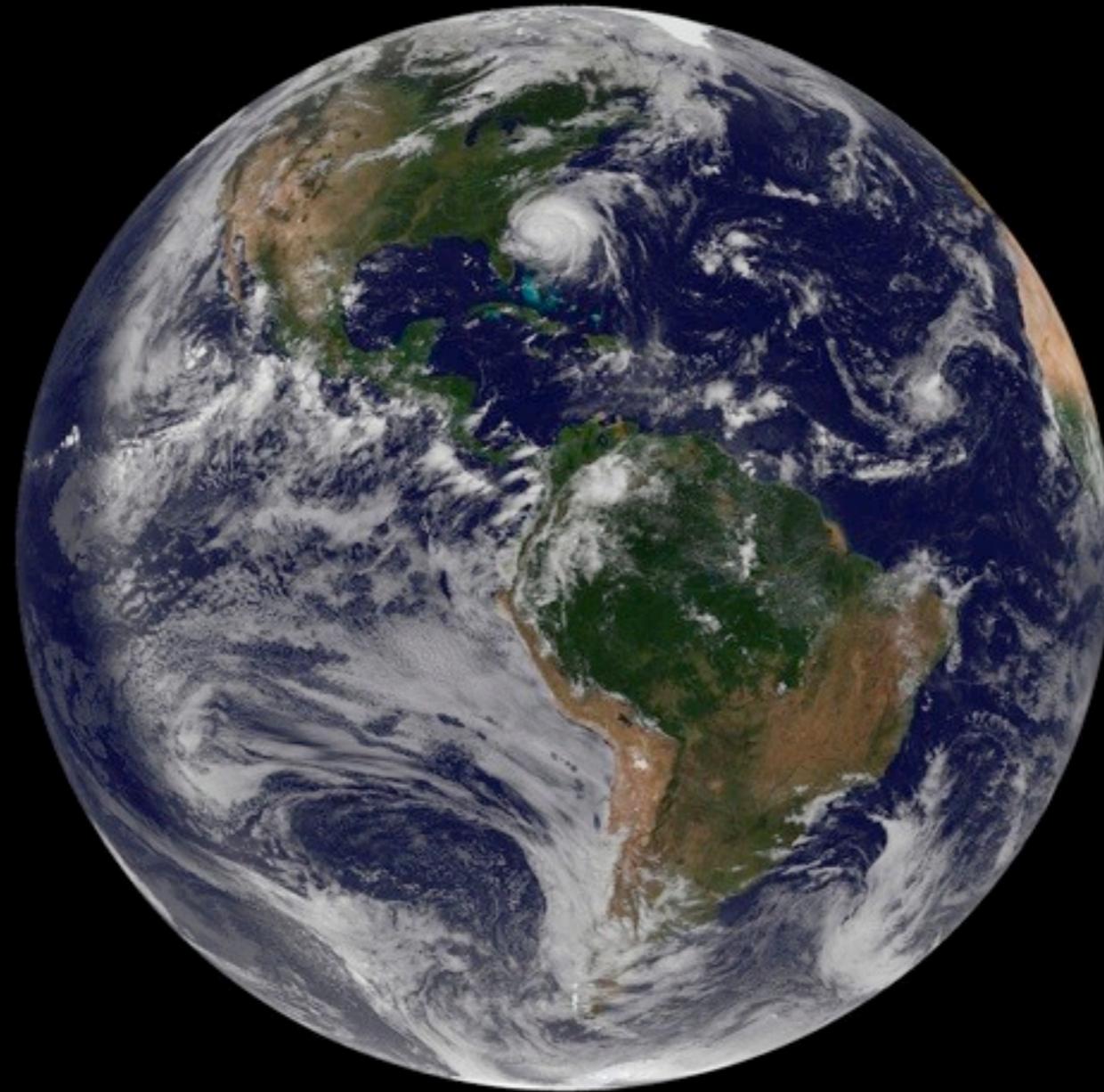
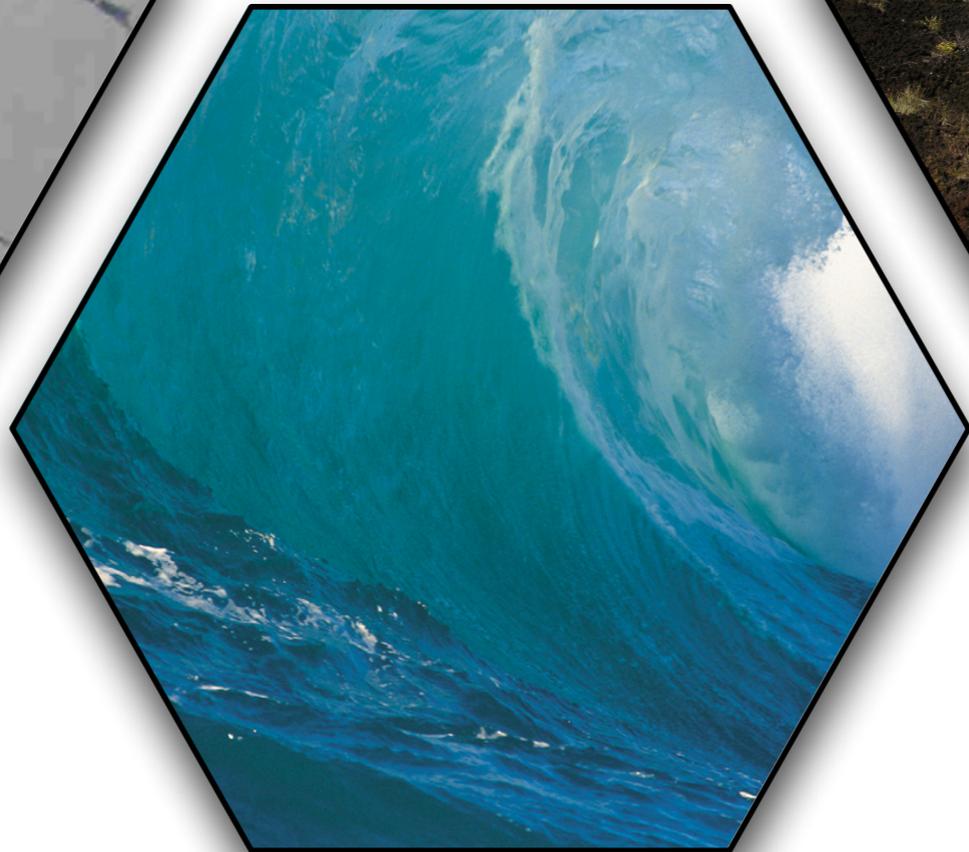
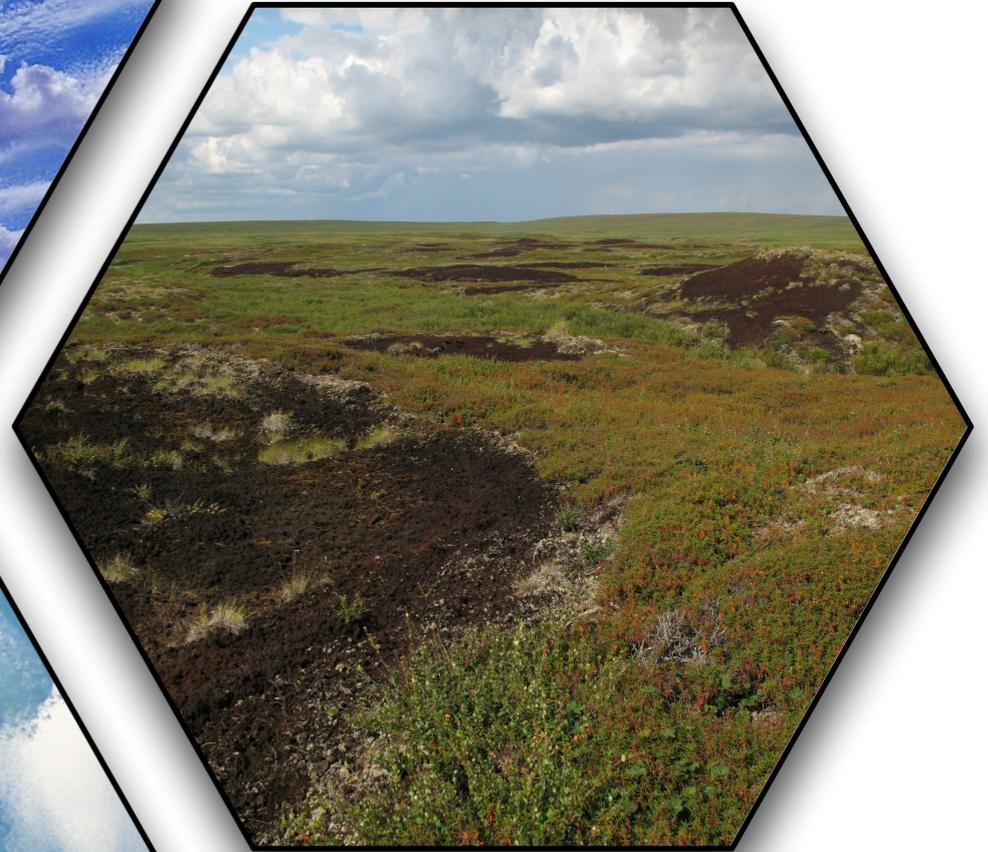
