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## Enhanced methane emissions from tropical wetlands during the 2011 La Niña

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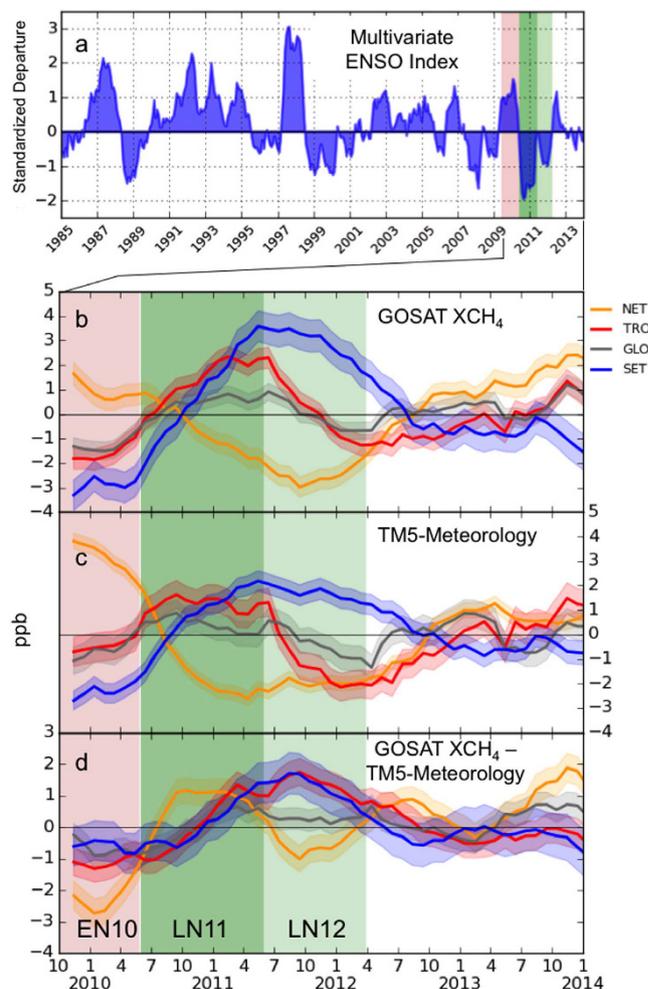
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Year-to-year variations in the atmospheric methane (CH<sub>4</sub>) growth rate show significant correlation with climatic drivers. The second half of 2010 and the first half of 2011 experienced the strongest La Niña since the early 1980s, when global surface networks started monitoring atmospheric CH<sub>4</sub> mole fractions. We use these surface measurements, retrievals of column-averaged CH<sub>4</sub> mole fractions from GOSAT, new wetland inundation estimates, and atmospheric δ<sup>13</sup>C-CH<sub>4</sub> measurements to estimate the impact of this strong La Niña on the global atmospheric CH<sub>4</sub> budget. By performing atmospheric inversions, we find evidence of an increase in tropical CH<sub>4</sub> emissions of ~6–9 TgCH<sub>4</sub> yr<sup>-1</sup> during this event. Stable isotope data suggest that biogenic sources are the cause of this emission increase. We find a simultaneous expansion of wetland area, driven by the excess precipitation over the Tropical continents during the La Niña. Two process-based wetland models predict increases in wetland area consistent with observationally-constrained values, but substantially smaller per-area CH<sub>4</sub> emissions, highlighting the need for improvements in such models. Overall, tropical wetland emissions during the strong La Niña were at least by 5% larger than the long-term mean.

CH<sub>4</sub> is the second most important anthropogenic greenhouse gas after CO<sub>2</sub>, accounting for 20% of direct anthropogenic radiative forcing<sup>1</sup>. CH<sub>4</sub> contributes strongly to anthropogenic climate change, directly through its radiative forcing as well as indirectly through impacts on atmospheric chemistry<sup>2</sup>. With a relatively short atmospheric lifetime of ~9 years, CH<sub>4</sub> is a primary target for global warming mitigation strategies<sup>3</sup>. Over the past decades, the atmospheric CH<sub>4</sub> growth rate has been highly variable<sup>3–7</sup>, including an approximate stabilization from 1999 to 2006 followed by a renewed growth since 2007<sup>8</sup>. Among a range of explanations that were proposed, some studies have suggested that more than 70% of the interannual variations of CH<sub>4</sub> can be explained by wetland CH<sub>4</sub> emissions<sup>9,10</sup>.

Wetland CH<sub>4</sub> emissions are highly sensitive to soil temperature and moisture<sup>11</sup>. Paleo records and studies of contemporary CH<sub>4</sub> suggest a strong positive feedback of wetlands to global warming through CH<sub>4</sub> emissions<sup>12,13</sup>. Proper quantification of this feedback is important for accurate future climate projections. Therefore, it is crucial to better understand the sensitivity of wetland CH<sub>4</sub> emissions to changes in climatic parameters. The El Niño Southern Oscillation (ENSO) is a major mode of variability of global precipitation and temperature, comprising alternating El Niño and La Niña phases<sup>14</sup>. Hodson *et al.*<sup>15</sup> estimated the influence of precipitation and temperature change, driven by ENSO, on wetland CH<sub>4</sub> emissions using a process-based wetland model. They found that

