

Supplementary

A new assessment of global and regional budgets, fluxes and lifetimes of atmospheric reactive N and S gases and aerosols

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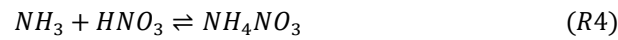
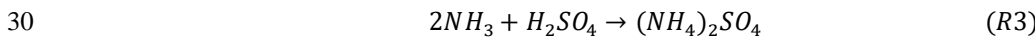
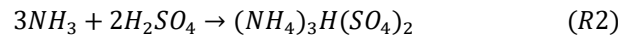
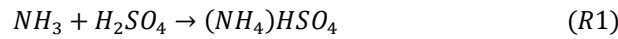
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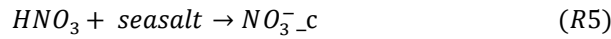
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S1. Model aerosol scheme

20 The model aerosol scheme is the Equilibrium Simplified Aerosol Model V4 (EQSAM4clim), which efficiently parameterises the aerosol water uptake and full gas-liquid-solid partitioning of mixtures of semi-volatile and non-volatile compounds. For environments with low NH₃ concentrations, sulfuric acid occurs in aerosol phase as H₂SO₄. When NH₃ is present, this dissolves irreversibly in the sulfate aerosol either totally, if limited by NH₃, or until titration of acidity (R1, R2, and R3), if limited by H₂SO₄. When H₂SO₄ is fully neutralised, residual NH₃ equilibrates with HNO₃ to form NH₄NO₃ aerosol
 25 (R4). Concentrations of NH₃ that exceed this full titration remain in the gas phase. Ammonium sulfates are considered thermally stable but the volatilisation of NH₃ and HNO₃ from NH₄NO₃ is a function of temperature and relative humidity. Both ammonium sulfate (from R1-R3) and ammonium nitrate (from R4) are classified as fine mode aerosol.



The HNO₃ also reacts with coarse mode sea salt and dust particles to produce coarse nitrate (R5 and R6).



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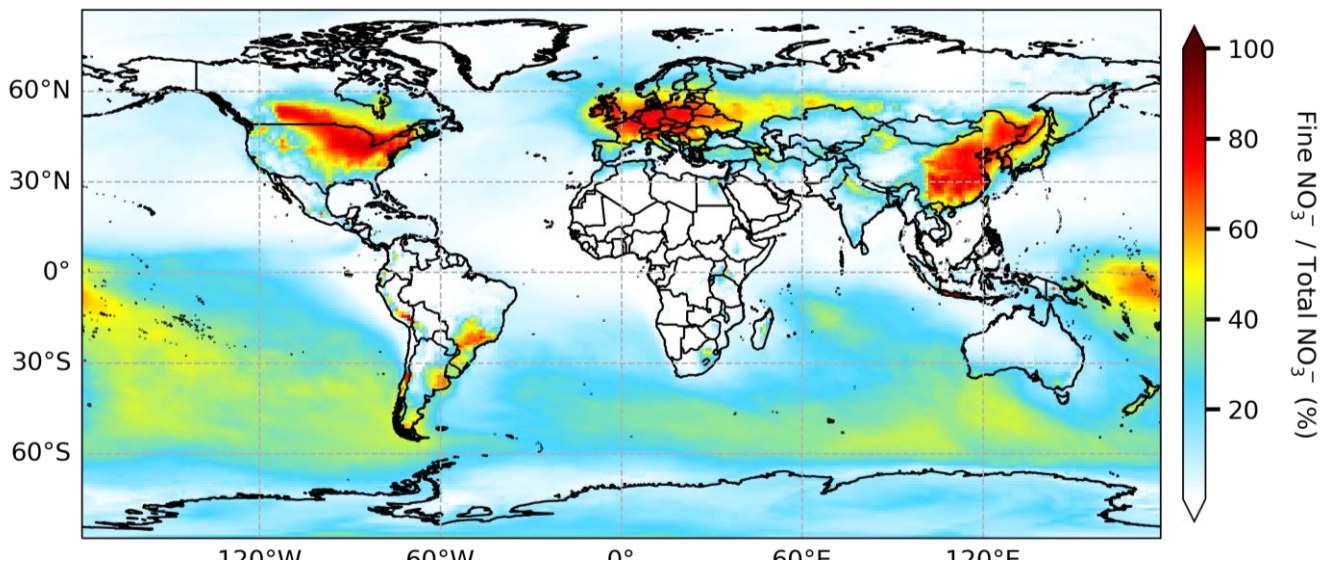
Table S1: Definition of RDN, OXN and OXS chemical groups.