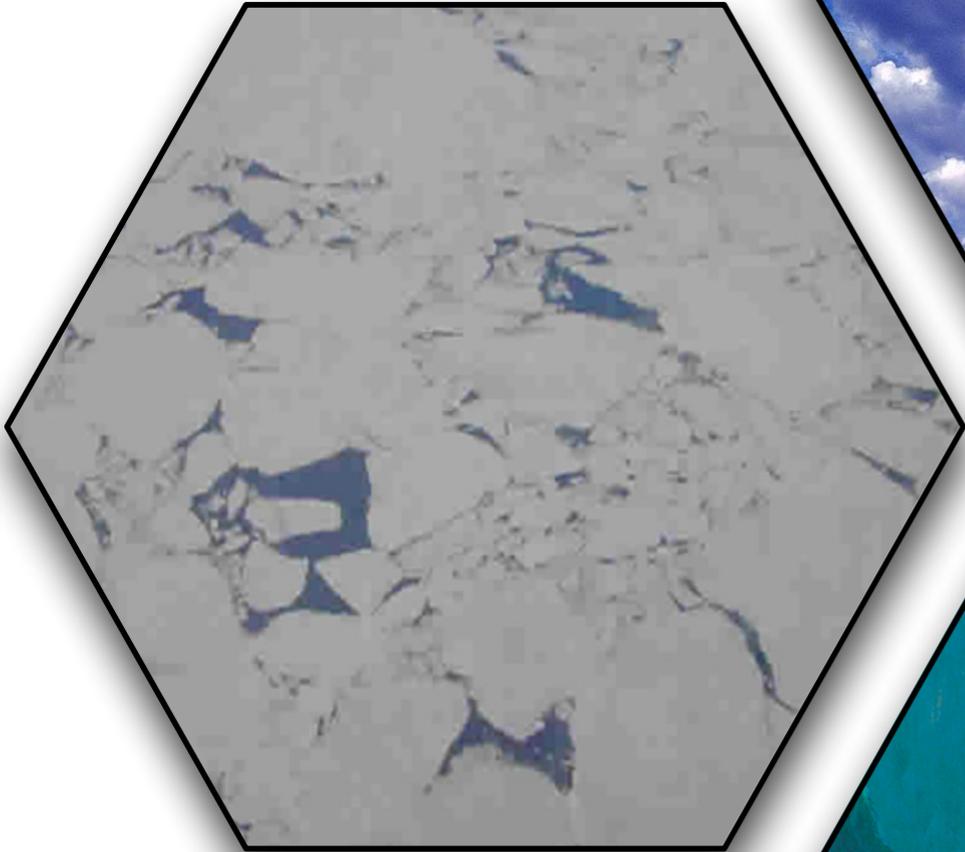
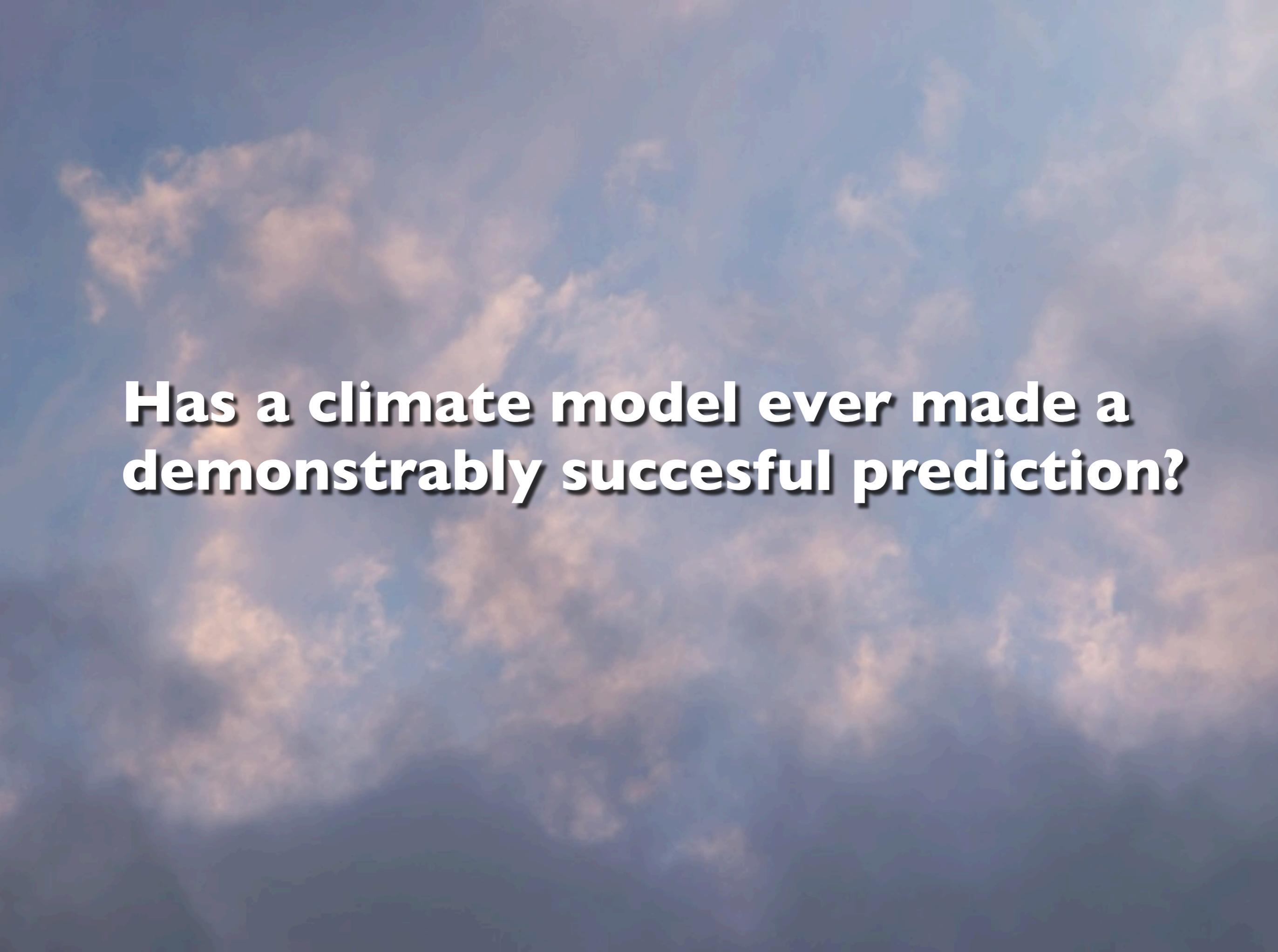