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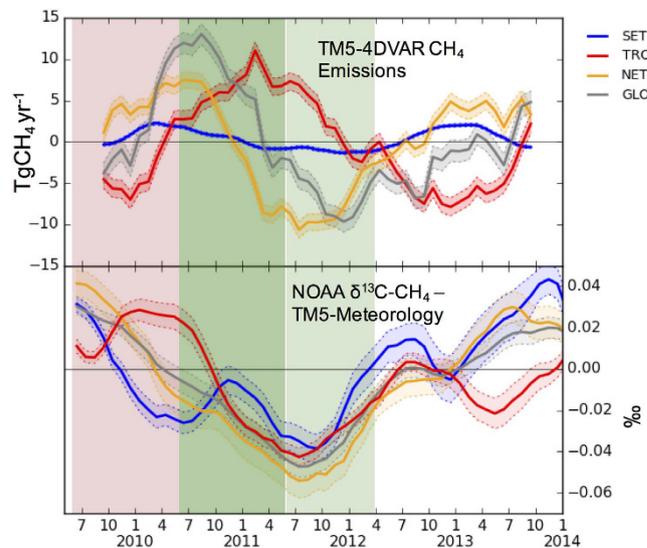
**Figure 1.** (a) Multivariate ENSO index (MEI<sup>53</sup>). The strong La Niña of 2011 (LN11) is shaded in dark green. The preceding El Niño of 2010 (EN10) and succeeding weak La Niña of 2012 (LN12) are shaded in lighter red and green colors, respectively. (b,c,d) Detrended and smoothed XCH<sub>4</sub> integrated over the large regions: (b) GOSAT FP XCH<sub>4</sub>, (c) TM5-Meteorology XCH<sub>4</sub>—that is, XCH<sub>4</sub> variability due to meteorological changes (TM5 is run with annually repeating emissions). (d) GOSAT FP XCH<sub>4</sub> corrected for the influence of meteorology (the difference between b and c). The light shaded regions represent the  $\pm 1\sigma$  uncertainty of the respective time series. NET, SET, TRO, and GLO are abbreviation of Northern Extra Tropics, Southern Extra Tropics, Tropics and Globe, respectively.

a large fraction of CH<sub>4</sub> variability is correlated with ENSO, with higher tropical wetland CH<sub>4</sub> emission during La Niña periods. However, this pattern has not been verified until now by atmospheric CH<sub>4</sub> measurements during a La Niña. Furthermore, La Niña periods have received less attention in studies of the atmospheric CH<sub>4</sub> budget than El Niño, since continued warming likely favors neutral or El Niño conditions<sup>16</sup>.

The La Niña of 2011 (LN11 hereafter) was the strongest since 1980 (see Fig. 1a) and offers the possibility to investigate the response of the atmospheric CH<sub>4</sub> budget to La Niña conditions. In this study, we investigate this response by combining different measurement dataset and model simulations. A brief overview of them is given in the next section.

## Method and Data

Atmospheric CH<sub>4</sub> measurements are available during the 2011 La Niña period from ground-based networks (NOAA-ESRL, CSIRO), and space (GOSAT, SCIAMACHY). The Greenhouse Gases Observing Satellite (GOSAT) has been measuring spectra for retrieval of the column average mole fraction of CH<sub>4</sub> (XCH<sub>4</sub>) since June 2009<sup>17</sup>. Onboard GOSAT is the Thermal And Near infrared Sensor for carbon Observation-Fourier Transform Spectrometer (TANSO-FTS), from which XCH<sub>4</sub> is obtained with high sensitivity to the lower troposphere, and hence, to surface emissions<sup>18</sup>. We analyze the interannual variability in GOSAT full-physics (FP) XCH<sub>4</sub>, obtained using the RemoteC algorithm<sup>19</sup>, and ground-based CH<sub>4</sub> flask-air measurements. Supplementary Material (SM) Section 9 further explains the FP retrieval method and justifies our choice of FP XCH<sub>4</sub> over XCH<sub>4</sub> derived from other retrieval algorithms.



**Figure 2.** (a) Detrended and smoothed  $\text{CH}_4$  surface emissions estimates from TM5-4DVAR for the same regions as in Fig. 1. The variability of GFED4s biomass burning emissions has been subtracted. (b)  $\delta^{13}\text{C}-\text{CH}_4$  measurements<sup>54</sup> corrected for the influence of transport using a meteorology-only TM5 simulation of  $\delta^{13}\text{C}-\text{CH}_4$ <sup>32</sup>. The light shaded regions represent the  $\pm 1\sigma$  uncertainty of the respective time series.

In addition to the surface emissions, changes in atmospheric transport can cause interannual variability in  $\text{CH}_4$ <sup>20</sup>. Large-scale transport patterns, including the strength of inter-hemispheric exchange and atmospheric temperature are influenced by ENSO<sup>3,21–23</sup>. To quantify the contribution of these meteorological parameters, we ran the Tracer Transport Model version 5 (TM5<sup>24</sup>) repeating surface emissions of 2008 for every year in 2009–2015. This simulation is referred to as TM5-Meteorology from hereon (see SM Section 1). To quantify the contribution of the surface emissions to  $\text{XCH}_4$  variability, we look at the difference between GOSAT FP  $\text{XCH}_4$  and  $\text{XCH}_4$  sampled from TM5-Meteorology.

Atmospheric inverse modeling systems are well established tools to convert atmospheric  $\text{CH}_4$  measurements into surface emissions<sup>25,26</sup>. We use the TM5-4DVAR (TM5-variational data assimilation system<sup>27</sup>) in combination with GOSAT FP  $\text{XCH}_4$ , and surface measurements from NOAA-ESRL<sup>3</sup> and CSIRO<sup>28</sup> to optimize surface  $\text{CH}_4$  emissions. Note that the inverse model makes use of actual meteorological fields from the ECMWF ERA-interim reanalysis to account for variability in the atmospheric transport of  $\text{CH}_4$ . Earlier studies have established the link between biomass burning  $\text{CH}_4$  emissions and ENSO<sup>29</sup>. To exclude the influence of biomass burning, fire related  $\text{CH}_4$  emissions from the Global Fire Emissions Database version 4s (GFED4s) inventory have been subtracted from the TM5-4DVAR emissions.

