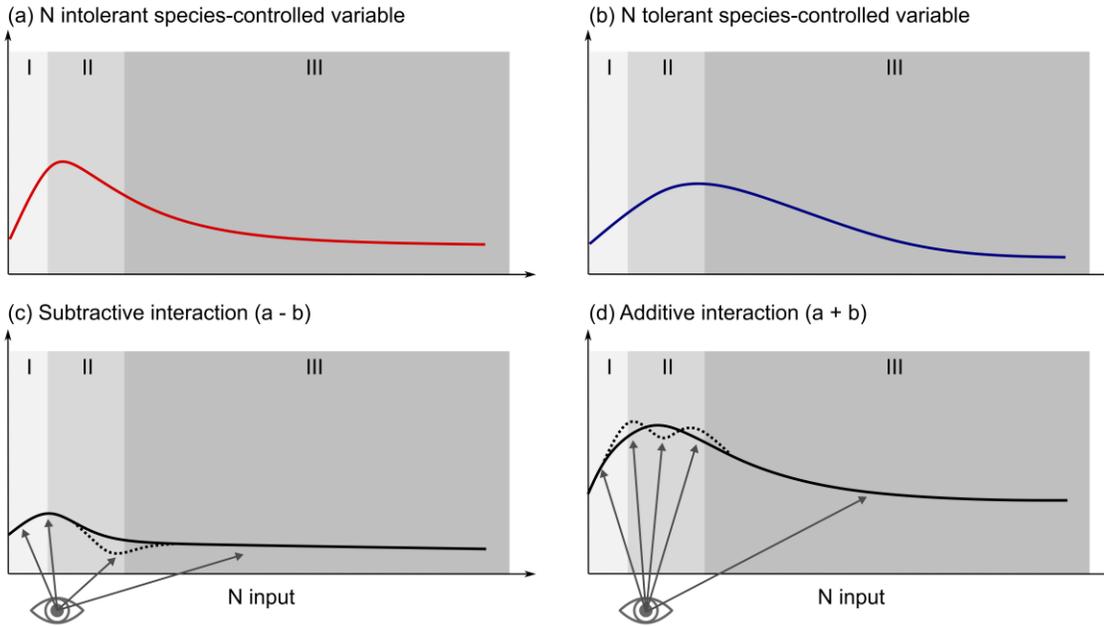
319 **4 Discussion**320 4.1 “Three stages” hypothesis generalizes response of forest soil CH₄ flux to N input

321 Both the exogenous N input level and the internal properties of forest ecosystems (such
322 as N availability) can influence the response of soil CH₄ flux to N input. Manipulative
323 experiments and meta-analyses have been conducted to examine changes of R_{CH_4} in response to
324 different N input levels (Aronson & Helliker, 2010; Chen et al., 2021). However, the spatially
325 varying responses of R_{CH_4} to N input in different forests remained unresolved. The absence of a
326 comprehensive framework for the effect of N input on R_{CH_4} has impeded the integration of site-
327 level observations and identification of a global pattern. In this study, we proposed a "three
328 stages" hypothesis to elucidate the relationship between R_{CH_4} and N input. It not only accounts
329 for the varying responses of R_{CH_4} to different levels of N input, but also explains the divergent
330 effects of N input on R_{CH_4} in N-limited and N-saturated forests.

331 The "three stages" concept is primarily determined by the biphasic dose-response
332 relationship between N input and biotic processes, exhibiting a stimulating effect at low doses
333 and a suppressing effect at high doses (referred to as the "hormesis" effect; (Agathokleous et al.,
334 2020). Additionally, the asynchronous responses of methane production and oxidation processes
335 to N input play a role; the hormesis effect leads to the transition from Stage I to subsequent
336 stages, and the transition from Stage II to Stage III occurs due to the lower tolerance of
337 methanotrophs to N input as compared to methanogens (Li et al., 2021). While methanotrophs
338 are generally sensitive to nitrogen addition (Nyerges & Stein, 2009), at least some methanogens
339 (such as hydrogenotrophic methanogens) are tolerant to high N and low soil pH (Horn Marcus et
340 al., 2003).

