

Supplementary Information A and B

A. Methods Extensions

S1. Redox titration

Redox titrations quantified the oxygen demand (dO_2) for sample combustion, reflecting the O_2 preserved by carbon burial. The method closely follows the protocol outlined in (1). Briefly, after sieving (250 μm) and homogenizing the freeze-dried peat material, an aliquot is placed in a quartz tube enclosing a container filled with a large excess of CuO— the oxygen donor. The reactor is then flame-sealed under vacuum (ca. 5 μbar) and gradually heated to 900 $^{\circ}C$ where it is left to react for 10 hours. At these nominal experimental conditions, the CuO-Cu₂O redox couple buffers the partial pressure of O_2 in the system (2) at $pO_2 \sim 10^{-6}$ bar within minutes. These conditions are sufficient to oxidize most redox active elements present in the sample, in particular C, S and Fe (1). A sample is considered fully oxidized when all of its elements have reached their highest oxidation state, C^{4+} , Fe^{3+} and S^{6+} . Weighing the O donor (CuO) prior to, and after, combustion provides a robust quantification of O consumption. This weight loss, corresponding to total mass of O consumed (nO mol), provides the absolute redox capacities, $\Delta O_2 = \frac{1}{2} nO$ (mol), and mass specific redox capacities, $dO_2 = \Delta O_2$ (mol)/ m (g), where m is the mass of the sample. Precision measured by repeated measurements of known standards is typically within 5% (1). The metric dO_2^* is obtained by normalizing the absolute redox capacity ΔO_2 to TOC ($dO_2^* = 100 \times \Delta O_2 / TOC$), not sample mass. Mass of O_2 reservoirs is provided in Pg (\equiv Gt), or Tg (\equiv Mt).

S2. Redox Stoichiometry:

The redox stoichiometry (0.9–1.15 mol O_2 /mol C) was calculated as:

$$O_2:C = 12 \times dO_2^*$$

where dO_2^* is in mol O_2 /g TOC (0.08–0.095), multiplied by the molar mass of C ($M_C = 12$ g/mol). This yields:

$$O_2:C \approx 0.9\text{--}1.15$$

This ratio reflects peat's oxygen-rich composition (e.g., cellulose, lignins), reducing external O_2 demand compared to pure carbon ($O_2:C = 1$). Unlike photosynthetic $O_2:C$ ratios (~ 1 for glucose), this method captures burial-specific stoichiometry, a novel contribution to atmospheric O_2 modeling.

S3. Novelty of Redox Stoichiometry

The redox stoichiometry is a novel metric, distinct from photosynthetic or biomass-based ratios, as it quantifies O_2 spared by burial. Unlike carbon-focused studies (3), this approach directly links terrestrial redox processes to atmospheric O_2 , offering new constraints for models (e.g., GEOCARBSULF;(4)).

S4. Materials and Thermal characteristics

The IMB100 peat core

The Imbonga (IMB100) peat core was recovered in 2017 from the Ingende Territory (Equateur Province) in the Democratic Republic of Congo (DRC) at the northern edge of a small basin filled with peat, around 800 m away from the Momboyo River, a tributary of the Ruki River (0.856693°S and 19.808078°E, 320 m; Fig. 1A). The sedimentology, age/depth model (^{14}C), and palaeo-environmental context inherent to core IMB100 are described extensively in (5). In summary, a total of 17 fine fraction samples ($<150\ \mu m$) were used to construct the age-depth model using rbacon (version 2.7) in R (version 4.1.0) (6). ^{14}C dates were calibrated using a mixed curve 50%:50% of north-south air-mass mixing derived from both the Northern Hemisphere curve (IntCal20, (7)) and the Southern Hemisphere curve (SHCal20, (8)).

The 237 cm-long IMB100 core is characterized by distinct phases of peat accumulation, initially transitioning from an organic carbon-rich floodplain to a forested marshland, which later became