Groups	Chemical species
RDN	NH ₃ , aerosol-phase NH ₄ ⁺
	HNO ₃
	NO _x
OXN	aerosol-phase NO ₃ ⁻
	NO, NO ₂
	NO ₃ ⁻ _f, NO ₃ ⁻ _c
	N ₂ O ₅ , HONO
	Rest OXN
	Other OXN species
	HO ₂ NO ₂ , SC ₄ H ₉ NO ₃ , PAN, MPAN, ISON, NALD
OXS	SO ₂ , aerosol-phase SO ₄ ²⁻ (latter includes H ₂ SO ₄ , NH ₄ HSO ₄ , (NH ₄) ₃ H(SO ₄) ₂ and (NH ₄) ₂ SO ₄)

40 In the chemical group of ‘Other OXN species’, PAN is peroxyacetyl nitrate, MPAN is peroxyacetyl nitrate, ISON is a lumped species representing first-generation nitrates from isoprene degradation, and NALD is used as a representative species for second- and subsequent-generation nitrates from isoprene degradation. The suffices ‘_f’ and ‘_c’ denote fine and coarse particle size fractions, respectively.

45 **Table S2: The allocation of IPCC reference regions (Iturbide et al., 2020) to the 10 world regions used in this study.**

World regions in this paper	IPCC region numbers (0-57)
Southeast Asia	38
East Asia	35
South Asia	37
Rest of Asia	28, 29, 30, 31, 32, 33, 34, 36
Euro_Medi	16, 17, 18, 19
North America	3, 4, 5, 6, 7, 8
South America	9, 10, 11, 12, 13, 14, 15
Africa	20, 21, 22, 23, 24, 25, 26, 27
Oceania	39, 40, 41, 42, 43
Rest of world	0, 1, 2, 44, 45, 46, 47, 48, 49, 50, 51, 52, 53, 54, 55, 56, 57



50 **Figure S1: The % contribution of fine NO_3^- to total NO_3^- (fine + coarse).**