341 More generally, We postulate that although the responses of many biological processes
342 (such as net primary production or N mineralization) across a wide range of N input is unimodal
343 (Aber et al., 1998), response patterns may differ for variables controlled by the interactions of
344 multiple functional groups (Fig. 5), such as soil CH₄ flux and soil respiration rate. Therefore,
345 experiments testing the response of forest soil CH₄ fluxes to N input may need to test at least
346 three levels of N additions, so as to capture the changes of response. Conducting experiments
347 with multiple N addition levels will facilitate a comprehensive understanding of changes in
348 biogeochemical cycles and the underlying mechanisms. Hypothesized response patterns of soil
349 variables controlled by additive or subtractive interactions (Fig. 5) may provide reference for
350 setting N addition levels in experiments.

351 Moreover, it should be noted that the calculated response factors showed high degrees of
352 uncertainty, due to the limited experimental data available. Additional experiments are required,
353 especially in boreal forests which are sensitive to future climate and N deposition change
354 (Fleischer et al., 2015; Galloway et al., 2004). On the basis, researchers will be able to reduce
355 uncertainties in the global forest soil CH₄ budget under spatially and temporally varying N loads
356 deposited from the atmosphere.



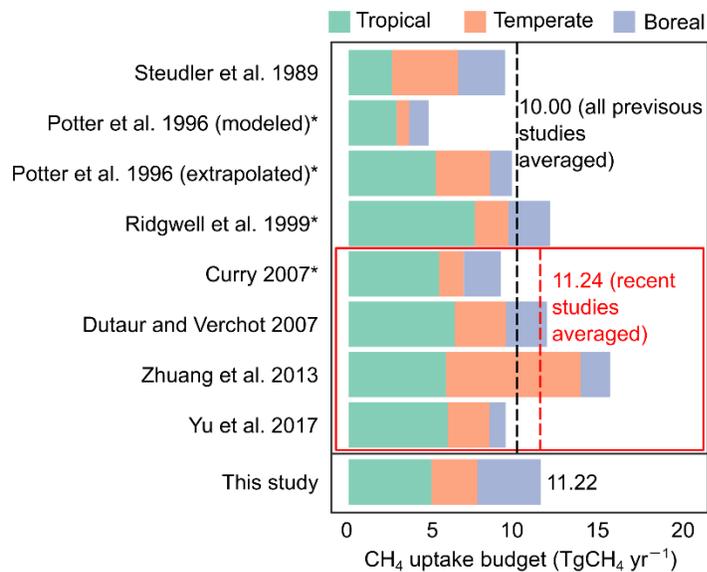
357

358 **Fig. 5.** Hypothesized effects of N input on variables controlled by N-tolerant or intolerant
 359 species, and their interactions. (a) Effect of N input on N intolerant species-controlled variables
 360 (such as CH_4 oxidation rate); (b) Effect of N input on N tolerant species-controlled variables
 361 (such as CH_4 production rate); (c) Effect of N input on variables controlled by subtractive
 362 interactions between different species (such as soil CH_4 uptake rate, which is the difference
 363 between CH_4 oxidation rate and CH_4 production rate); (d) Effect of N input on variables
 364 controlled by additive interactions between different species (such as soil respiration rate, which
 365 is the sum of plant-root respiration and the respiration of various soil microbes). In panels c and
 366 d, dashed curves illustrate the alternative responses of the interaction-controlled variables to N
 367 input, depending on the relative importance of the participating species for their interactions;
 368 arrows indicate critical stages and transitions in the response curves, which should ideally be
 369 captured in experiments aiming to fully reveal the changes in responses.

370

371 4.2 Effects of N deposition and forest N status on soil CH₄ flux

372 The global map presented in Figure 4 illustrates the distribution of soil CH₄ uptake, with
 373 hotspots predominantly located in temperate forests. This can be attributed to favorable
 374 conditions in temperate forests, such as optimal soil moisture levels for aeration and suitable
 375 temperatures for enhanced methanotrophic activity, both of which promote CH₄ uptake by soils
 376 (Castro et al., 1995). Meanwhile, soils in central Amazon rainforest, tropical forests in Southeast
 377 Asia, and boreal forests in Siberia and northwestern Canada were predicted to be CH₄ sources,
 378 which is consistent with field observations (Melling et al., 2005; Pangala et al., 2017; Rask et al.,
 379 2002). The net emission of CH₄ is probably caused by submerged soils widespread in these
 380 regions, which favors methanogenesis and hinders methanotrophy. The estimated global budget
 381 for CH₄ uptake by forest soils in this study is 11.2 TgCH₄ yr⁻¹. This aligns well with estimations
 382 from previous studies using data extrapolation or modeling approaches (as shown in Fig. 6;
 383 (Curry, 2007; Dutaur & Verchot, 2007; Potter et al., 1996; Ridgwell et al., 1999; Steudler et al.,
 384 1989; L. J. Yu et al., 2017; Zhuang et al., 2013).