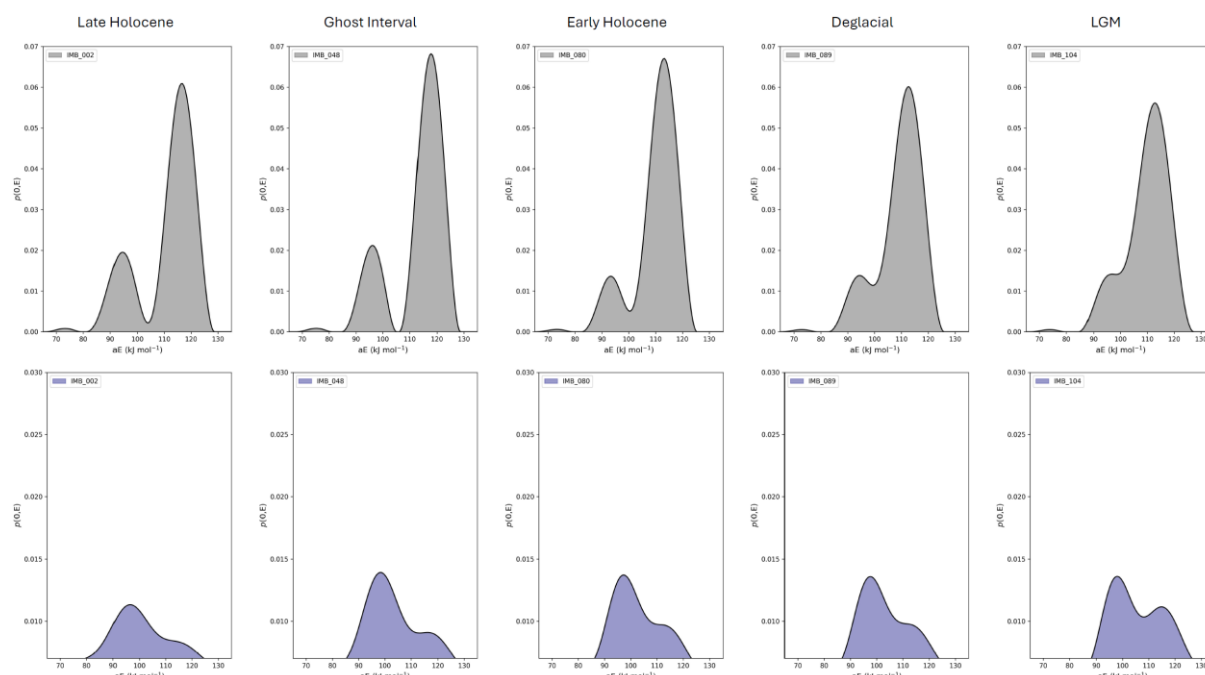
disconnected from the river, and finally to a closed-canopy swamp forest. Peat development began at ~10.6 kyr BP in response to increasing precipitation as inferred from plant wax hydrogen isotope (δD_n -C29) data (5). By ~8.7 kyr BP, the terrestrialization of the peatland coincided with the establishment of a closed canopy forest. Before ~10.6 kyr BP, old (of up to 38 kyr BP) and scattered radiocarbon (^{14}C) ages suggest that allochthonous material was incorporated into the fluvial sediments making the chronology uncertain (5). Between 124 cm and 100 cm (i.e. 7.5 kyr - 2.3 kyr BP), the slope of the modelled age/depth profile is 6 to 10 times lower than in the peat sections immediately below and above. This interval has been referred as the "Ghost Interval", an interval of extensively decomposed peat material observed across the central Congo peatlands (9)

TGA. The thermal and calorimetric properties of the samples were measured using a hybridized TGA/DSC-microGC system at the University of Lausanne (UNIL). The TGA/DSC3+ (Mettler Toledo) is coupled to a 990 microGC (Agilent) via a transfer line maintained at 80 °C. Samples were heated at 10 °C/min from 100 to 1000 °C under a 10:1 N_2 - O_2 atmosphere. All gases used were of high purity (> 5.8) and supplied by Carbagas and Linde. Thermograms were processed using a kinetic inversion code developed by Hemingway et al. (10), adapted for the inhouse analytical setup. By converting the thermograms into kinetograms, the inversion provides apparent activation energies (aEa), assuming a constant prefactor in the Arrhenius model. We found that a prefactor of $10^{8.5}$ which reliably reproduces the known activation energy of calcite (170-190 kJ/mol) and graphite (180-190 kJ/mol) using a Model-Free Kinetics (MFK) approach. All kinetograms presented here were obtained with a smoothing factor of 4 (SI Fig. 1), which optimally balances spectral detail and smoothness.

TOC data have been obtained using Rock-Eval pyrolysis at UNIL. The TOC concentrations were obtained on the same sample material that has been used for the dO_2 analyses (and hence for establishing the TOC vs. dO_2 correlation). The principle of Rock-Eval analysis is explained in details elsewhere (11, 12). In brief, two heating ramps are determined, the first under N_2 atmosphere and the second under air. The gas

produced during the first ramp (200 to 650 °C) is analyzed for hydrocarbons by a flame ionization detector (FID) and for CO₂ (and CO) using an infrared (IR) detector. The residue of the first pyrolysis step is introduced in a combustion chamber, and the evolved gas is analyzed for CO₂ (and CO) (11, 12). The total organic carbon (TOC) parameter is obtained following a standard procedure as revised by (13). In the absence of carbonates, the organic C content corresponds to the sum of all the carbon moieties (TOC and MINC, which represent the organic and mineral C content, respectively) released during pyrolysis and oxidation. Ultimately, a correcting factor of 1.166256 is applied to biopolymer-rich samples such as forest soil litter or peat material. This correction to the ‘hydrocarbon yield factor’ (14) ensures compatibility between Rock-Eval TOC- (TOC-RE) and EA TOC- (TOC-EA) derived values (13). This also ensures consistency with the basin wide TOC data published in (9, 15) that are also used here for the basin wide GIS O₂ mapping.