The Physical Climate System







Has a climate model ever made a demonstrably successful prediction?

The Effects of Doubling the CO₂ Concentration on the Climate of a General Circulation Model¹

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(Manuscript received 6 June 1974, in revised form 8 August 1974)

ABSTRACT

An attempt is made to estimate the temperature changes resulting from doubling the present CO₂ concentration by the use of a simplified three-dimensional general circulation model. This model contains the following simplifications: a limited computational domain, an idealized topography, no heat transport by ocean currents, and fixed cloudiness. Despite these limitations, the results from this computation yield some indication of how the increase of CO₂ concentration may affect the distribution of temperature in the atmosphere. It is shown that the CO₂ increase raises the temperature of the model troposphere, whereas it lowers that of the model stratosphere. The tropospheric warming is somewhat larger than that expected from a radiative-convective equilibrium model. In particular, the increase of surface temperature in higher latitudes is magnified due to the recession of the snow boundary and the thermal stability of the lower troposphere which limits convective heating to the lowest layer. It is also shown that the doubling of carbon dioxide significantly increases the intensity of the hydrologic cycle of the model.

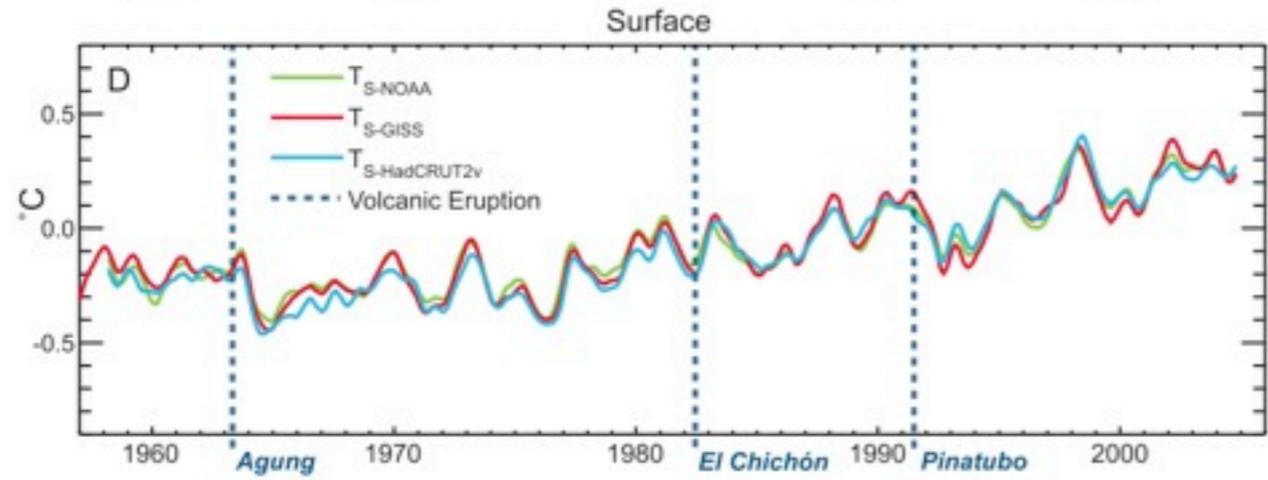
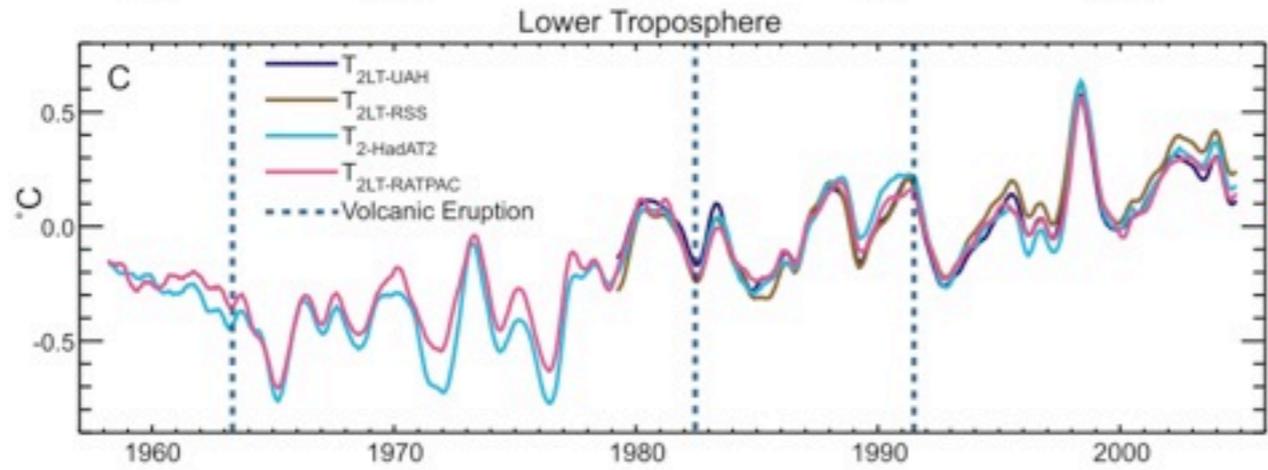
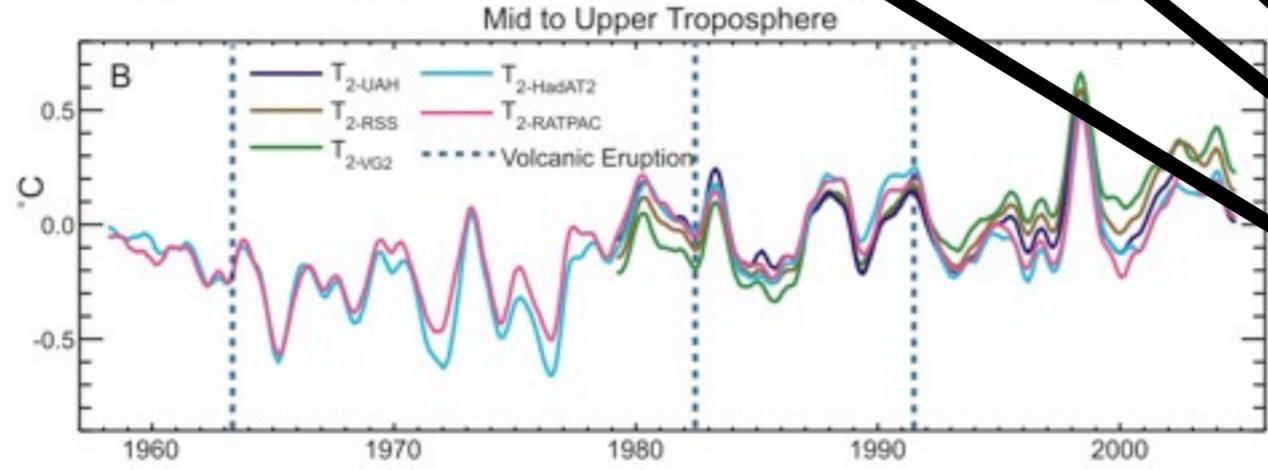
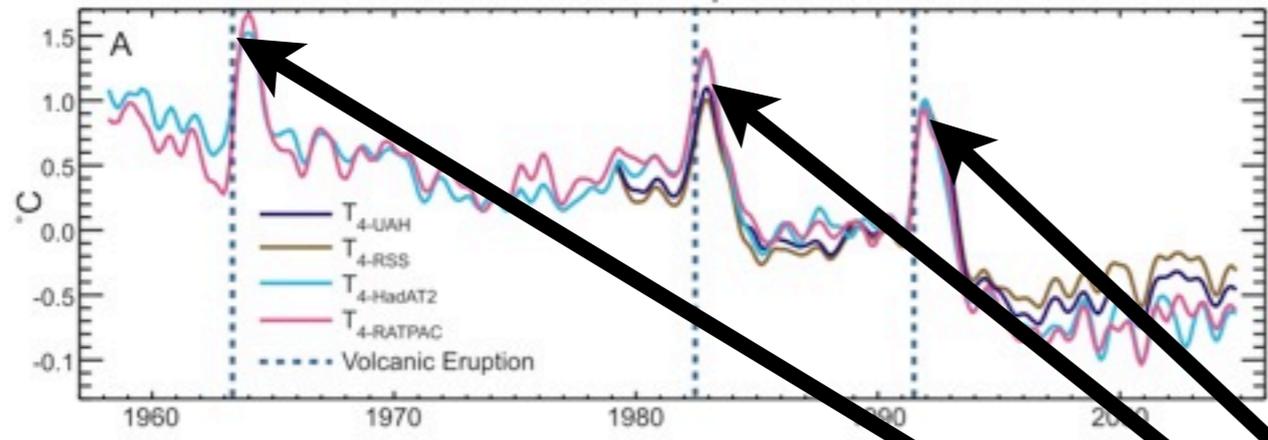
Manabe & Wetherald predicted:

- **Warming lower atmosphere**
- **Greater warming near the poles**
- **Cooling stratosphere**
- **More rain and higher humidity**



All of these things have now happened.

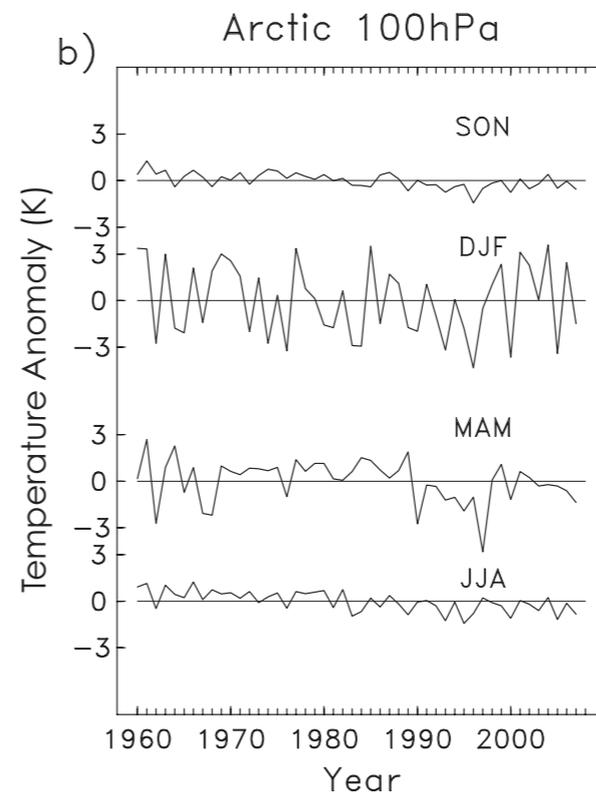
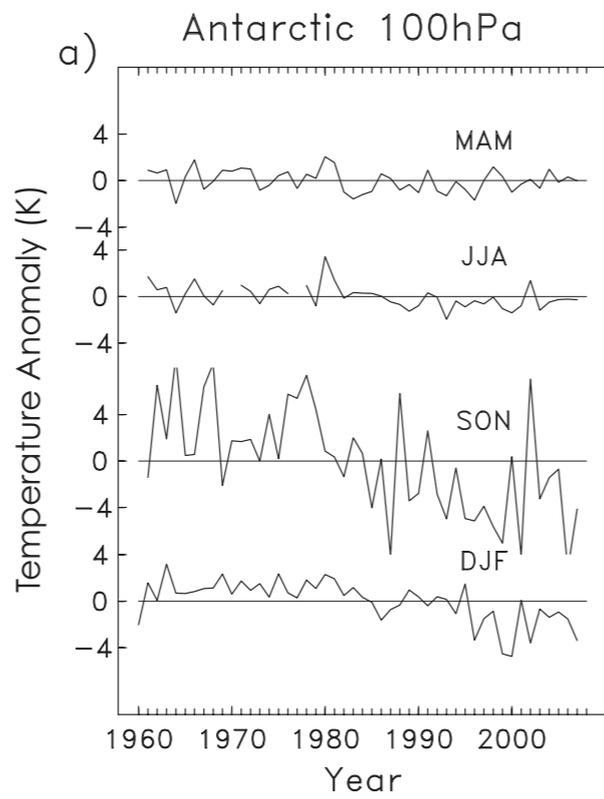
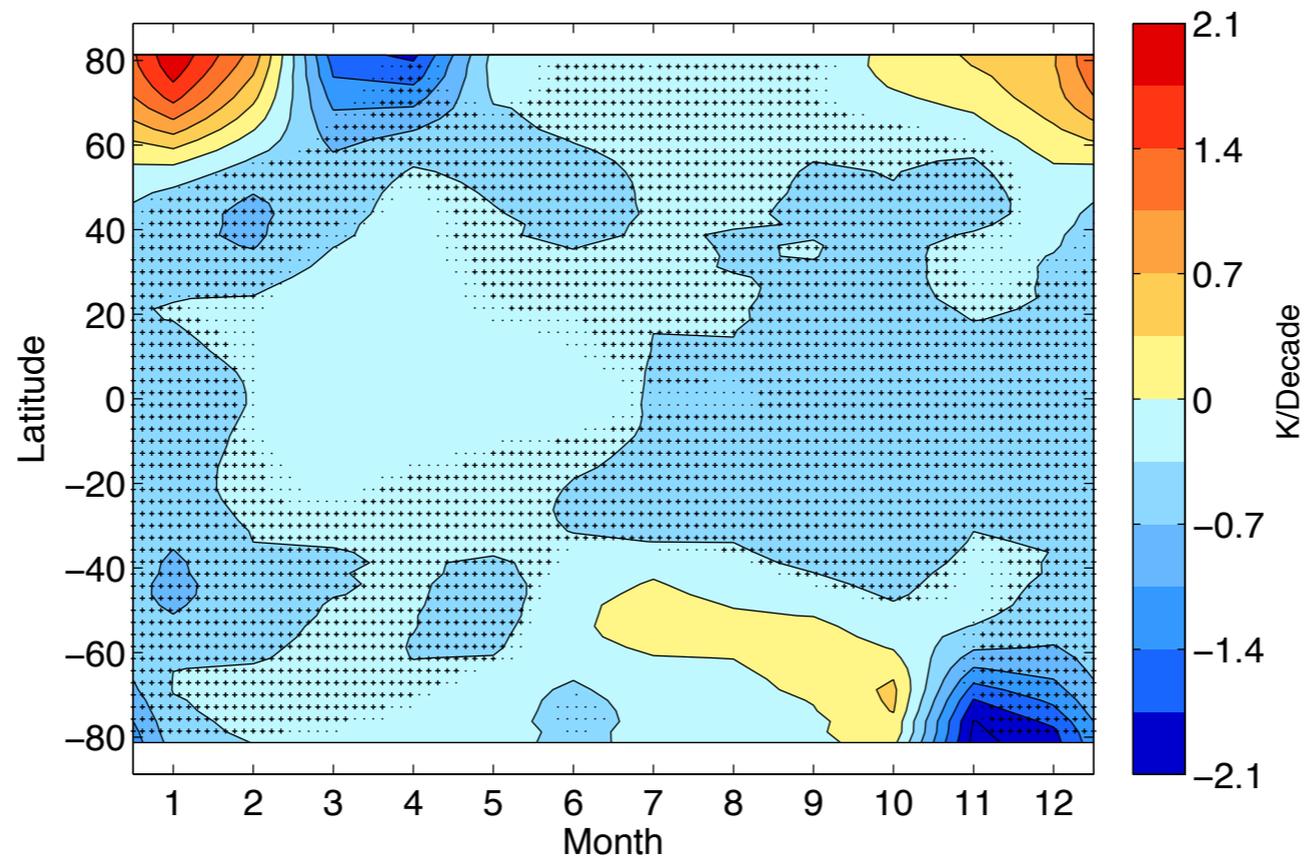
Global Anomalies