385

386 **Fig. 6.** Comparison of the estimated global forest soil CH₄ uptake budgets from previous studies
387 with the findings of this study. It should be noted that the global forest area used in four earlier
388 studies ($\sim 6 \times 10^9$ ha) significantly exceeded the currently accepted value ($\sim 4 \times 10^9$ ha). To facilitate
389 accurate comparison, we rectified the estimates to account for the differences in forest area. The
390 rectified estimates are indicated with asterisks (*).

391

392 N deposition impacts the capacity of forest soils to absorb atmospheric CH₄. N deposition
393 enhances plant growth in N-limited ecosystems, leading to increased root exudates, which adds
394 to the substrates and anoxic microsites for methanogenesis. Moreover, deposited N may
395 stimulate the activity of methanogens, thereby accelerating the rate of CH₄ production. The
396 produced CH₄ can either diffuse into the soil or be released into the near-surface atmosphere
397 through tree stems and leaves. Elevated CH₄ concentrations promote methanotrophy, the process
398 of CH₄ oxidation (Carmichael et al., 2014; Covey & Megonigal, 2019; Le Mer & Roger, 2001).
399 Methanotrophs present in near-surface soils can be stimulated by atmospheric N deposition,
400 further enhancing CH₄ oxidation. On the other hand, long-term high N deposition can drive
401 forests towards a state of N saturation (Aber et al., 1998; Ågren & Bosatta, 1988). Additional N
402 input to N-saturated forests may suppress plant and microbial activities, leading to a decrease in
403 the rate of CH₄ oxidation. Furthermore, the deposited ammonium may compete with CH₄ for
404 oxidants, further reducing CH₄ uptake by soils (Schnell & King, 1994).

405 Despite of the mechanistic relevance between N saturation status and soil CH₄ uptake,
406 previous studies were unable to separately analyze the N effect on R_{CH_4} in N-limited and N-
407 saturated forests, owing to the lack of site-level N status information or a global map of the N
408 saturation status of forests. We innovatively determined the N limitation or saturation status of

409 global forest ecosystems using the sensitivity of soil N₂O emissions to N deposition as an
410 indicator, which was estimated from soil N₂O emission data measured in global N addition
411 experiments (Fig. 2a; see Supporting Text S1 for details). Globally-distributed, coordinated
412 manipulative experiments can help reveal spatially-varying sensitivities of ecosystems to
413 environmental changes, facilitating biogeochemical and global change studies.

414

415 4.3 Suppressing effect of N deposition on global soil CH₄ uptake depends on forest N status

416 Findings in this study (Fig. 2e,f; Fig. 3) suggest that the current level of N deposition (<
417 40 kgN ha⁻¹ yr⁻¹ in the majority of forests) primarily stimulates soil CH₄ uptake in N-limited
418 forests whereas suppressing soil CH₄ uptake in N-saturated forests. Globally, we revealed that N
419 deposition decreased forest soil CH₄ uptake. However, the extent of this suppression effect varies
420 across different biomes depending on the N limitation or saturation status of the forests.

421 The most pronounced suppression effect was observed in temperate forests (Table 1),
422 likely due to the transition of many forests in this region from a N-limited to a N-saturated status
423 caused by N deposition. At this stage, N input suppresses CH₄ uptake (refer to Supporting Fig.
424 S5). In contrast, tropical forests naturally exist in or near N saturation (Lu et al., 2021; Matson et
425 al., 2002), resulting in a weakening response of R_{CH_4} to additional N input. Boreal forests, mostly
426 N-limited by nature, exhibit a stimulated CH₄ uptake in response to N deposition (Supporting
427 Fig. S5).