Sedimentary material thermal characteristics obtained by TGA-microGC. Probability density distribution, $p(o,E)$, based on mass loss characteristics of the sample in oxidizing atmosphere.



Thermogravimetric analysis (TGA) identified two organic matter pools in the IMB100 core (Fig. 1A,B): a labile fraction (decomposing at 370°C, activation energy E = 90 kJ/mol), comprising fresh plant litter and carbohydrates, and a refractory fraction (450°C, E = 110 kJ/mol), including lignocellulosic and aromatic compounds. The labile-to-refractory (L/R) ratio declined from 0.3 in surface peats to 0.2 at depth, reflecting selective preservation. During the Ghost Interval, both fractions showed a 5 kJ/mol E increase, indicating drier conditions favored recalcitrant compounds.

S5. Supplemental Table 1

Samples	Depth (cm)	Depth_mean (cm)	Thickness (cm)	Age med. (yr)	TOC (raw %)	TOC (corr %)	Peak_1_Ea (kJ/mol)	peak_1_area AU	Peak_2_Ea (kJ/mol)	peak_2_area AU	P1/P2 ratio	dO ₂ (μmol/g)	dO ₂ * (μmol/g TOC)	Stoichiometric ratio (O ₂ :C mol/mol)	Time interval
IMB1.002	16-18	17	19.0	215	39.1	47.4	94.2	0.2	116.8	0.8	0.3	44021	92965	1.12	Late Holocene
IMB1.004	20-22	21	4.0	320	44.0	52.8	95.7	0.2	117.8	0.8	0.3	48271	91358	1.10	Late Holocene
IMB1.006	24-26	25	4.0	442	42.0	50.8	95.7	0.2	117.3	0.8	0.3	46832	92242	1.11	Late Holocene
IMB1.008	28-30	29	4.0	570	43.3	52.0	92.7	0.2	114.3	0.8	0.3	48998	94145	1.13	Late Holocene
IMB1.010	32-34	33	5.0	703	44.5	53.9	92.7	0.2	114.3	0.8	0.2	48114	89325	1.07	Late Holocene
IMB1.013	38-40	39	6.0	905	44.0	53.7	93.7	0.2	115.3	0.8	0.3	46111	85858	1.03	Late Holocene
IMB1.016	44-46	45	6.0	1007	41.9	50.9	95.2	0.2	116.8	0.8	0.3	46875	92121	1.11	Late Holocene
IMB1.019	50-52	51	6.0	1122	41.6	50.2	95.7	0.2	117.8	0.8	0.3	47286	94138	1.13	Late Holocene
IMB1.022	56-58	57	6.0	1240	45.8	55.5	95.7	0.2	117.3	0.8	0.3	46038	82986	1.00	Late Holocene
IMB1.025	62-64	63	5.0	1360	40.8	49.7	94.7	0.2	116.8	0.8	0.3	na	na	na	Late Holocene
IMB1.027	66-68	67	4.0	1435	42.1	51.1	94.7	0.2	116.3	0.8	0.3	47627	93282	1.12	Late Holocene
IMB1.029	70-72	71	4.0	1509	42.9	52.6	94.7	0.2	115.8	0.8	0.3	47154	89713	1.08	Late Holocene
IMB1.031	74-76	75	5.0	1572	41.9	50.8	95.2	0.2	116.8	0.8	0.3	46538	91535	1.10	Late Holocene
IMB1.034	80-82	81	6.0	1676	42.3	51.8	95.2	0.2	116.8	0.8	0.3	46767	90310	1.08	Late Holocene
IMB1.037	86-88	87	6.0	1782	41.7	50.8	95.2	0.2	116.8	0.8	0.3	46616	91824	1.10	Late Holocene
IMB1.040	92-94	93	4.0	1882	43.9	53.0	95.7	0.2	117.8	0.8	0.3	na	na	na	Late Holocene
IMB1.041	94-96	95	3.0	1917	44.6	54.1	95.2	0.2	117.3	0.8	0.3	44097	81471	0.98	Late Holocene
IMB1.043	98-100	99	3.0	2056	43.0	52.3	95.2	0.2	116.8	0.8	0.3	48117	92053	1.10	Ghost Interval
IMB1.044	100-102	101	2.0	2257	43.3	52.6	95.7	0.2	117.3	0.8	0.3	48924	92966	1.12	Ghost Interval
IMB1.045	102-104	103	2.0	2590	46.1	56.0	95.2	0.2	116.8	0.8	0.3	48473	86558	1.04	Ghost Interval
IMB1.046	104-106	105	2.0	2905	44.5	53.6	95.7	0.2	117.3	0.8	0.3	49042	91425	1.10	Ghost Interval
IMB1.047	106-108	107	2.0	3042	47.4	57.0	96.2	0.2	117.3	0.8	0.3	50205	89082	1.06	Ghost Interval