Stratospheric Cooling

Major volcanoes





Unprecedented Arctic ozone loss in 2011

Gloria L. Manney^{1,2}, Michelle L. Santee¹, Markus Rex³, Nathaniel J. Livesey¹, Michael C. Pitts⁴, Pepijn Veefkind^{5,6}, Eric R. Nash⁷, Ingo Wohltmann³, Ralph Lehmann³, Lucien Froidevaux¹, Lamont R. Poole⁸, Mark R. Schoeberl⁹, David P. Haffner⁷, Jonathan Davies¹⁰, Valery Dorokhov¹¹, Hartwig Gernandt³, Bryan Johnson¹², Rigel Kivi¹³, Esko Kyrö¹³, Niels Larsen⁴, Pieternel F. Levelt^{5,6,15}, Alexander Makshtas¹⁶, C. Thomas McElroy¹⁰, Hideaki Nakajima¹⁷, Maria Concepción Parrondo¹⁸, David W. Tarasick¹⁰, Peter von der Gathen³, Kaley A. Walker¹⁹ & Nikita S. Zinoviev¹⁶

Chemical ozone destruction occurs over both polar regions in local winter–spring. In the Antarctic, essentially complete removal of lower-stratospheric ozone currently results in an ozone hole every year, whereas in the Arctic, ozone loss is highly variable and has until now been much more limited. Here we demonstrate that chemical ozone destruction over the Arctic in early 2011 was—for the first time in the observational record—comparable to that in the Antarctic ozone hole. Unusually long-lasting cold conditions in the Arctic lower stratosphere led to persistent enhancement in ozone-destroying forms of chlorine and to unprecedented ozone loss, which exceeded 80 per cent over 18–20 kilometres altitude. Our results show that Arctic ozone holes are possible even with temperatures much milder than those in the Antarctic. We cannot at present predict when such severe Arctic ozone depletion may be matched or exceeded.

Since the emergence of the Antarctic ‘ozone hole’ in the 1980s¹ and elucidation of the chemical mechanisms^{2–5} and meteorological conditions⁶ involved in its formation, the likelihood of extreme ozone depletion over the Arctic has been debated. Similar processes are at work in the polar lower stratosphere in both hemispheres, but differences in the evolution of the winter polar vortex and associated polar temperatures have in the past led to vastly disparate degrees of spring-time ozone destruction in the Arctic and Antarctic. We show that chemical ozone loss in spring 2011 far exceeded any previously observed over the Arctic. For the first time, sufficient loss occurred to reasonably be described as an Arctic ozone hole.

Arctic polar processing in 2010–11

In the winter polar lower stratosphere, low temperatures induce condensation of water vapour and nitric acid (HNO₃) into polar stratospheric clouds (PSCs). PSCs and other cold aerosols provide surfaces for heterogeneous conversion of chlorine from longer-lived reservoir species, such as chlorine nitrate (ClONO₂) and hydrogen chloride (HCl), into reactive (ozone-destroying) forms, with chlorine monoxide (ClO) predominant in daylight^{5,7}.

In the Antarctic, enhanced ClO is usually present for 4–5 months (through to the end of September)^{8–11}, leading to destruction of most of the ozone in the polar vortex between ~14 and 20 km altitude⁷. Although ClO enhancement comparable to that in the Antarctic occurs at some times and altitudes in most Arctic winters⁹, it rarely persists for more than 2–3 months, even in the coldest years¹⁰. Thus chemical ozone loss in the Arctic has until now been limited, with largest previous losses observed in 2005, 2000 and 1996^{7,12–14}.

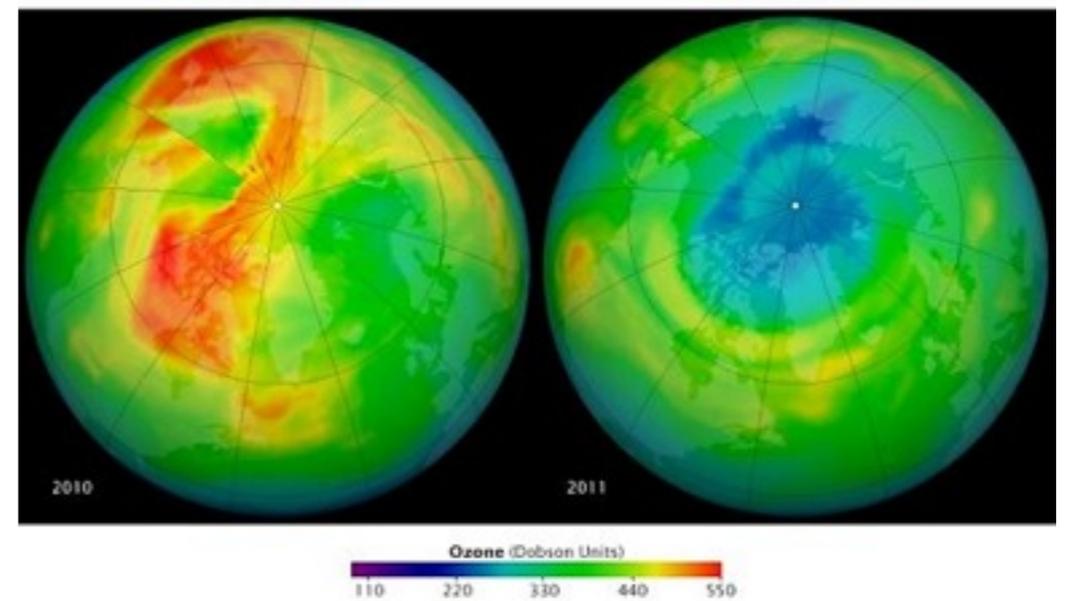
The 2010–11 Arctic winter–spring was characterized by an anomalously strong stratospheric polar vortex and an atypically long continuously cold period. In February–March 2011, the barrier to

transport at the Arctic vortex edge was the strongest in either hemisphere in the last ~30 years (Fig. 1a, Supplementary Discussion).

The persistence of a strong, cold vortex from December through to the end of March was unprecedented. In the previous years with most ozone loss, temperatures (T) rose above the threshold associated with chlorine activation (T_{act} , near 196 K, roughly the threshold for the potential existence of PSCs) by early March (Fig. 1b, Supplementary Figs 1, 2). Only in 2011 and 1997 have Arctic temperatures below T_{act} persisted through to the end of March, sporadically approaching a vortex volume fraction similar in size to that in some Antarctic winters (Fig. 1b). In 1996–97, however, the cold volume remained very limited until mid-January and was smaller than that in 2011 at most times during late January through to the end of March (Fig. 1b, Supplementary Figs 1, 2).