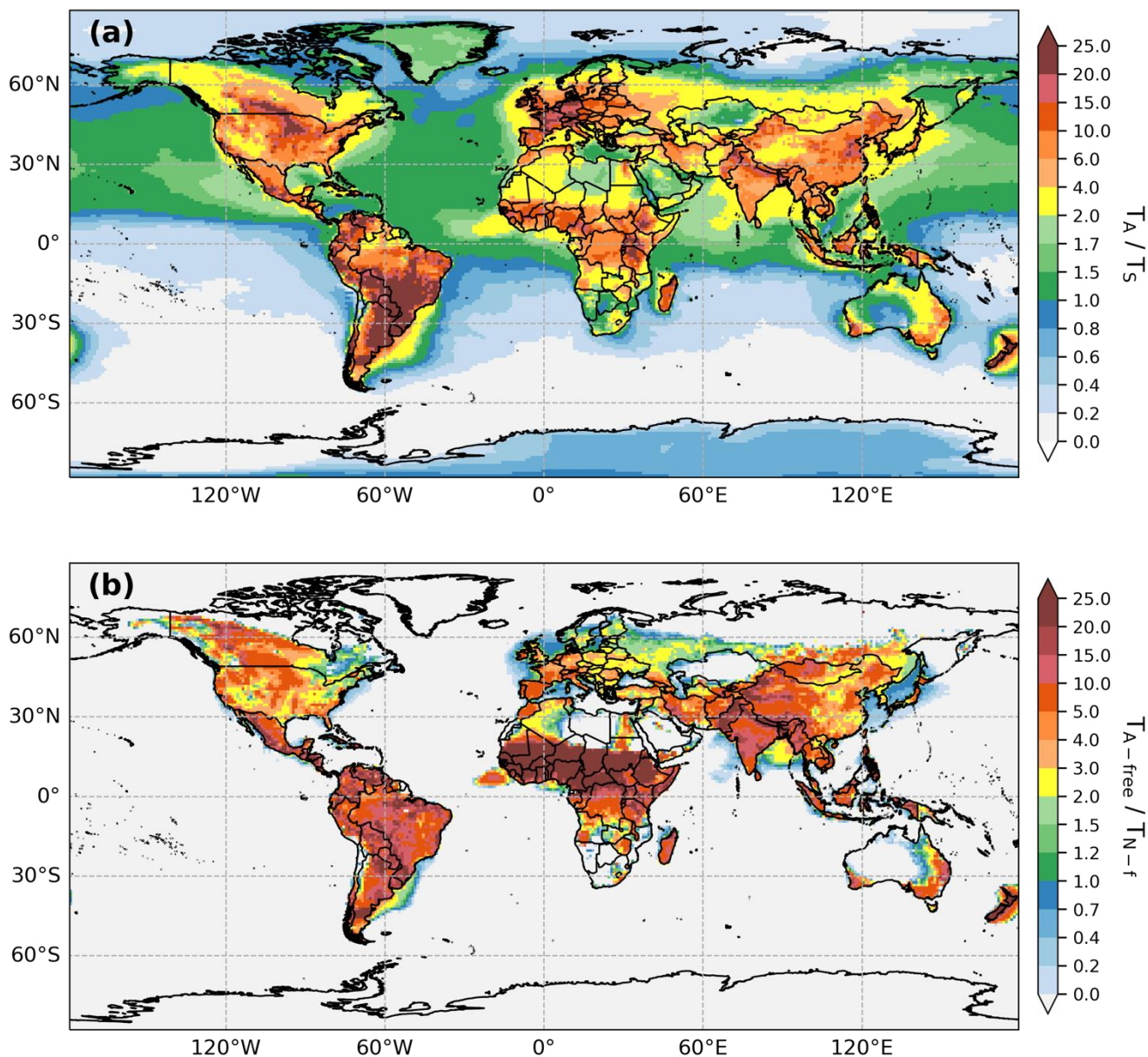


Figure S2: Maps of (a) the ratio of annual surface total ammonia T_A ($= [\text{NH}_3] + [\text{NH}_4^+]$) to total sulfate T_S ($= [\text{SO}_4^{2-}]$) molar concentrations; (b) the ratio of free ammonia T_{A-free} ($= T_A - (2 \times T_S)$) to total nitrate (fine) T_{N-f} ($= [\text{HNO}_3] + [\text{NO}_3^-_{f}]$). These maps provide the quantitative species ratios that underpin the categorisation into the four chemical domains shown in Fig. 3 of the main paper. In panel (a), wherever $\frac{T_A}{T_S} > 2$ all sulfate is fully neutralised to $(\text{NH}_4)_2(\text{SO}_4)_2$. Locations where $1 < \frac{T_A}{T_S} < 2$ are characterised as ‘ SO_4^{2-} rich’, and locations where $\frac{T_A}{T_S} < 1$ are characterised as ‘ SO_4^{2-} very rich’. In panel (b) there are coloured areas only where $\frac{T_A}{T_S} > 2$ in panel (a), i.e. the grey colour in panel (b) shows locations where the sulfate is not fully neutralised with ammonia and no free ammonia exists. Locations in panel (b) where $0 < \frac{T_{A-free}}{T_{N-f}} < 1$ indicate where NH_3 is the limiting factor for NH_4NO_3 formation and are characterised as ‘ NO_3^- rich.’ Locations in panel (b) where $\frac{T_{A-free}}{T_{N-f}} > 1$ are characterised as ‘ NH_3 very rich’, and are where it is availability of nitrate rather than NH_3 that limits NH_4NO_3 formation. See main paper for further description.