IMB1.048	108-110	109	2.0	3178	47.1	56.6	96.2	0.2	117.8	0.8	0.3	51913	91729	1.10	Ghost Interval
IMB1.049	110-112	111	3.0	3578	46.8	56.6	96.2	0.2	117.8	0.8	0.3	51313	90693	1.09	Ghost Interval
IMB1.051	114-116	115	3.0	4890	48.0	58.2	95.2	0.2	116.8	0.8	0.2	51270	88158	1.06	Ghost Interval
IMB1.052	116-118	117	2.0	5621	47.2	57.1	96.2	0.2	117.8	0.8	0.2	50465	88372	1.06	Ghost Interval
IMB1.053	118-120	119	3.0	6359	45.6	55.0	95.2	0.2	116.8	0.8	0.3	48693	88589	1.06	Ghost Interval
IMB1.055	122-124	123	4.0	7496	44.1	53.5	94.7	0.2	115.8	0.8	0.2	46984	87839	1.05	Ghost Interval
IMB1.057	126-128	127	3.0	8251	44.2	53.9	94.7	0.2	116.3	0.8	0.2	46949	87113	1.05	Early Holocene
IMB1.058	128-130	129	2.0	8439	43.2	52.6	94.2	0.2	115.8	0.8	0.2	na	na	na	Early Holocene
IMB1.059	130-132	131	3.0	8561	42.9	52.3	94.7	0.2	115.8	0.8	0.3	46858	89527	1.07	Early Holocene
IMB1.061	134-136	135	5.0	8649	42.8	52.0	94.2	0.2	115.8	0.8	0.2	45887	88280	1.06	Early Holocene
IMB1.064	140-142	141	4.0	8789	43.6	53.1	94.2	0.2	115.8	0.8	0.2	44072	82965	1.00	Early Holocene
IMB1.065	142-144	143	3.0	8837	43.1	52.3	94.7	0.2	115.8	0.8	0.3	45369	86790	1.04	Early Holocene
IMB1.067	146-148	147	3.0	8934	44.7	54.5	94.2	0.2	115.3	0.8	0.2	45036	82599	0.99	Early Holocene
IMB1.068	148-150	149	4.0	8989	44.2	54.3	94.2	0.2	115.3	0.8	0.2	44784	82542	0.99	Early Holocene
IMB1.071	154-156	155	6.0	9242	45.0	54.4	92.7	0.2	113.3	0.8	0.2	41573	76387	0.92	Early Holocene
IMB1.074	160-162	161	6.0	9571	42.3	51.6	92.7	0.2	113.3	0.8	0.2	40488	78480	0.94	Early Holocene
IMB1.077	166-168	167	6.0	9903	42.3	51.4	99.7	0.1	119.8	0.9	0.2	na	na	na	Early Holocene
IMB1.080	172-174	173	6.0	10227	42.4	51.5	93.2	0.1	113.3	0.9	0.2	42722	82945	1.00	Early Holocene
IMB1.083	178-180	179	4.0	10542	44.1	53.7	94.2	0.1	113.3	0.9	0.2	na	na	na	Early Holocene
IMB1.084	180-182	181	3.0	10711	42.5	51.5	94.2	0.1	113.8	0.8	0.2	42605	82673	0.99	Early Holocene
IMB1.086	184-186	185	5.0	11132	37.6	45.6	94.7	0.1	113.3	0.9	0.2	na	na	na	Deglacial
IMB1.089	190-192	191	4.0	18424	36.7	44.6	94.2	0.1	112.8	0.8	0.2	40221	90209	1.08	Deglacial
IMB1.090	192-194	193	3.0	19695	34.3	41.7	96.7	0.1	114.8	0.8	0.2	na	na	na	Deglacial
IMB1.092	196-198	197	5.0	21118	32.7	39.7	95.2	0.1	112.8	0.8	0.2	na	na	na	Last Glacial Maximum
IMB1.095	202-204	203	4.0	21475	26.6	32.5	96.2	0.1	112.3	0.9	0.1	27634	85086	1.02	Last Glacial Maximum
IMB1.096	204-206	205	3.0	21561	24.8	30.2	95.7	0.1	112.3	0.8	0.2	22711	75221	0.90	Last Glacial Maximum
IMB1.098	208-210	209	5.0	21762	21.6	26.4	97.2	0.1	113.3	0.9	0.1	22711	86115	1.03	Last Glacial Maximum
IMB1.101	214-216	215	4.0	22626	23.0	28.1	96.2	0.1	112.8	0.8	0.2	na	na	na	Last Glacial Maximum
IMB1.102	216-218	217	3.0	22790	21.1	25.7	96.2	0.1	113.3	0.8	0.2	na	na	na	Last Glacial Maximum
IMB1.104	220-222	221	5.0	23240	22.1	26.9	96.7	0.1	112.8	0.9	0.1	22845	84786	1.02	Last Glacial Maximum
IMB1.107	226-228	227	4.0	24337	20.3	24.7	101.3	0.2	119.8	0.8	0.2	na	na	na	Last Glacial Maximum
IMB1.108	228-230	229	3.0	24675	20.6	25.1	97.2	0.1	113.8	0.9	0.1	21618	86296	1.04	Last Glacial Maximum
IMB1.110	232-234	233	2.0	25405	18.8	23.1	96.7	0.1	112.8	0.9	0.1	na	na	na	Last Glacial Maximum