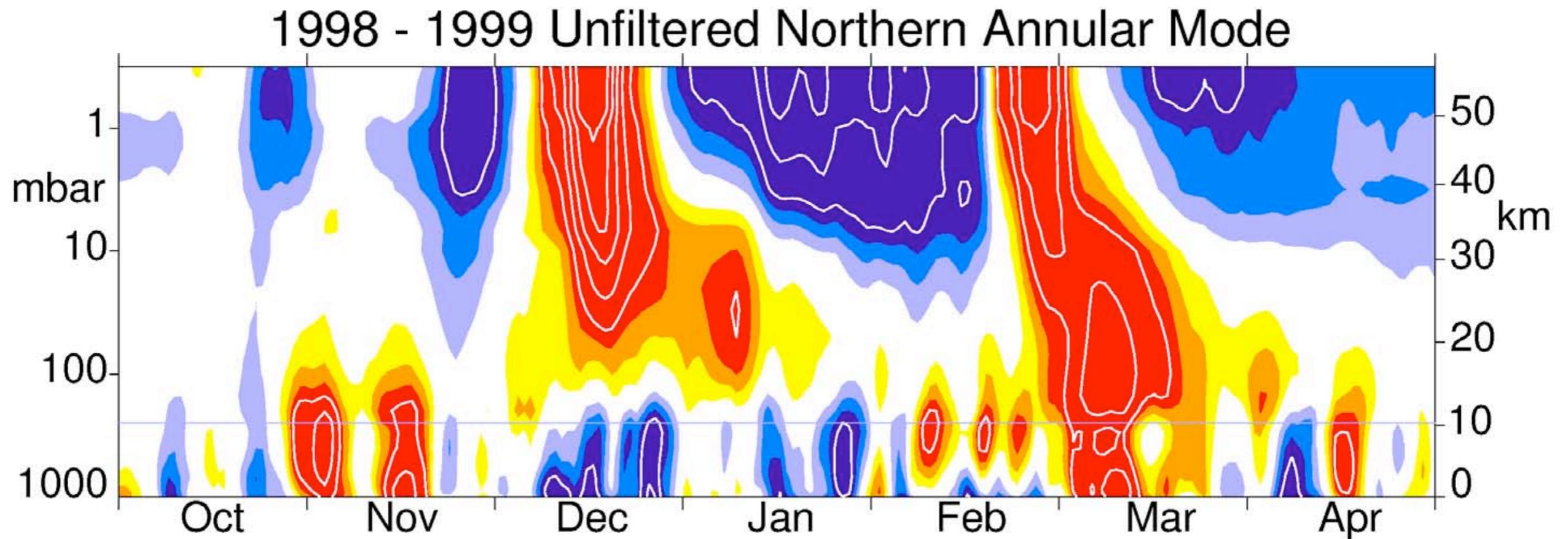
Daily minimum temperatures in the 2010–11 Arctic winter were not unusually low, but the persistently cold region was remarkably deep (Supplementary Figs 1, 2). Temperatures were below T_{act} for more than 100 days over an altitude range of ~15–23 km, compared to a similarly prolonged cold period over only ~20–23 km altitude in 1997; below ~19 km altitude, $T < T_{act}$ continued for ~30 days longer in 2011 than in 1997 (Supplementary Fig. 1b). In 2005, the previous year with largest Arctic ozone loss⁷, $T < T_{act}$ occurred for more than 100 days over ~17–23 km altitude, but all before early March.

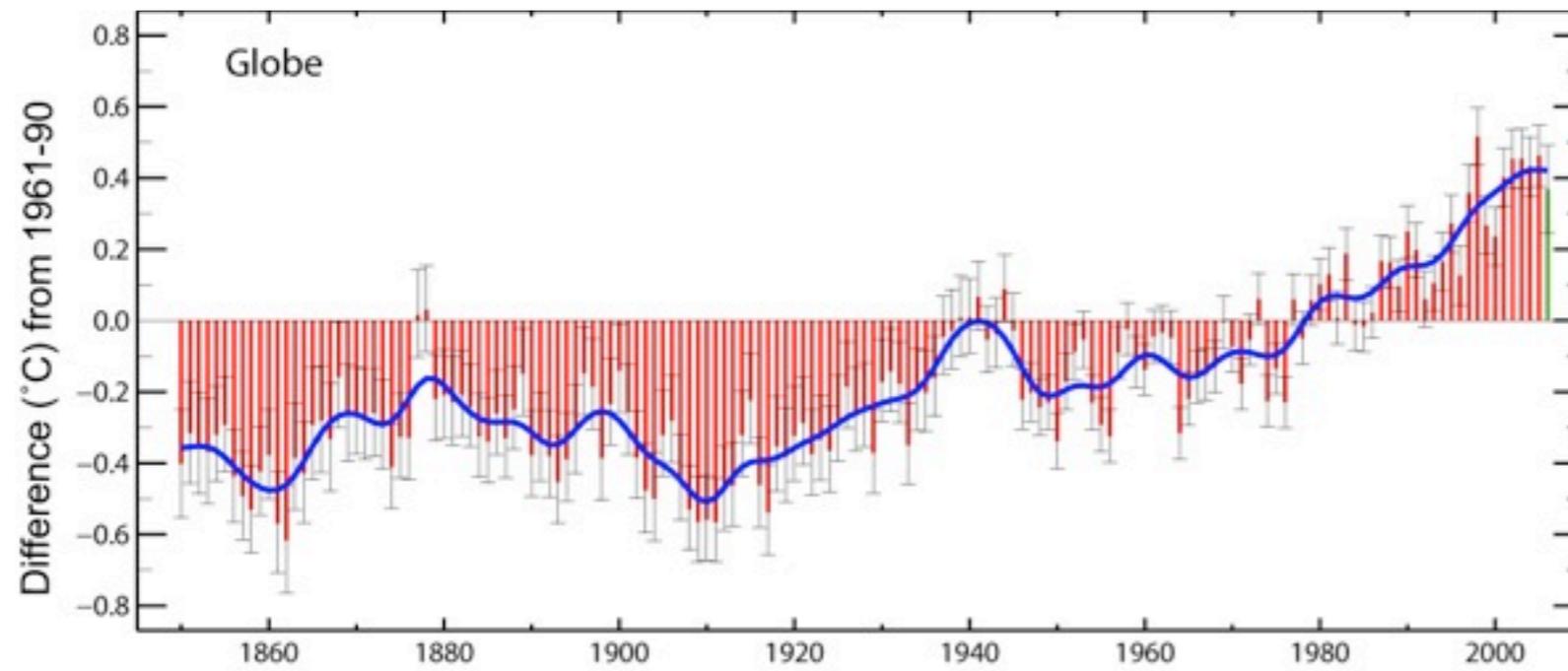
The winter mean volume of air in which PSCs may form (that is, with $T < T_{act}$), V_{psc} , is closely correlated with the potential for ozone loss^{7,15–17}. In 2011, V_{psc} (as a fraction of the vortex volume) was the largest on record (Fig. 1c). Both large V_{psc} and cold lingering well into spring are important in producing severe chemical loss^{7,15,16}, and 2010–11 was the only Arctic winter during which both conditions have been met. Much lower fractional V_{psc} in 1997 than in 1996, 2000, 2005 or 2011 (Fig. 1c) is consistent with less ozone loss that year^{16,17}.