65 **Table S3: Maximum, median, and mean $\frac{T_{A-free}}{T_{N-f}}$ ratios in the 10 world regions defined in Figure 1 of the main paper. The max and median values refer to an individual model grid within that region, while the mean ratio is calculated using the full-region area-weighted T_{A-free} divided by the full-region area-weighted T_{N-f} .**

	Maximum	Median	Mean
Southeast Asia	56	0.0	1.9
East Asia	17	2.4	3.0
South Asia	87	9.9	9.6
Rest of Asia	93	-2.6	4.1
Euro_Medi	18	1.3	2.5
North America	68	2.0	3.1
South America	57	7.4	8.6
Africa	730	2.8	7.2
Oceania	138	-1.5	0.7
Rest of world	31	-7.7	-2.8

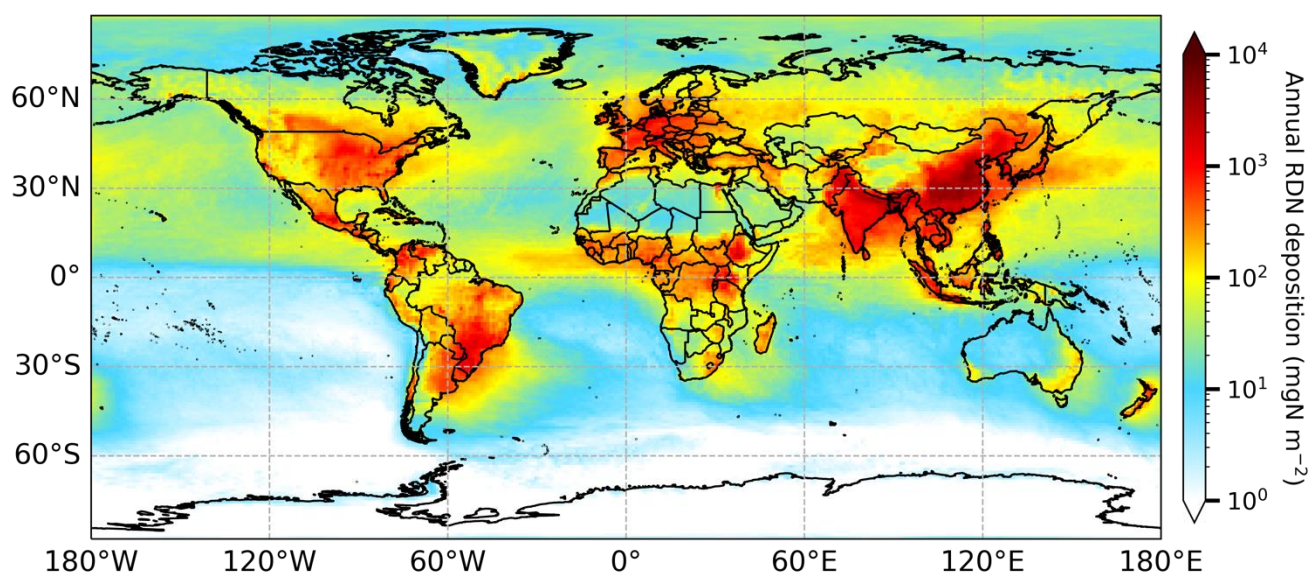


Figure S3: Annual total deposition (wet + dry) of RDN for 2015. Note the logarithmic scale.