The origin of an atmospheric  $\text{CH}_4$  anomaly can be identified using  $\text{CH}_4$  stable isotope measurements. We look at measurements of  $^{13}\text{C}/^{12}\text{C}$  in  $\text{CH}_4$  (expressed in  $\delta$ -notation as  $\delta^{13}\text{C}-\text{CH}_4$ ) analyzed by INSTAAR in samples from the NOAA-ESRL (ref. 30, see Fig. 2b).  $\delta^{13}\text{C}-\text{CH}_4$  of atmospheric  $\text{CH}_4$  (global average in 2009 =  $-47.14\text{‰}$ ) is controlled by the relative contribution from different source types with distinct isotopic signatures. The mean isotopic signatures of the biogenic category is  $\sim -60\text{‰}$  (includes wetlands, agriculture, waste), for the thermogenic category it is  $\sim -37\text{‰}$  (includes fossil-fuels) and for pyrogenic category it is  $\sim -22\text{‰}$  (includes biomass burning)<sup>31</sup>. To account for impact of meteorological variability on  $\delta^{13}\text{C}-\text{CH}_4$ , a meteorology simulation of  $\delta^{13}\text{C}-\text{CH}_4$  was performed using TM5<sup>32</sup>.

To identify factors which might have altered the wetland emissions, we look at the variability in land precipitation and temperature data in CRU-TS version 3.23 (Climatic Research Unit-time series<sup>33</sup>). We also analyze  $\text{CH}_4$  emission and surface inundation extent from two process-based wetland models: LPJ-wsl<sup>15,34</sup> and CLM4.<sup>5,35,36</sup> Additionally, we derive an independent estimate of inundation extent from remotely sensed Surface Water Microwave Product Series (SWAMPs<sup>37</sup>).

The primary sink of  $\text{CH}_4$  is the reaction with OH in the troposphere ( $\sim 454\text{--}617 \text{ TgCH}_4 \text{ yr}^{-1}$ ), and inter-annual variations in OH can also contribute to the observed  $\text{CH}_4$  variability. Tropospheric OH concentrations are influenced by many factors, including temperature, water vapor,  $\text{O}_3$ ,  $\text{NO}_x$ ,  $\text{CH}_4$ , CO, and the overhead stratospheric ozone column<sup>38</sup>. The TM5-4DVAR inversions performed in this study make use of OH fields from ref. 39, which vary seasonally, but are the same each year. To investigate possible variations caused by the OH sink, we analyze posterior  $\text{CH}_4$  emissions of LMDz-PYVAR-SACS inversion<sup>40–42</sup>. In this inversion, the OH fields were optimized simultaneously using methyl chloroform (MCF) measurements (see SM Section 1.3).

**Data Analysis.** The above mentioned measurements and model outputs have been analyzed by taking their monthly averages and integrating them over three zones over the globe (GLO): Tropics (TRO:  $30^\circ\text{S}$  to  $30^\circ\text{N}$ ), Northern Extra Tropics (NET:  $30^\circ\text{N}$  to  $90^\circ\text{N}$ ), Southern Extra Tropics (SET:  $90^\circ\text{S}$  to  $30^\circ\text{S}$ ). The time series of these monthly averages have been detrended and smoothed using a 12 month running mean. The resulting

Region	Phase	$\Delta_r^q$			$\mu_r^q$	$\Delta_r^q$
		GOSAT (ppb)	TM5 (ppb)	GOSAT-TM5 (ppb)	TM5-4DVAR (TgCH <sub>4</sub> yr <sup>-1</sup> )	$\delta^{13}\text{C-CH}_4$ (‰)
NET:	EN10	-0.85	-1.60	0.74	4.58	-0.05
	LA11	-2.66	-3.86	1.20	0.01	-0.03
	LA12	0.31	0.38	-0.08	-8.91	0.03
TRO:	EN10	0.86	0.61	0.24	-5.76	0.01
	LA11	2.42	0.38	2.04	5.94	-0.06
	LA12	-3.57	-3.32	-0.25	3.94	0.02
SET:	EN10	0.55	0.77	-0.23	0.21	-0.06
	LA11	5.62	3.70	1.92	0.63	-0.01
	LA12	-1.73	-0.78	-0.94	-1.14	0.03
GLO:	EN10	0.56	1.18	-0.61	-0.96	-0.03
	LA11	0.91	-0.50	1.41	7.58	-0.03
	LA12	-1.59	-1.54	0.05	-6.11	0.03

**Table 1.**  $\Delta_r^q$  (i.e. the sum of the derivative) or  $\mu_r^q$  (i.e. mean) of the times series of quantity  $q$ , averaged over region  $r$ , as shown in Figs 1 and 2 during different ENSO phases. Please note that the GOSAT and TM5-4DVAR time series do not cover the whole EN10 period, as continuous GOSAT measurements are only available since June 2009, and the 12-month smoothing causes data points loss. Only  $\delta^{13}\text{C-CH}_4$  values cover the whole EN10.

time series of GOSAT FP XCH<sub>4</sub> from this method is less influenced by systematic errors, which often affect the satellite retrievals<sup>9,43,44</sup>.

Further, based on the Multivariate ENSO index (MEI) we define three periods: 1) La Niña in 2011 as LN11; 2) The preceding El Niño of 2010 as EN10; 3) succeeding weak La Niña of 2012 as LN12 (see Fig. 1a). Time series of measurements and model outputs are analyzed during these periods using the following parameters:

1.  $\Delta_r^q$ : Defined as the change of a quantity  $q$  in region  $r$  during an ENSO phase.
2.  $\mu_r^q$ : Defined as the mean of a quantity  $q$  in region  $r$  during an ENSO phase.