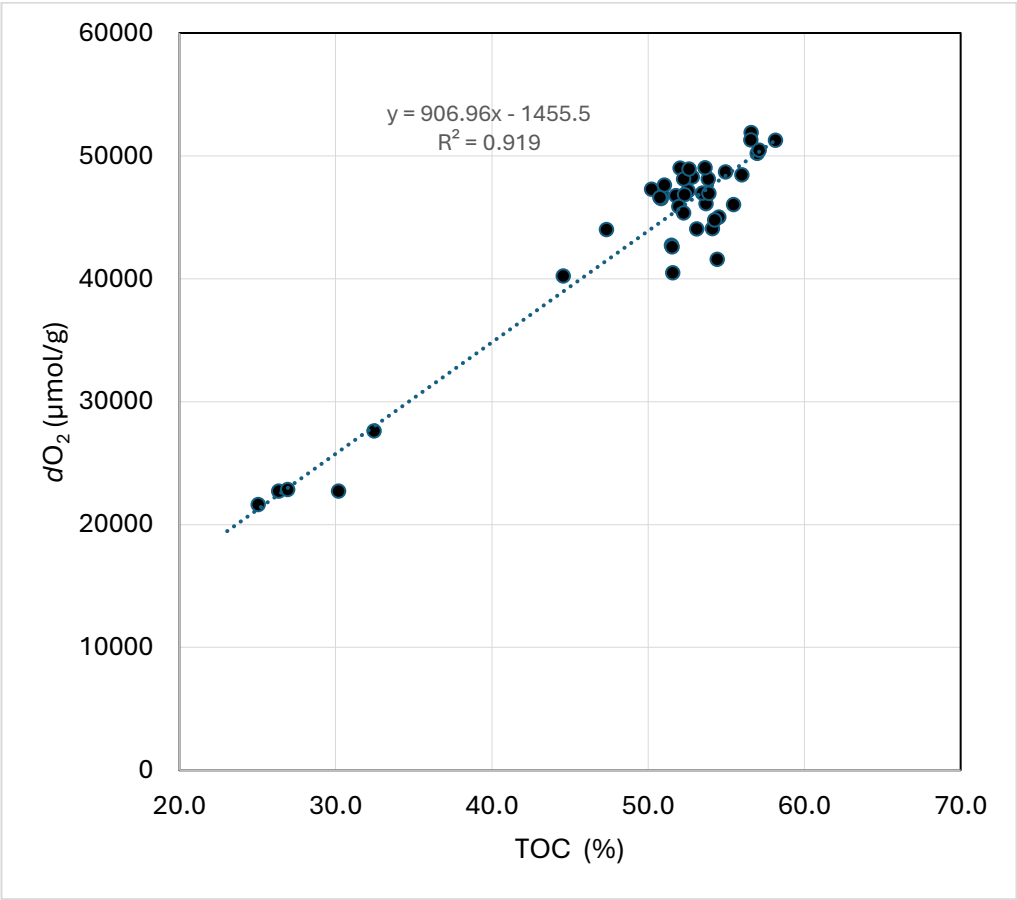
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101 S6. Geospatial analysis: TOC vs dO_2

102 Correlation between TOC (corrected) values for the IMB core (this work) and dO_2 values (SI Table 1) used

103 for O_2 production reconstructions.