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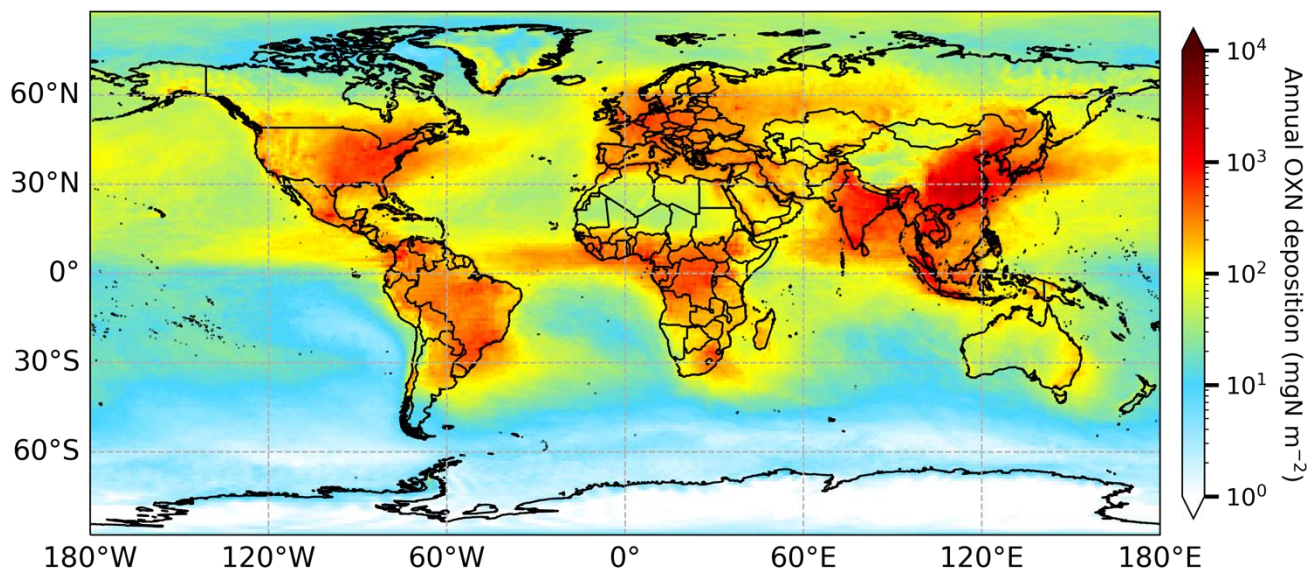


Figure S4: Annual total deposition (wet + dry) of OXN for 2015. Note the logarithmic scale.