Table 1 summarizes  $\Delta_r^q$  or  $\mu_r^q$  of time series shown in Figs 1 and 2.

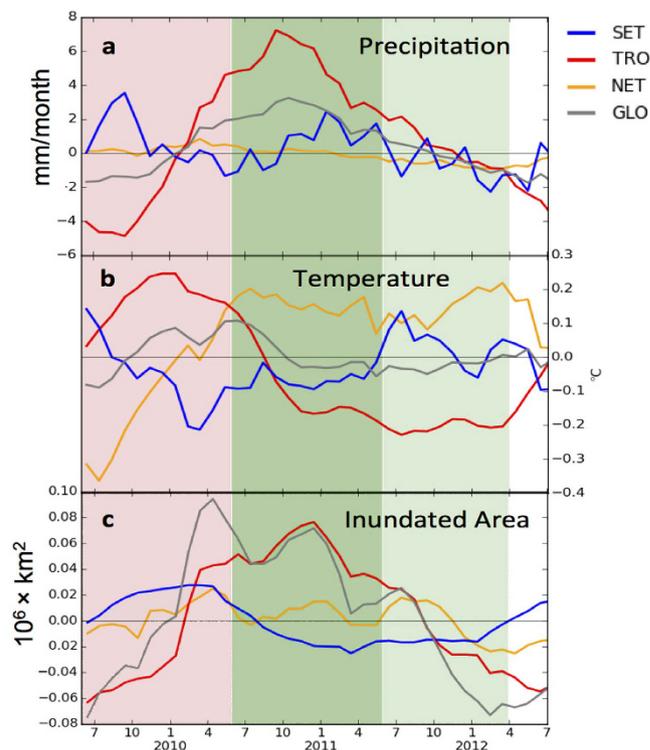
## Results and Discussion

**GOSAT observations.** Figure 1b shows detrended and smoothed time series of GOSAT FP XCH<sub>4</sub>. During LN11, SET and TRO have increasing XCH<sub>4</sub> ( $\Delta_{\text{SET}}^{\text{XCH}_4} = 5.6$  ppb and  $\Delta_{\text{TRO}}^{\text{XCH}_4} = 2.4$ ) contrasted by a decrease in NET ( $\Delta_{\text{NET}}^{\text{XCH}_4} = -2.7$  ppb). During LN12, the opposite trends are found: XCH<sub>4</sub> gradually reduced in SET and TRO and increased in NET.

Figure 1c shows the results of TM5-Meteorology simulation. The modeled XCH<sub>4</sub> decreases over NET and increases over SET ( $\Delta_{\text{SET}}^{\text{XCH}_4} = 3.7$  ppb,  $\Delta_{\text{NET}}^{\text{XCH}_4} = -3.9$  ppb). These trends can be attributed to the faster inter-hemispheric exchange during La Niña conditions, which transferred additional CH<sub>4</sub> rich air from the Northern Hemisphere to the Southern Hemisphere. Francey *et al.*<sup>21</sup> found the strongest inter-hemispheric transport of CH<sub>4</sub> during LN11 since 1990. Previous studies have reported such enhancement in inter-hemispheric transport, and consequent increase in CH<sub>4</sub> in the Southern Hemisphere, during the La Niña of 2007–2008<sup>3</sup> and 1989<sup>23</sup>. During LN12, the strength of anomalies over both regions weakens as inter-hemispheric exchange returns to its normal strength. Modeled XCH<sub>4</sub> over TRO show less variation during LN11. During LN12, the modeled XCH<sub>4</sub> ( $\Delta_{\text{TRO}}^{\text{XCH}_4} = -3.3$  ppb) explains a major fraction of GOSAT XCH<sub>4</sub> ( $\Delta_{\text{TRO}}^{\text{XCH}_4} = -3.6$  ppb) variability. It is noteworthy that OH concentrations are not affected in our meteorology simulation as TM5 uses OH fields from Spivakovsky *et al.*<sup>39</sup>. Eventhough these fields don't vary interannually, temperature variations can still cause variability in the atmospheric CH<sub>4</sub> sink, as the rate constant for reaction with OH is temperature dependent<sup>45</sup>.

Figure 1d shows the GOSAT FP XCH<sub>4</sub> after correction for variations in atmospheric transport using the TM5-Meteorology simulation, and indicates the fraction of CH<sub>4</sub> variability that can be attributed to variability in CH<sub>4</sub> sources and sinks. An analogous plot showing surface flask-air measurements is given in the SM section 4 and shows similar patterns. During LN11, the largest transport-corrected  $\Delta_{\text{TRO}}^{\text{XCH}_4}$  is seen in TRO ( $\Delta_{\text{TRO}}^{\text{XCH}_4} = 2.0$  ppb), pointing to higher CH<sub>4</sub> emissions from the tropical continents. As SET does not have large CH<sub>4</sub> surface emissions, it is caused most likely by transport from TRO. This means the transport-corrected anomaly of TRO is transferred to SET.