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105 S7. Geospatial analysis

106 Correlation between TOC and peat thickness

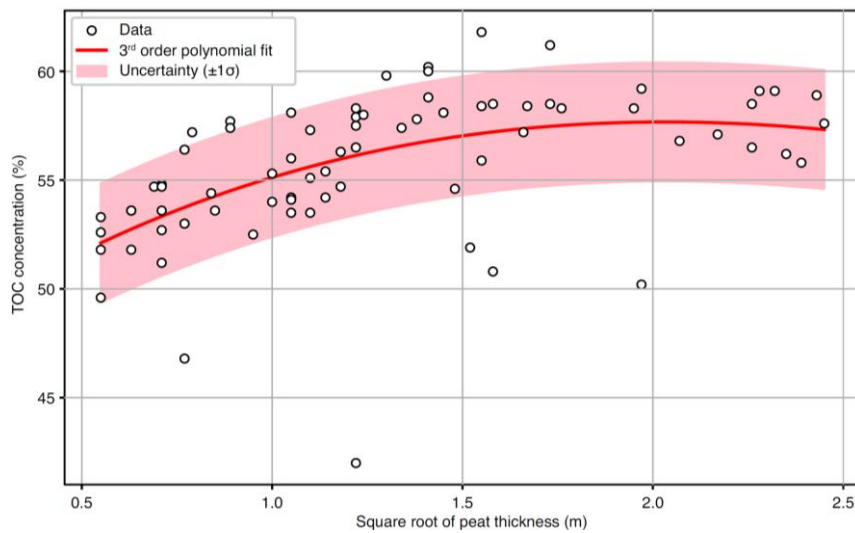
107 Using field data such as peat thickness (Fig 1B), sediment bulk density and organic carbon

108 concentrations, (15) have established a relationship between peat thickness and peat carbon density for

109 the central Congo Basin. In the same way, we used field data from (15) to construct a peat-thickness-total

110 organic carbon regression (SI Fig 2) to spatially model peat TOC content (Fig. 1C). We then applied the

regression between TOC values for the IMB100 core (see S6) and dO_2 values to the TOC content map to spatially model dO_2 at the basin scale (Fig. 3B).



S8. Flux uncertainties ($\pm 10 \text{ Pg O}_2$) was computed with a Monte Carlo simulation (2000 iterations) which incorporated variable peat density (normal distribution with mean 0.17 g/cm^3 , $1\sigma = 0.01 \text{ g/cm}^3$, providing values normally distributed in between minima at ca. 0.15 and maxima at ca. 0.19), dO_2 ($\pm 5\%$), and spatial variability (uniform sampling between 167600 and 185900 km^2).

B. Global Context

S9. Amazon O_2 production

We estimate the Amazon's net O₂ production at ~20–40 Tg O₂ yr⁻¹, derived from its net ecosystem production (NEP; 0.05–0.1 Pg C yr⁻¹, (16, 17) and stoichiometric conversion using an O₂:C molar ratio of ~1.1 mol/mol. This estimate converts NEP carbon fluxes to O₂ production, accounting for photosynthetic O₂ release relative to carbon fixation, with uncertainties reflecting variability in NEP and stoichiometric assumptions.

S10. Marine DOC O₂ sink

The highly recalcitrant dissolved organic carbon (DOC) reservoir in the global ocean, primarily the refractory DOC (DOCR) pool, contains approximately 640 Pg C, constituting ~97% of the total marine DOC pool of ~662 Pg C. This reservoir, concentrated in the deep ocean, exhibits remarkable stability with radiocarbon ages ranging from 4,000 to 6,000 years, reflecting its resistance to microbial degradation. Its vast size, comparable to atmospheric CO₂, and millennial persistence make it a critical component of long-term carbon sequestration, as quantified by (18) and (19) using global oceanographic data.

S11. Global peatland and geological O₂ flux estimates

Peatlands are critical carbon sinks, contributing to atmospheric O₂ through Holocene (11,700 years) carbon sequestration. Tropical peatlands (380,000–657,000 km²) store 110–140 Pg C, including 81–92 Pg from Southeast Asian and Peruvian Amazon peatlands (3, 20) and 30 Pg from the Congo Basin (3, 21). Boreal peatlands (3,100,000–3,900,000 km²) store 265–565 Pg C, with ~300 Pg accumulated during the Holocene (22, 23). Global peat carbon stocks are estimated at 598–618 Pg (best estimate: 610 Pg), excluding Congo (20). O₂ fluxes are calculated assuming an O₂:C stoichiometry of 0.9–1.15 mol/mol.