428 It is important to note that maximizing soil CH₄ uptake might suggest maintaining a
429 relatively high N deposition level around the transition point between Stage I and Stage II.
430 However, this approach should consider the potential acceleration of N₂O emissions resulting

431 from N deposition (Cen et al., 2022). Further research is required to evaluate whether
432 maintaining a relatively high N deposition rate can effectively reduce combined greenhouse gas
433 emissions from soils. Additionally, the shift between N-limited and N-saturated status driven by
434 N deposition can lead to systematic changes in the structure and function of plant and microbial
435 communities. This can at least transiently reduce ecosystem resilience and increase
436 environmental risks such as species invasion. Therefore, maintaining forest ecosystems near the
437 threshold for N status change may not be a feasible strategy for climate change mitigation.

438 To ensure long-term environmental health, it is crucial to regulate and address N
439 pollution. Western Europe and the eastern United States have witnessed a decrease in
440 atmospheric N deposition due to the reduction in anthropogenic N emissions (Ackerman et al.,
441 2019). Similarly, China has stabilized its N deposition through effective N pollution control
442 measures (G. Yu et al., 2019), although many parts of the country are still under a relatively high
443 level of N deposition ($> 20 \text{ kgN ha}^{-1} \text{ yr}^{-1}$). These countries and regions have experienced
444 significant anthropogenic impacts, leading to large areas of N-saturated forests. With continued
445 efforts to control N pollution effectively, a decline in N deposition is anticipated. Consequently,
446 soils in N-saturated forests in these counties and regions are likely to absorb more atmospheric
447 CH_4 , thus contributing to global warming mitigation.

448 In this study, we computed the response factors of soil CH_4 flux to N input by utilizing
449 data from global N addition experiments. We quantified the impact of N deposition on soil CH_4
450 uptake in forests worldwide. It is important to note that the majority of the experiments were
451 conducted over a short-term period (approximately 85% of the data in the CH_4 _exp dataset
452 comprised forest sites where N addition experiments lasted no longer than 2 years). Therefore,
453 the derived response factors primarily reflect the short-term influence of N deposition on soil

454 CH₄ flux. They may not provide insights into the long-term adaptation of plants and microbes to
455 altered N deposition regimes. Consequently, our results should be interpreted as the short-term
456 direct effect of N deposition on soil CH₄ uptake. If future research aims to estimate or predict the
457 influence of N deposition on soil CH₄ uptake over a long period of time (e.g., on a centennial
458 scale), additional observational data from long-term experiments will be necessary. These data
459 should encompass the adaptive changes in soil microbial communities (especially methanogens
460 and methanotrophs), as well as the quantity and quality of plant root exudates.

461

462 **5 Conclusions**

463 Using compiled data from N addition experiments in global forests, we validated a
464 “stimulating-suppressing-weakening” (“three stages”) response pattern of soil CH₄ uptake to N
465 input, which could generalize the diverse effects of N input on soil CH₄ flux in N-limited and N-
466 saturated forests. On the basis, we quantified that on global level, current level of N deposition
467 suppressed forest soil CH₄ uptake by ~3%. The suppressing effect, however, differs among
468 biomes, because of the different proportions of N-saturated forests in different biomes. Our
469 findings imply that by controlling N pollution and reducing N deposition, soil CH₄ uptake in N-
470 saturated forests (mostly in tropical and temperate biomes) are expected to increase, potentially
471 mitigating global warming. Due to the limitations of available data, our result could only show
472 the short-term effect of N deposition on global forest soil CH₄ flux. In the future when more
473 long-term experimental data become available, researchers could further study the adaptations of
474 methanogens and methanotrophs to long-term N addition, thus improving predictions of N
475 deposition-induced change in the global methane budget.

476

477 **Acknowledgments**

478 We thank Dr. P. Vitousek for helpful comments and suggestions. We are grateful to all the
479 researchers who spent time and efforts on manipulative experiments and observations, and made
480 the data publicly accessible. This work was financially supported by the National Natural
481 Science Foundation of China (42141004, 32171544), the National Key R&D Program of China
482 (2022YFF080210102), and CAS Project for Young Scientists in Basic Research (YSBR-037).
483 KBB received funding by the Danish National Research Foundation (DNRF), Grant Number P2
484 “Pioneer Center for Research in Sustainable Agricultural Futures (Land-CRAFT). The authors
485 declare no conflict of interest.

486

487 **Open Research**

488 All the source data and R code used for data analysis in this study have been uploaded as
489 supporting information for peer review purposes, which will be archived in a publicly accessible
490 repository prior to publication.

491

492

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