**Emissions and source attribution.** Detrended and smoothed time series of the posterior TM5-4DVAR emissions are shown in Fig. 2a. The CH<sub>4</sub> emissions over TRO show a positive anomaly during LN11 ( $\mu_{\text{TRO}}^{\text{emission}} = 5.9$  TgCH<sub>4</sub> yr<sup>-1</sup>).  $\mu_{\text{TRO}}^{\text{emission}}$  is higher by 11.7 TgCH<sub>4</sub> yr<sup>-1</sup> in LN11 than in the preceding EN10 ( $\mu_{\text{TRO}}^{\text{emission}} = -5.8$  TgCH<sub>4</sub> yr<sup>-1</sup>). This can be due to higher CH<sub>4</sub> emissions from tropical wetlands during the La Niña, as suggested by ref. 15. During LN12, the CH<sub>4</sub> emission enhancement over TRO is weaker ( $\mu_{\text{TRO}}^{\text{emission}} = 3.9$



**Figure 3.** (a,b) Detrended and smoothed regionally averaged precipitation and temperature measurements over land in CRU-TS version 3.23 (Climatic Research Unit-time series<sup>33</sup>). (c) Anomalies in the total inundated area estimated by SWAMPS<sup>37</sup>.

TgCH<sub>4</sub> yr<sup>-1</sup>). NET has a sharp decrease in CH<sub>4</sub> emissions during LN11 ( $\Delta_{\text{NET}}^{\text{emission}} = -13 \text{ TgCH}_4 \text{ yr}^{-1}$ ), which shifts the maximum of the global CH<sub>4</sub> emission anomaly towards the beginning of LN11.

Detrended and smoothed time series of  $\delta^{13}\text{C}-\text{CH}_4$  is shown in Fig. 2b. During LN11,  $\delta^{13}\text{C}-\text{CH}_4$  decreased over each region. Over TRO, we observe a decrease of 0.06‰, which is a larger decrease than the GLO  $\delta^{13}\text{C}-\text{CH}_4$  decrease by 0.03‰. An isotope mass balance calculation shows that if the increase in TRO CH<sub>4</sub> emissions of 11.7 TgCH<sub>4</sub> yr<sup>-1</sup> (change from EN10 with  $\mu_{\text{TRO}}^{\text{emission}} = -5.8 \text{ TgCH}_4 \text{ yr}^{-1}$  to LN11 with  $\mu_{\text{TRO}}^{\text{emission}} = 5.9 \text{ TgCH}_4 \text{ yr}^{-1}$ ), is attributed to a biogenic source, it would cause a drop in  $\delta^{13}\text{C}-\text{CH}_4$  of similar magnitude. This indicates that the source of the LN11 CH<sub>4</sub> anomaly in the Tropics is of biogenic origin. The increase of GLO  $\delta^{13}\text{C}-\text{CH}_4$  ( $\approx 0.03\text{‰}$ ) over LN12 can be explained by reduced biogenic emissions ( $\mu_{\text{GLO}}^{\text{emission}}(\text{LN12}) - \mu_{\text{GLO}}^{\text{emission}}(\text{LN11}) = \sim -14 \text{ TgCH}_4 \text{ yr}^{-1}$ ) and increased biomass burning emissions ( $\mu_{\text{GLO}}^{\text{GFED4s}}(\text{LN12}) - \mu_{\text{GLO}}^{\text{GFED4s}}(\text{LN11}) = 2.2 \text{ TgCH}_4 \text{ yr}^{-1}$ ) in comparison to LN11.

Tropical Biomass burning is strongly influenced by ENSO<sup>46</sup>. Globally, GFED4s biomass burning CH<sub>4</sub> emissions indicate a decrease of 0.67 TgCH<sub>4</sub> yr<sup>-1</sup> from EN10 to LN11 (see SM Section 5). The effect of this change on the isotopic composition is only  $-0.003\text{‰}$ , thus much smaller than the observed trend. According to GFED4s, CH<sub>4</sub> emissions from biomass burning over TRO during LN11 are close to average over the whole period ( $\mu_{\text{TRO}}^{\text{GFED4s}} = -0.23 \text{ TgCH}_4 \text{ yr}^{-1}$ ). In LN12, these emissions were higher in NET and TRO ( $\mu_{\text{TRO}}^{\text{GFED4s}} = 1.09 \text{ TgCH}_4 \text{ yr}^{-1}$  and  $\mu_{\text{NET}}^{\text{GFED4s}} = 1.32 \text{ TgCH}_4 \text{ yr}^{-1}$ ). This increase may be explained by higher fuel availability due to enhanced biomass growth during the preceding LN11. Ref. 47 suggested a similar impact of Australian biomass burning on CO<sub>2</sub> emissions.

Figure 3a and b show monthly anomalies recorded in climate parameters. A significant redistribution of heat and precipitation is seen during the different phases of ENSO.  $\mu_{\text{TRO}}^{\text{precipitation}}$  was  $-1.72$ ,  $4.90$ , and  $0.64 \text{ mm}$  during EN10, LN11, and LN12, respectively. During LN11 the precipitation anomaly in TRO (and in GLO) was the highest since the onset of the 21st century (see SM Figure 10). Regions like Australia had six consecutive seasons of increased rainfall over the La Niña of 2011 and 2012<sup>48</sup>. Higher temperatures were observed in NET during LN11 ( $\mu_{\text{NET}}^{\text{temperature}} = 0.18 \text{ °C}$ ), favoring increased biomass burning, for example, near Moscow during the summer of 2010<sup>49,50</sup>. Mean temperatures during LN11 ( $\mu_{\text{TRO}}^{\text{temperature}} = -0.05 \text{ °C}$ ) were in between those during EN10 ( $\mu_{\text{TRO}}^{\text{temperature}} = 0.15 \text{ °C}$ ) and LN12 ( $\mu_{\text{TRO}}^{\text{temperature}} = -0.22 \text{ °C}$ ).