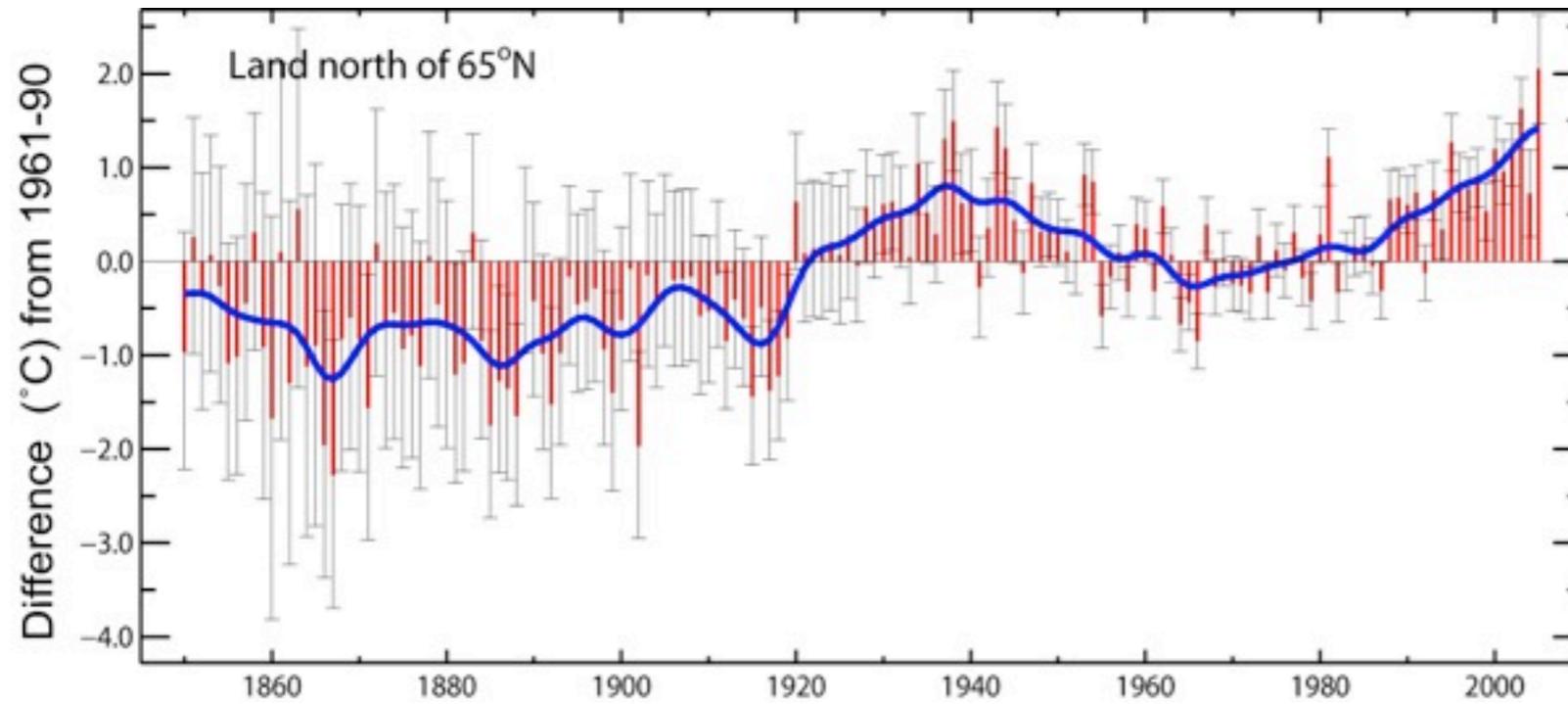
¹Jet Propulsion Laboratory, California Institute of Technology, Pasadena, California 91109, USA. ²New Mexico Institute of Mining and Technology, Socorro, New Mexico 87801, USA. ³Alfred Wegener Institute for Polar and Marine Research, D-14473 Potsdam, Germany. ⁴NASA Langley Research Center, Hampton, Virginia 23681, USA. ⁵Royal Netherlands Meteorological Institute, 3730 AE De Bilt, The Netherlands. ⁶Delft University of Technology, 2600 GA Delft, The Netherlands. ⁷Science Systems and Applications, Inc., Lanham, Maryland 20706, USA. ⁸Science Systems and Applications, Inc., Hampton, Virginia 23666, USA. ⁹Science and Technology Corporation, Lanham, Maryland 20706, USA. ¹⁰Environment Canada, Toronto, Ontario, Canada M3H 5T4. ¹¹Central Aerological Observatory, Dolgoprudny 141700, Russia. ¹²NOAA Earth System Research Laboratory, Boulder, Colorado 80305, USA. ¹³Arctic Research Center, Finnish Meteorological Institute, 99600 Sodankylä, Finland. ¹⁴Danish Climate Center, Danish Meteorological Institute, DK-2100 Copenhagen, Denmark. ¹⁵Eindhoven University of Technology, 5600 MB Eindhoven, The Netherlands. ¹⁶Arctic and Antarctic Research Institute, St Petersburg 199397, Russia. ¹⁷National Institute for Environmental Studies, Tsukuba-city, 305-8506, Japan. ¹⁸National Institute for Aerospace Technology, 28850 Torrejón De Ardoz, Spain. ¹⁹University of Toronto, Toronto, Ontario, Canada M5S 1A7.

Stratosphere-troposphere coupling

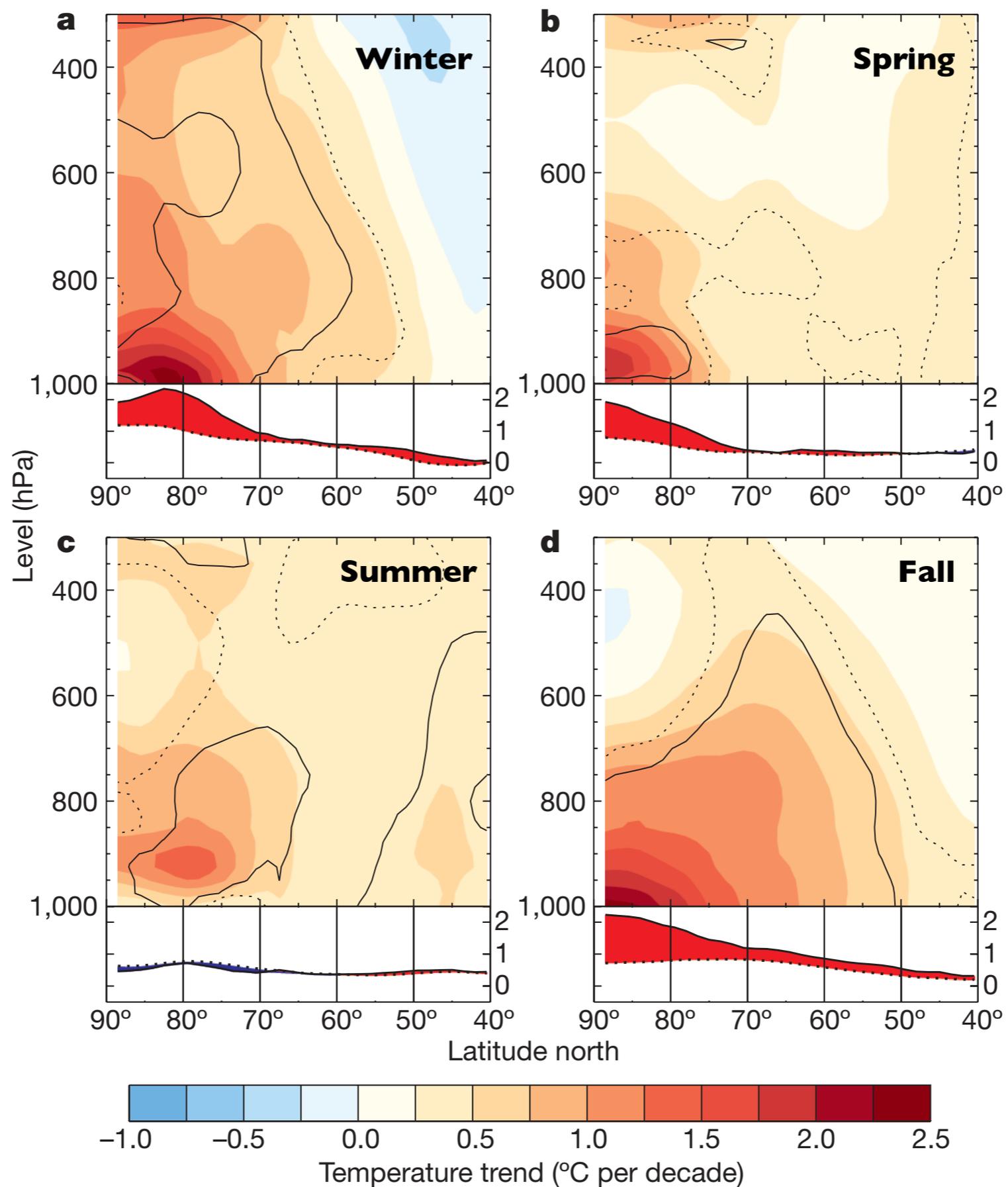




Warming in the Arctic is roughly double that for the whole Earth.