For example, Southeast Asian peatlands (67.5 Pg C, mid-range) sequester 5.769×10^{12} g C yr⁻¹ (6.75×10^{16} g ÷ 11,700 yr), equivalent to 4.808×10^{11} mol C yr⁻¹ (÷ 12 g/mol). With a 1.025 O₂/C molar ratio, this produces 4.928×10^{11} mol O₂ yr⁻¹, or 15.8 Tg O₂ yr⁻¹ (× 32 g/mol), yielding 184.5 Pg O₂ over 11,700 years.

Table S5 summarizes carbon stocks, accumulation rates, and O₂ fluxes, with a global O₂ contribution of ~1161.6 Pg.

Global estimated oxygen production of 99.3 [84.1-115.4] represents about 30 % of the global marine O₂ net production flux, estimated at ~336 Tg O₂ yr⁻¹ [126 Tg C yr⁻¹] by (24).

S12. Oxidative Weathering O₂ Sinks

- OC_{petro} weathering

Hilton and West (25) estimate modern OC_{petro} weathering at ~70 Tg C/yr (40–100 Tg C/yr), releasing CO₂ and consuming O₂ via equation 1, main text. With a 1:1 O₂:C mol/mol ratio (2.6667 g O₂/g C):

O₂ consumption: 70 × 2.6667 = 187 Tg O₂ yr⁻¹ (107–267 Tg O₂ yr⁻¹).

- Sulfide (FeS₂) Weathering O₂ Sinks

Burke et al. (2018) estimate modern sulfide weathering at 1.3 ± 0.2 Tmol S/yr (41.6 Tg S/yr), via FeS₂ + 15/4 O₂ + 7/2 H₂O = FeOH₃ + 2 H₂SO₄

O₂ consumption: 41.6 × 1.875 = 78 Tg O₂ yr⁻¹ (66–90 Tg O₂ yr⁻¹).

Process	C/S Flux (Tg yr ⁻¹)	O ₂ :Substrate Ratio	O ₂ Consumption (Tg yr ⁻¹)	Source
Petrogenic OC	70 [40–100] C	2.67 (g O ₂ /g C)	187 [106–267]	Hilton & West (2020)
Sulfide (FeS ₂)	42 [33–50] S	1.875 (g O ₂ /g S)	78 [66–90]	Burke et al. (2018)

Notes: Values are midpoints with ranges in parentheses. O₂ fluxes assume an O₂/C stoichiometry of 0.9–1.15 mol/mol. Global Total sums Total Tropical and Boreal contributions. Total tropical peatlands includes additional, smaller peatlands areas not reported in this table, but presented a thoroughly reviewed in (Page, 2011 #6167). OC = Organic carbon. Ranges reflect 1σ uncertainties. Stoichiometric ratios account for complete oxidation (OC + O₂ → CO₂; FeS₂ + 3/4 O₂ + 3/2 H₂O → FeO(OH) + 2H⁺ + 2SO₄²⁻ (1.875:1 mass O₂:S)). 1 mol FeS₂ (120g) consumes 1.875 mol O₂ (60g). Thus 41.6 Tg S/yr × (32g S/mol)/(120g FeS₂/mol) × 1.875 × 32g O₂/mol = 78 Tg O₂/yr

S13. Synthesis of O₂ and CO₂ Deficits During the Ghost Interval

The Ghost Interval (GI; 7.5–2.3 kyr BP) caused an 80% decline in the Congo Basin's peatland O₂ flux (from 25–30 Tg O₂ yr⁻¹ to 5–6 Tg O₂ yr⁻¹), which impacted both O₂ and CO₂ budgets:

1. O₂ Deficit

- Total O₂ production during GI: 30–36 Pg O₂ (6 kyr × 5–6 Tg O₂ yr⁻¹).

- Expected O₂ without GI: 150–180 Pg O₂ (assuming sustained pre-GI rates).
- O₂ deficit: 120–144 Pg O₂ (missed production due to aridity).
 - Equivalent to ~50% of potential Holocene output (203–227 Pg O₂ if GI had not occurred).

2. CO₂ Sequestration Deficit

Using the measured O₂/C stoichiometry (1.025 mol O₂/mol C):

- C loss: 120–144 Pg O₂ deficit × 0.366 Tg C/Tg O₂ = 44–53 Pg C.
- CO₂ equivalent: 44–53 Pg C × 3.67 = 161–194 Pg CO₂.