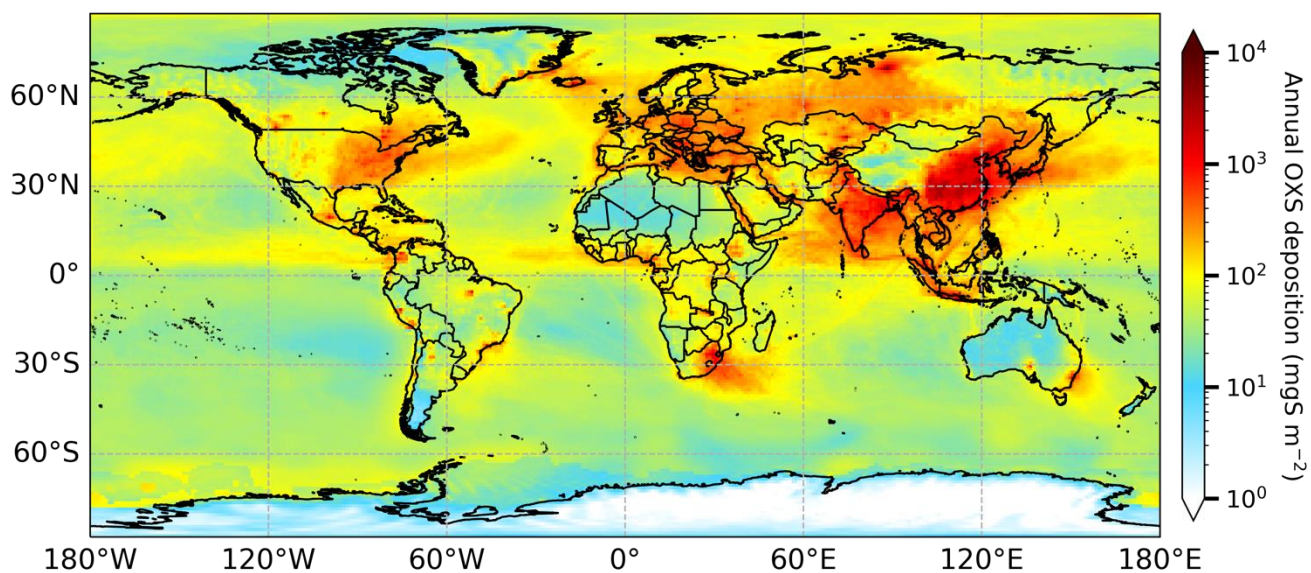


Figure S5: Annual total deposition (wet + dry) of OXS for 2015. Note the logarithmic scale.

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Table S4: Annual global deposition of gaseous and aerosol-phase RDN, OXN, and OXS species. DDEP represents dry deposition, and WDEP represents wet deposition. The ‘Amount’ represents the absolute value of deposition of one component, while the ‘Proportion’ represents the contribution of that component to global total RDN/OXN/OXS deposition.

(TgN/TgS yr ⁻¹)	Gaseous DDEP		Aerosol DDEP		Gaseous WDEP		Aerosol WDEP	
	Amount	Proportion	Amount	Proportion	Amount	Proportion	Amount	Proportion
RDN	22.4	40%	5.02	9%	12.2	22%	15.8	28%
OXN	16.0	28%	8.71	15%	10.5	18%	22.7	39%
OXS	17.9	35%	5.24	10%	5.44	11%	22.0	44%