An increase in total inundated area is observed in the remotely sensed SWAMPS data (see Fig. 3c). The total inundated area estimated by the wetland models LPJ-wsl and CLM4.5 also show a similar increase. However, these wetland models estimate a relatively weaker enhancement in CH<sub>4</sub> emissions with  $\mu_{\text{TRO}}^{\text{emission}} = 1.54 \text{ TgCH}_4 \text{ yr}^{-1}$  for LPJ-wsl and  $\mu_{\text{TRO}}^{\text{emission}} = 2.38 \text{ TgCH}_4 \text{ yr}^{-1}$  for CLM4.5 during LN11 (see SM Section 8).

To further investigate the relation between inversion-estimated CH<sub>4</sub> emissions and potential climatic drivers, we examine their correlation coefficients (R) [see SM Figure 11]. CH<sub>4</sub> emission anomalies (as shown in

Fig. 2a) correlate stronger with precipitation anomalies than with temperature anomalies (as shown in Fig. 3) in both NET  $R_{\text{precipitation,emission}} = 0.86$ ,  $R_{\text{temperature,emission}} = -0.52$ ) and TRO ( $R_{\text{precipitation,emission}} = 0.85$ ,  $R_{\text{temperature,emission}} = -0.62$ ). This points to precipitation as the more important driver of the CH<sub>4</sub> anomaly in TRO during LN11, supported further by the correlation with inundated area ( $R_{\text{inundation,emission}} = 0.67$ ). This is consistent with the findings of Bloom *et al.*<sup>51</sup>, who show that precipitation plays a more dominant role than temperature in determining anomalous CH<sub>4</sub> variability in the Tropics.

To investigate possible variations caused by the OH sink, we analyzed optimized CH<sub>4</sub> emissions from a LMDz-PYVAR-SACS inversion, in which OH fields were also optimized. During LN11, the results of this inversion suggest  $\mu_{\text{TRO}}^{\text{emission}}$  of 9.1 TgCH<sub>4</sub> yr<sup>-1</sup>, compared to TM5-4DVAR  $\mu_{\text{TRO}}^{\text{emission}}$  of ~6 TgCH<sub>4</sub> yr<sup>-1</sup> (see SM Section 6). The differences in interannual variations of the emission estimates of the two inversions are mainly caused by their different treatment of OH sink. Assuming that the MCF-optimized OH sink of LMDz-PYVAR-SACS is more accurate than TM5, the sink was stronger than normal during LN11 by ~3 TgCH<sub>4</sub> yr<sup>-1</sup>. This is consistent with the hypothesis of an increased CH<sub>4</sub> sink during La Niña and a weaker sink during El Niño<sup>52</sup>.

## Conclusion

Our inversion results, supported by  $\delta^{13}\text{C}-\text{CH}_4$  measurements, provide strong evidence of enhanced tropical biogenic CH<sub>4</sub> emissions by ~6–9 TgCH<sub>4</sub> yr<sup>-1</sup>, during the La Niña of 2011. Wetlands were the likely cause of this anomaly as a simultaneous increase in total inundated area is shown by remote sensing observations and hydrological models. The increase in inundated area was in response to La Niña induced increase in precipitation. 2011 experienced the strongest La Niña event in the past 4 decades as well as since the onset of modern atmospheric CH<sub>4</sub> measurements. It is noteworthy that during this La Niña the increase in global CH<sub>4</sub> mole fractions were not as pronounced due to a simultaneous decrease in the CH<sub>4</sub> emissions in the Northern Extra Tropics. Our analysis presents the first evidence of the large-scale response of wetland CH<sub>4</sub> emissions to ENSO variability using satellite retrievals.

## Data Availability

We use Level 2 SRFP XCH4 v2.3.7 GOSAT XCH4 retrievals that are publicly available from ESA's Climate Change Initiative website ([www.esa-ghg-cci.org/](http://www.esa-ghg-cci.org/)). NOAA CH4 and INSTAAR  $\delta^{13}\text{C}-\text{CH}_4$  measurements are freely available from NOAA's public ftp server (<ftp://aftp.cmdl.noaa.gov/data>). CSIRO CH4 measurements can be downloaded from the WDCGG (World Data Centre for Green-house Gases) website. GFED4s CH4 emissions can be downloaded from <http://daac.ornl.gov>. CRU TS3.23). Precipitation and temperature data are held at British Atmospheric Data Centre, RAL, UK (<http://badc.nerc.ac.uk/data/cru/>). SWAMP wetlands fraction data can be downloaded from <http://wetlands.jpl.nasa.gov> after a short registration. CLM4.5 and LPJ-wsl CH4 emissions and wetlands fractions can be obtained by contacting William J. Riley and B. Poulter, respectively.