S14. Carbon Cycle Perturbation Thresholds

The ghost interval’s carbon perturbation, which we estimated at $\Delta m_{GI} = 44\text{--}53$ Pg C over $\tau_{env} = 6$ kyr, is evaluated against Rothman’s (2017) critical size ($M_c \approx 0.0082 \pm 0.0041$, equivalent to $\Delta m_c \approx 310 \pm 155$ Pg C) and critical rate ($r_c \approx 0.0016 \pm 0.0005$ kyr⁻¹) thresholds for catastrophic carbon cycle destabilization on long timescales as observed in four of the “Big Five” mass extinctions (e.g., end-Permian, $M = 0.243$, $\tau_{env} = 60$ kyr; Fig. 2B, p. 4). The critical rate r_c indicates that carbon additions exceeding 0.16% of the ocean’s carbon mass per thousand years (~61 Pg C/kyr) on short timescales risk catastrophic outcomes.

The normalized mass of C for the Ghost Interval in Congo, $M_{GI} = \Delta m_{GI} / m^*$ ($m^* \approx 38,000$ Pg C, the modern ocean inorganic carbon), ranges from 0.00116 (44 Pg C) to 0.00140 (53 Pg C), well below M_c , indicating that the perturbation’s magnitude was indeed insufficient to trigger global catastrophic change. However, the normalized perturbation rate, M_{GI} / τ_{env} , ranges from 0.193 to 0.233 kyr⁻¹ (ie. 7.33–8.83 Pg C/kyr, the raw rate of carbon addition), exceeding r_c by two orders of magnitude, even surpassing the normalized perturbation rate of the Big Five’s rates (0.0023–0.0048 kyr⁻¹, (26)).

The ghost interval’s rate is extremely rapid relative to the ocean’s carbon mass, suggesting potential for destabilization if sustained for ~35–42 kyr (approaching $M_c \sim 310$ Pg C imbalance). However, the Ghost Interval’s brief duration (6 kyr) and limited mass, coupled with possible compensation from other global regions, averted catastrophe. In contrast, the end-Permian (~252 Ma, $M = 0.24$, $\tau = 60$ kyr, $r = 0.00405$

kyr⁻¹) and end-Triassic (~201 Ma, $M = 0.095$, $\Delta m = 3600$ Pg C, $\tau = 20$ kyr, $r = 0.00475$ kyr⁻¹) perturbations rates (26) exceeded the critical rate and, with durations of 60 kyr and 20 kyr, amassed carbon imbalances ($\Delta m_{\text{Permian}} = 9234$ Pg C, $\Delta m_{\text{Triassic}} = 3610$ Pg) that crossed M_c , triggering catastrophic carbon cycle shifts.

S15. Future O₂ flux losses (33–80%, ~33–80 Tg O₂ yr⁻¹ globally) were estimated by scaling historical flux reductions (80% during the mid-Holocene Ghost Interval, ~7.5–2.3 ka BP, linked to increased aridity in the central Congo Basin (9)). (9) provided quantitative estimates of precipitation with a reduction of annual precipitation of ~35–50 % (if precipitation seasonality is kept invariant) between 5.5 and 2.3 kyr BP, which led to the formation of the Ghost Interval. Other paleoclimate proxy data from East, West, and Central Africa, indicating comparable drying trends during the Ghost Interval. In East Africa, lake sediment records from Lake Chala suggest a 20–40% reduction in annual rainfall relative to early Holocene levels, driven by weakened monsoon intensity (27). In West Africa, pollen and lake-level reconstructions from Lake Bosumtwi indicate a 15–35% rainfall decline, associated with a southward shift of the Intertropical Convergence Zone (ITCZ) (28). For Central Africa, marine sediment cores off the Congo River mouth show a 20–45% reduction in terrestrial runoff, reflecting decreased precipitation in the Congo Basin (29). These regional estimates (15–45% rainfall reduction) align with the Congo Basin’s aridification during the Ghost Interval (9), halving carbon burial rates and causing the observed 80% O₂ flux decline (this work).

These historical drying trends are consistent with CMIP6-projected rainfall declines for Central Africa (20–40% by 2100 under SSP5-8.5 (30)), which predict increased aridity due to weakened monsoon circulation and ITCZ contraction. Assuming a linear relationship between rainfall reduction and O₂ flux loss, a 20–40% rainfall decline by 2100 could reduce global peatland O₂ fluxes by 28–93% (~27.8–92.3 Tg O₂ yr⁻¹), as reduced inundation accelerates organic matter oxidation. These projections are conservative, as nonlinear feedbacks (e.g., vegetation shifts) may amplify flux losses.

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