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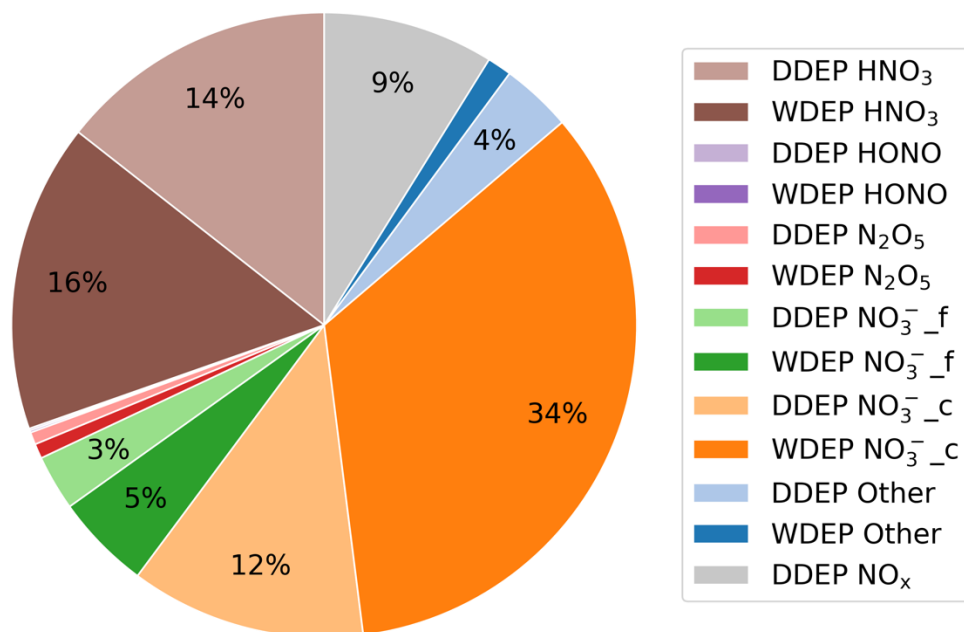
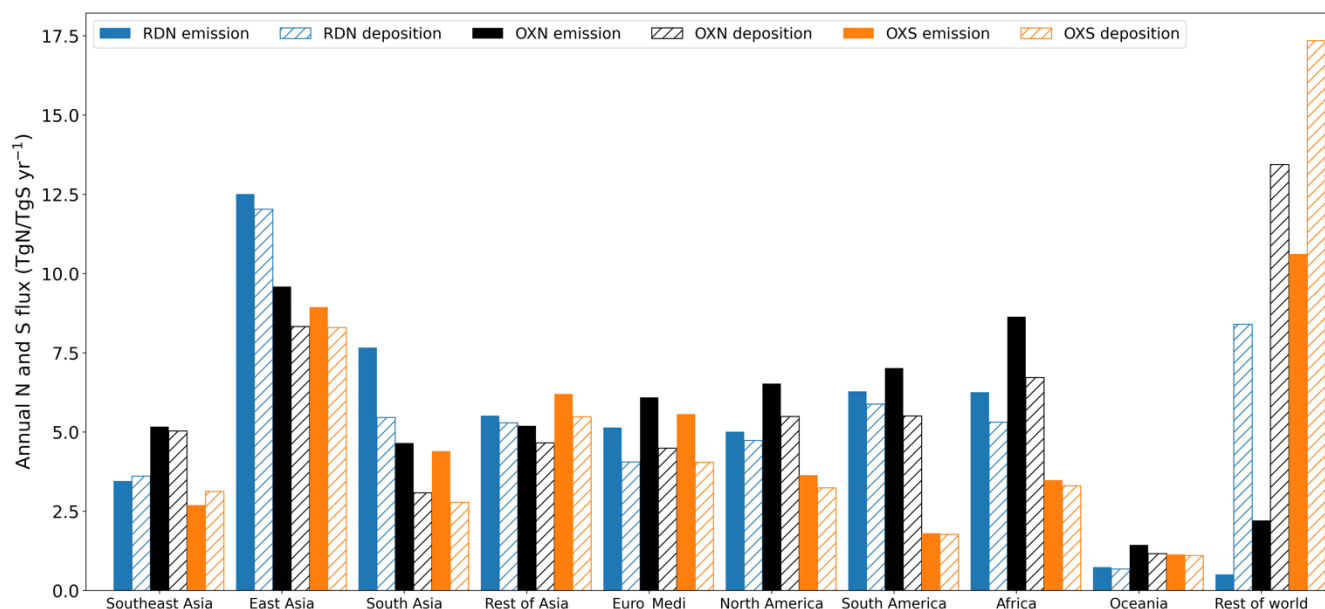
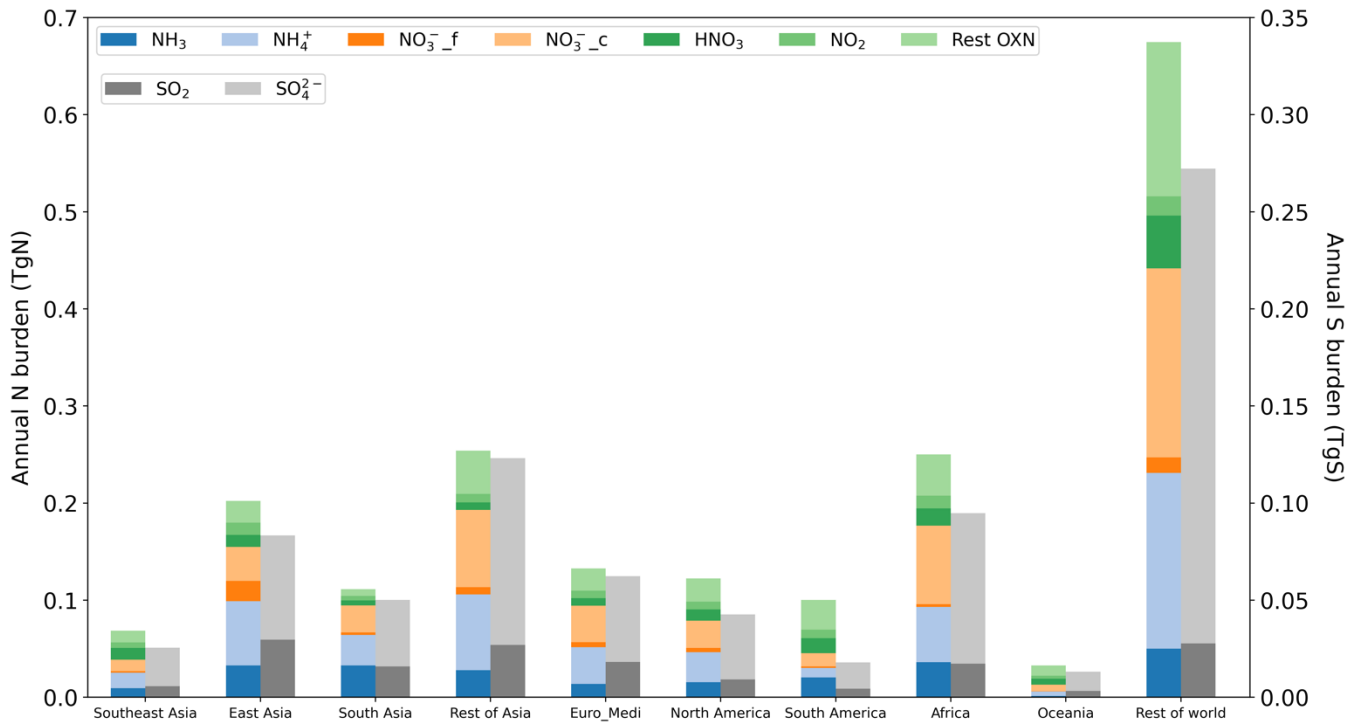