## References

- Myhre, G. *et al.* Anthropogenic and Natural Radiative Forcing. In Stocker, T. F. D., Qin, G.-K., Plattner, M., Tignor, S. K., Allen, J., Boschung, A., Nauels, Y., Xia, V. B. & (eds) J., P. M. (eds) *Climate Change 2013: The Physical Science Basis. Contribution of Working Group I to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change*, 659–740 (Cambridge University Press, Cambridge, United Kingdom and New York, NY, USA, 2013).
- Kirschke, S. *et al.* Three decades of global methane sources and sinks. *Nature Geoscience* **6**, 813–823 (2013).
- Dlugokencky, E. J. *et al.* Observational constraints on recent increases in the atmospheric CH<sub>4</sub> burden. *Geophysical Research Letters* **36**, L18803 (2009).
- Heimann, M. Atmospheric science: Enigma of the recent methane budget. *Nature* **476**, 157–158 (2011).
- Kai, F. M., Tyler, S. C., Randerson, J. T. & Blake, D. R. Reduced methane growth rate explained by decreased Northern Hemisphere microbial sources. *Nature* **476**, 194–7 (2011).
- Aydin, M. *et al.* Recent decreases in fossil-fuel emissions of ethane and methane derived from firn air. *Nature* **476**, 198–201 (2011).
- Ferretti, D. F. *et al.* Unexpected changes to the global methane budget over the past 2000 years. *Science (New York, N.Y.)* **309**, 1714–7 (2005).
- Nisbet, E., Dlugokencky, E. J. & Bousquet, P. Methane on the Rise—Again. *Science* **343**, 493–495 (2014).
- Bousquet, P. *et al.* Contribution of anthropogenic and natural sources to atmospheric methane variability. *Nature* **443**, 439–43 (2006).
- Chen, Y. H. & Prinn, R. G. Estimation of atmospheric methane emissions between 1996 and 2001 using a three-dimensional global chemical transport model. *Journal of Geophysical Research Atmospheres* **111**, 1–25 (2006).
- Christensen, T. R. Factors controlling large scale variations in methane emissions from wetlands. *Geophysical Research Letters* **30**, 10–13 (2003).
- Nisbet, E. G. & Chappellaz, J. Shifting Gear, Quickly. *Science* **324**, 477–478 (2009).
- Petrenko, V. V. *et al.* <sup>14</sup>CH<sub>4</sub> Measurements in Greenland Ice: Investigating Last Glacial Termination CH<sub>4</sub> Sources. *Science* **324**, 506–508 (2009).
- Walker, G. & Bliss, E. World weather V. *Memoirs of the Royal Meteorological Society* **4**, 53–84 (1932).
- Hodson, E. L., Poulter, B., Zimmermann, N. E., Prigent, C. & Kaplan, J. O. The El Niño–Southern Oscillation and wetland methane interannual variability. *Geophysical Research Letters* **38**, 3–6 (2011).
- Cai, W. *et al.* Increasing frequency of extreme El Niño events due to greenhouse warming. *Nature Climate Change* **5**, 1–6 (2014).
- Yokota, T. *et al.* Global Concentrations of CO<sub>2</sub> and CH<sub>4</sub> Retrieved from GOSAT: First Preliminary Results. *SOLA* **5**, 160–163 (2009).
- Kuze, A., Suto, H., Nakajima, M. & Hamazaki, T. Thermal and near infrared sensor for carbon observation Fourier-transform spectrometer on the Greenhouse Gases Observing Satellite for greenhouse gases monitoring. *Applied optics* **48**, 6716–33 (2009).
- Butz, A., Hasekamp, O. P., Frankenberg, C., Vidot, J. & Aben, I. CH<sub>4</sub> retrievals from space-based solar backscatter measurements: Performance evaluation against simulated aerosol and cirrus loaded scenes. *Journal of Geophysical Research* **115**, D24302 (2010).
- Warwick, N., Bekki, S., Kaw, K., Nisbet, E. & Pyle, J. The impact of meteorology on the interannual growth rate of atmospheric methane. *Geophysical Research Letters* **29**, 1947 (2002).
- Francey, R. J. & Frederiksen, J. S. The 2009–2010 step in atmospheric CO<sub>2</sub> interhemispheric difference. *Biogeosciences* **13**, 873–885 (2016).
- Prinn, R. *et al.* Global average concentration and trend for hydroxyl radicals deduced from ALE/GAGE trichloroethane (methyl chloroform) data for 1978–1990. *Journal of Geophysical Research* **97**, 2445–2461 (1992).

