Pitch







Supersonic jet would fly from NYC to London in just 80 minutes

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What will it cost us?









EĜU













Thank you for your attention!

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Interactive component







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Non-CO₂ emissions of supersonic aircraft have multiple ways of affecting the Ozone layer and climate [1-7].





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scenario

Impact

To evaluate the atmospheric impacts of supersonic emissions, we need chemistry transport & chemistry climate models. These model chemistry and transport for hundreds of chemical species across the atmosphere.

We make use of the GEOS-Chem model for our simulations.

Modelling approach





Overview of processes chemistry transport models solve for thousands of atmospheric cells

Due to their complexity, these models need high-performance computer clusters to use.

Even then, a single multi-year model evaluation can take weeks to evaluate.

A small part of the Snellius supercomputer, which we use for our research





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We study the impact of supersonic emissions in two flight regions: the transatlantic corridor (TAC) and south arabian sea (SAS).



In these regions we introduce emissions representing 8 Tg of annual fuel burn of a hypothetical supersonic aircraft.

Emissions profile



With the GEOS-Chem Chemistry Transport Model we evaluate the impact of these emissions on a modern atmosphere over the course

of 10 years.

Model setup



We combine over 24 variations of these scenarios to calculate first- and second-order sensitivities of the global ozone response in these regions.

Parametric study



Two regions of anticipated use for supersonic flight are selected: the transatlantic corridor (TAC) and south arabian sea (SAS).



These regions are chosen as they are likely to be used for supersonic aviation, even if overland supersonic flight restrictions are considered [1,5]. Within these regions we introduce emissions representative of 8Tg of annual

supersonic fuel burn.

Emissions profile

These regions are located in different parts of the Brewer-Dobson circulation, affecting how emissions and impacts from these regions propagate. This is demonstrated below with a simplified lagrangian model using MERRA-2 meteorology:



Trajectories show average transport of emissions in june from the **SAS** and **TAC**. Vector field shows wind fields averaged over 3 years of data.







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*The emission of CO_2 is not incorporated in the GEOS-Chem model.

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Emissions are introduced as a box volume in a 2014 atmosphere. For anthropogenic emissions the CEDS v2 we use anthropogenic surface emissions inventory [8] and subsonic aviation emissions estimated from ADSB data [9]. Vemaps.com Change in global column ozone (%)from TAC emissions, evaluated by GEOS-Chem 2014-01-01 -0.4 -0.2 0.0 0.2 0.4 Δ%



Part of the Snellius supercomputer \sim



We use v13.3.1 of the GEOS-Chem chemistry transport model to evaluate the impact of these emissions over the course of 10 years. We use a global resolution of $4^{\circ} \times 5^{\circ}$ (lat,lon) with 72 altitude layers and 20 minute timesteps.

The model is ran using the Dutch national supercomputer Snellius, with support of the dutch national e-infrastructure.







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Results are grouped in several categories:



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Generally the introduction of the emissions leads to increases in global ozone across all scenarios with altitudes below 19.3 km. Across all 20.4 km scenarios ozone depletion outweighs lower-stratospheric production instead.

Average 2021-2024 change in global column ozone [DU]

	Base		$+NO_x$		$+SO_x$		$+H_2O$	
	TAC	SAS	TAC	SAS	TAC	SAS	TAC	SAS
16.2 km	0.3105	0.2870	0.4009	0.3724	0.2788	0.2628	0.3062	0.2871
18.3 km	0.1350	0.1025						
20.4 km	-0.3121	-0.7160	-0.4136	-0.9014	-0.3619	-0.7500	-0.3359	-0.7270
	$+\mathrm{HC}$	+CO	$+NO_x+SO_x$		$+SO_x+H_2O$		+H ₂ O+NO	
16.2 km	0.3118	0.3110						
20.4 km			-0.4576	-0.9532	-0.3850	-0.8073	-0.4353	-0.9331
							(preliminary)	

The SAS region is more sensitive to global ozone depletion. At 20.4 km we find over double the ozone loss compared to equivalent emissions in TAC.



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Emissions from the TAC region stay mostly in the northern hemisphere, also containing their impact. Supersonic emissions lead to **ozone increases** in the lower stratosphere, and **depletion** at higher altitudes. The latter ends up larger, decreasing global column ozone.

From the SAS region emissions are transported to the upper stratosphere more effectively. This increases ozone depletion by NO_x as well as mixing between hemispheres. Averaged globally, can lead to up to double the ozone loss.

Transport differences

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Spread of emitted black carbons

2014-01-01



Here we show the spread of emitted black carbon (soot) to highlight differences in transport. Notice that in the SAS region (lower) it takes around 2 years for the soot to saturate the stratosphere, a year faster than TAC emissions.

The faster and more effective transport to the upper stratosphere enhances ozone depletion to NO_x . Better hemispheric mixing also contributes to stronger global impacts.

First-order sensitivities









stratospheric transport.

2014-01-01



Emissions of NO_x , SO_x , and H_2O interact through heterogeneous chemistry. In this reaction NO_x is converted to HNO_3 , suppressing ozone loss. This interaction results in nonlinearities, which we quantify through cross-sensitivities.

Here we show global ozone loss for 3 scenarios. The bottom right panel shows changes in ozone impacts from interactions between NO_x and SO_x . The magnitude of this has been multiplied by 50 for visualisation.

$$\blacktriangleright NL = \Delta O_3(NO_x, SO_x) - (\Delta O_3(NO_x) + \Delta O_3(SO_x))$$

After some time you can see that the interactions between the NO_x and SO_x emissions "increase" ozone columns, ozone loss is reduced because more NO_x is converted to HNO_3 .



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Cross-sensitivities characterise how the response to one emission species changes with other emissions.



If H_2O and SO_x emissions both increase by 30%, their interactions increase O_3 depletion by -0.0716 DU (10%!) in the SAS region. In the TAC region this would decrease ozone loss by less than 1% instead!

Sensitivities of atmospheric ozone to supersonic emissions

In this figure we show the magnitude of the 2nd order sensitivities between $NO_{x_{y}} SO_{x'}$ and H_2O emissions.

In the TAC region interactions through chemistry **reduce ozone loss** by accelerated conversion of NO_x . For SAS emissions these interactions **accelerate ozone loss instead**, with a considerably larger effect.

The large differences in these interactions between locations may represent a considerable challenge for surrogate modelling! Results

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The region of supersonic emissions greatly affects their impact on global ozone levels. Over the south arabian sea we find over double to ozone depletion from supersonic cruise emissions compared to the transatlantic corridor, primarily due to NO_x emissions.

General Ozone response First-order sensitivities

We observe large differences in NO_x -SO_x-H₂O cross-sensitivities between the studied regions. Above the atlantic corridor cross-sensitivites dampen ozone depletion, the opposite occurs above the south-arabian sea.

Chemical cross-sensitivies

 NO_x - SO_x - H_2O may have a considerable effect on the impact of emissions in some regions, making it important to include them in surrogate modelling. Before this can be undertaken we first need to better understand geospatial dependencies of these interactions.

Read more about the MORE & LESS project

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