Figure S6: Relative contributions to annual global OXN deposition. Labels for individual contributions <2% are omitted to aid clarity. The darker shade colours collectively indicate wet deposition (WDEP), whilst the lighter shade colours collectively indicate dry deposition (DDEP). The ‘Other’ OXN species is predominantly PAN (full compositional details in Table S1).

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90 Figure S7: Emission and deposition fluxes of RDN (TgN yr⁻¹), OXN (TgN yr⁻¹), and OXS (TgS yr⁻¹) for the 10 world regions defined in Fig. 1. The constituents of RDN, OXN and OXS are listed in Table S1.



95 **Figure S8: Compositions of the tropospheric burdens in 2015 of total N (RDN + OXN) and total S (OXS) for the 10 world regions defined in Fig. 1. ‘Rest OXN’ includes HONO, N₂O₅ and other OXN species (full listing given in Table S1). Note the different scales for the N and S burdens.**

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Table S5: Regional atmospheric lifetimes for RDN, OXN and OXS calculated using either total emission or deposition fluxes. Q is the burden, P is the production pathway (i.e., emissions), R is the removal pathway (i.e., depositions), and τ is atmospheric lifetime. These regional lifetimes are not fully accurate because they do not take into account the net transport of RDN, OXN or OXS across the boundary of the region, but these are relatively small for all regions except Rest of world.

(days)	Southeast Asia	East Asia	South Asia	Rest of Asia	Euro_Medi	North America	South America	Africa	Oceania	Rest of world
RDN										
$\tau = \frac{Q}{P}$	2.7	2.9	3.1	7.0	3.7	3.4	1.8	5.4	2.9	164
$\tau = \frac{Q}{R}$	2.6	3.0	4.3	7.3	4.6	3.6	1.9	6.4	3.2	10
OXN										
$\tau = \frac{Q}{P}$	3.1	3.9	3.7	10	4.8	4.2	3.6	6.6	6.8	73
$\tau = \frac{Q}{R}$	3.1	4.5	5.6	12	6.6	5.0	4.6	8.5	8.4	12
OXS										
$\tau = \frac{Q}{P}$	3.4	3.4	4.2	7.2	4.1	4.3	3.6	9.9	4.2	9.4
$\tau = \frac{Q}{R}$	3.0	3.7	6.6	8.2	5.6	4.8	3.7	10	4.4	5.7

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