23. Steele, L. *et al.* Slowing down of the global accumulation of atmospheric methane during the 1980s. *Nature* **358**, 313–316 (1992).
24. Krol, M. *et al.* The two-way nested global chemistry-transport zoom model TM5: algorithm and applications. *Atmospheric Chemistry and Physics* **5**, 417–432 (2005).
25. Hein, R., Crutzen, P. J. & Heimann, M. An inverse modeling approach to investigate the global atmospheric methane cycle. *Global Biogeochemical Cycles* **11**, 43–76 (1997).
26. Houweling, S., Kaminski, T., Dentener, F., Lelieveld, J. & Heimann, M. Inverse modeling of methane sources and sinks using the adjoint of a global transport model. *Journal of Geophysical Research* **104**, 26137–26160 (1999).
27. Meirink, J. F., Bergamaschi, P. & Krol, M. C. Four-dimensional variational data assimilation for inverse modelling of atmospheric methane emissions: method and comparison with synthesis inversion. *Atmospheric Chemistry and Physics Discussions* **8**, 12023–12052 (2008).
28. Francey, R. J., Steele, L. P., Langenfelds, R. L. & Pak, B. C. High Precision Long-Term Monitoring of Radiatively Active and Related Trace Gases at Surface Sites and from Aircraft in the Southern Hemisphere Atmosphere. *Journal of Atmospheric Sciences* **56**, 279–285 (1999).
29. Worden, J. *et al.* El Nino, the 2006 Indonesian peat fires, and the distribution of atmospheric methane. *Geophysical Research Letters* **40**, 4938–4943 (2013).
30. Miller, J. B. *et al.* Development of analytical methods and measurements of <sup>13</sup>C/<sup>12</sup>C in atmospheric CH<sub>4</sub> from the NOAA-CMDL global air sampling network. *J. Geophys. Res.* **107** (2002).
31. Dlugokencky, E. J., Nisbet, E. G., Fisher, R. & Lowry, D. Global atmospheric methane: budget, changes and dangers. *Philosophical transactions. Series A, Mathematical, physical, and engineering sciences* **369**, 2058–2072 (2011).
32. Monteil, G. *et al.* Interpreting methane variations in the past two decades using measurements of CH<sub>4</sub> mixing ratio and isotopic composition. *Atmospheric Chemistry and Physics* **11**, 9141–9153 (2011).
33. Harris, I., Jones, P., Osborn, T. & Lister, D. Updated high-resolution grids of monthly climatic observations - the CRU TS3.10 Dataset. *International Journal of Climatology* **34**, 623–642 (2014).
34. Zhang, Z., Zimmermann, N. E., Kaplan, J. O. & Poulter, B. Modeling spatiotemporal dynamics of global wetlands: Comprehensive evaluation of a new sub-grid TOPMODEL parameterization and uncertainties. *Biogeosciences* **13**, 1387–1408 (2016).
35. Riley, W. J. *et al.* Barriers to predicting changes in global terrestrial methane fluxes: Analyses using CLM4Me, a methane biogeochemistry model integrated in CESM. *Biogeosciences* **8**, 1925–1953 (2011).
36. Xu, X. *et al.* A multi-scale comparison of modeled and observed seasonal methane cycles in northern wetlands. *Biogeosciences* **13**, 5043–5056 (2016).
37. Schroeder, R. *et al.* Development and evaluation of a multi-year fractional surface water data set derived from active/passive microwave remote sensing data. *Remote Sensing* **7**, 16688–16732 (2015).
38. Dalsoren, S. *et al.* Atmospheric methane evolution the last 40 years. *Atmospheric Chemistry and Physics Discussions* **16**, 3099–3126 (2016).
39. Spivakovsky, C. M. *et al.* Three-dimensional climatological distribution of tropospheric OH: Update and evaluation. *Journal of Geophysical Research-Atmospheres* **105**, 8931–8980 (2000).
40. Locatelli, R., Bousquet, P., Saunio, M., Chevallier, F. & Cressot, C. Sensitivity of the recent methane budget to LMDz sub-grid-scale physical parameterizations. *Atmospheric Chemistry and Physics* **15**, 9765–9780 (2015).
41. Chevallier, F. *et al.* Inferring CO<sub>2</sub> sources and sinks from satellite observations: Method and application to TOVS data. *Journal of Geophysical Research: Atmospheres* **110**, 1–13 (2005).
42. Hourdin, F. *et al.* The LMDZA general circulation model: climate performance and sensitivity to parametrized physics with emphasis on tropical convection. *Climate Dynamics* **27**, 787–813 (2006).
43. Basu, S. *et al.* Global CO<sub>2</sub> fluxes estimated from GOSAT retrievals of total column CO<sub>2</sub>. *Atmospheric Chemistry and Physics* **13**, 8695–8717 (2013).
44. Schepers, D. *et al.* Methane retrievals from Greenhouse Gases Observing Satellite (GOSAT) shortwave infrared measurements: Performance comparison of proxy and physics retrieval algorithms. *Journal of Geophysical Research* **117**, D10307 (2012).
45. Sander, S. P. *et al.* Chemical Kinetics and Photochemical Data for Use in Atmospheric Studies: Evaluation Number 14. *JPL Publication 02-25* **14**, 1–334 (2003).
46. van der Werf, G. R. *et al.* Interannual variability in global biomass burning emissions from 1997 to 2004. *Atmospheric Chemistry and Physics* **6**, 3423–3441 (2006).
47. Detmers, R. G. *et al.* Anomalous carbon uptake in Australia as seen by GOSAT. *Geophysical Research Letters* **42**, 8177–8184 (2015).
48. Poulter, B. *et al.* Contribution of semi-arid ecosystems to interannual variability of the global carbon cycle. *Nature* **509**, 600–603 (2014).
49. Kononov, I. B., Beekmann, M., Kuznetsova, I. N., Yurova, A. & Zvyagintsev, A. M. Atmospheric impacts of the 2010 Russian wildfires: Integrating modelling and measurements of an extreme air pollution episode in the Moscow region. *Atmospheric Chemistry and Physics* **11**, 10031–10056 (2011).
50. Krol, M. C. *et al.* Correction to “Interannual variability of carbon monoxide emission estimates over South America from 2006 to 2010”. *Journal of Geophysical Research: Atmospheres* **118**, 5061–5064 (2013).
51. Bloom, A. A., Palmer, P. I., Fraser, A., Reay, D. S. & Frankenberg, C. Large-Scale Controls of Methanogenesis Inferred from Methane and Gravity Spaceborne Data. *Science* **327**, 322–327 (2010).
52. Elshorbany, Y. F., Duncan, B. N., Strode, S. A., Wang, J. S. & Kouatchou, J. The description and validation of the computationally Efficient CH<sub>4</sub>-CO-OH (ECCOHv1.01) chemistry module for 3-D model applications. *Geoscientific Model Development* **9**, 799–822 (2016).
53. Wolter, K. The Southern Oscillation in Surface Circulation and Climate over the Tropical Atlantic, Eastern Pacific, and Indian Oceans as Captured by Cluster Analysis (1987).
54. Fisher, R., Lowry, D., Wilkin, O., Sriskantharajah, S. & Nisbet, E. G. High-precision, automated stable isotope analysis of atmospheric methane and carbon dioxide using continuous-flow isotope-ratio mass spectrometry. *Rapid communications in mass spectrometry: RCM* **20**, 200–8 (2006).

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## Author Contributions

S.P., S.H., T.R., I.A. and M.C.K. conceived the research. P.B., S.P., N.B., G.M., W.J.R., X.X., B.P. and Z.Z. developed and ran the computations models. J.W.C.W., E.J.D., R.D., I.A., K.C.M. and O.T. provided data. S.P. and M.C.K.

analyses the data and model outputs. S.P., S.H., M.C.K. and T.R. wrote the manuscript with suggestion from all the authors.

### Additional Information

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