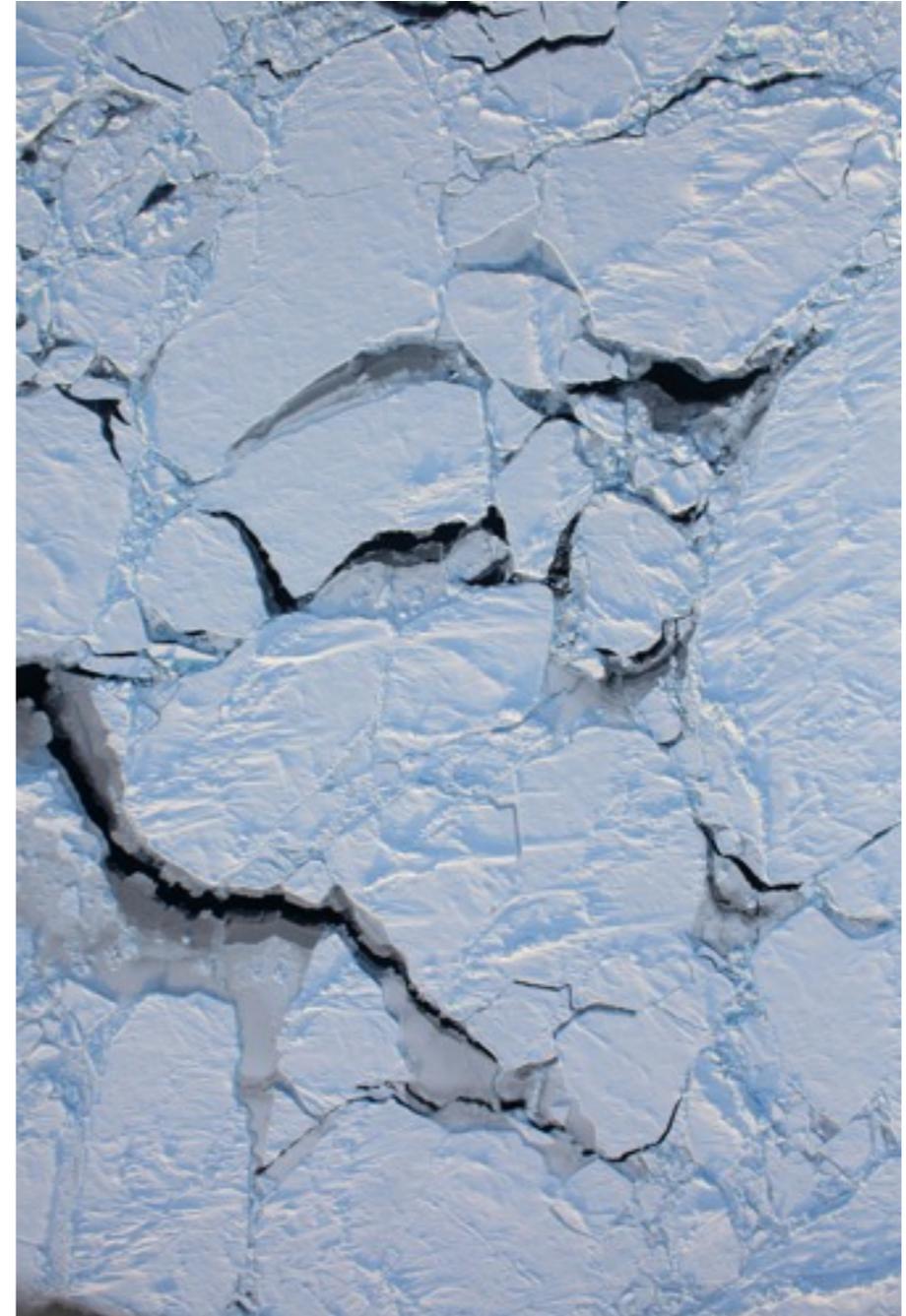


Note different scales.

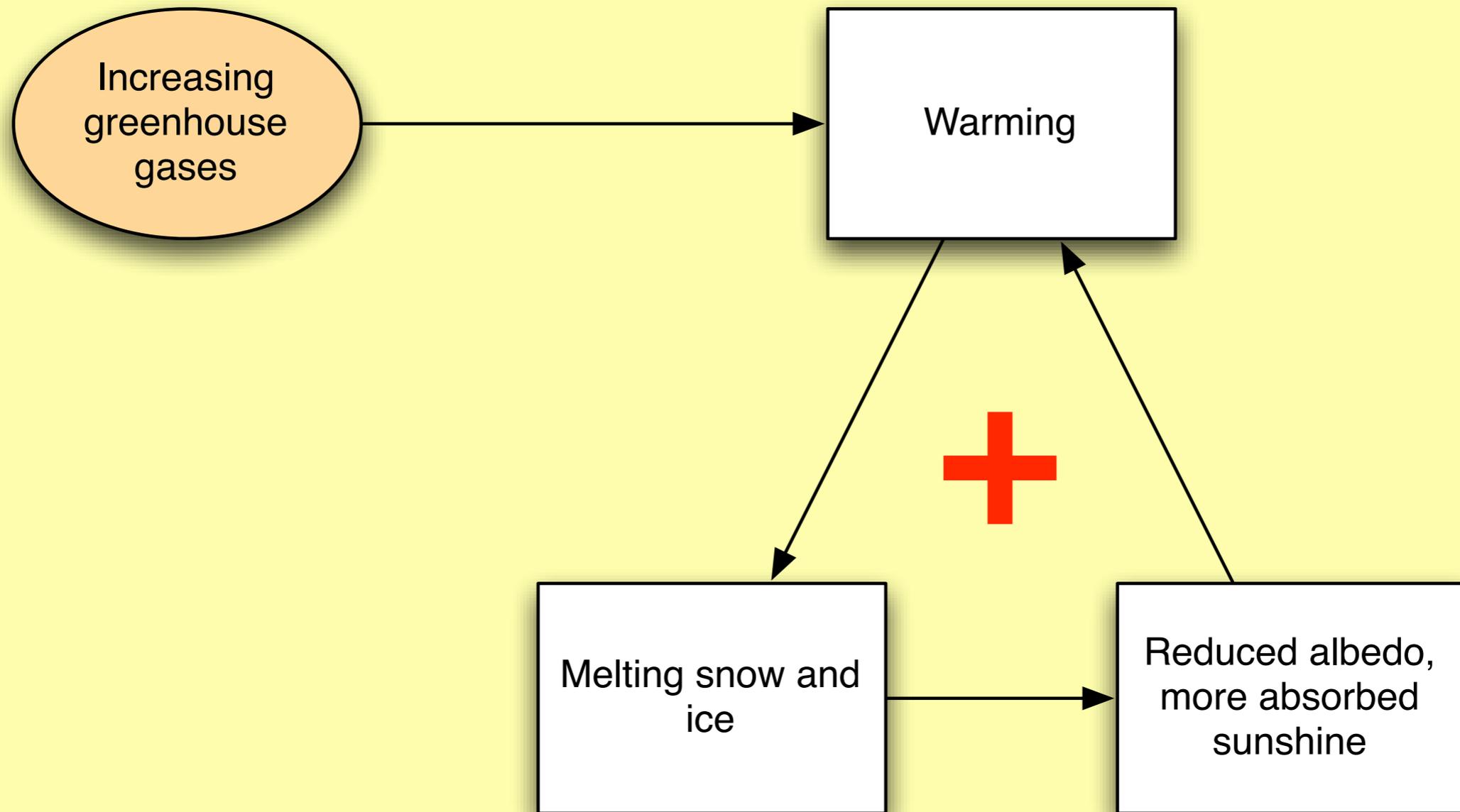


Sea ice

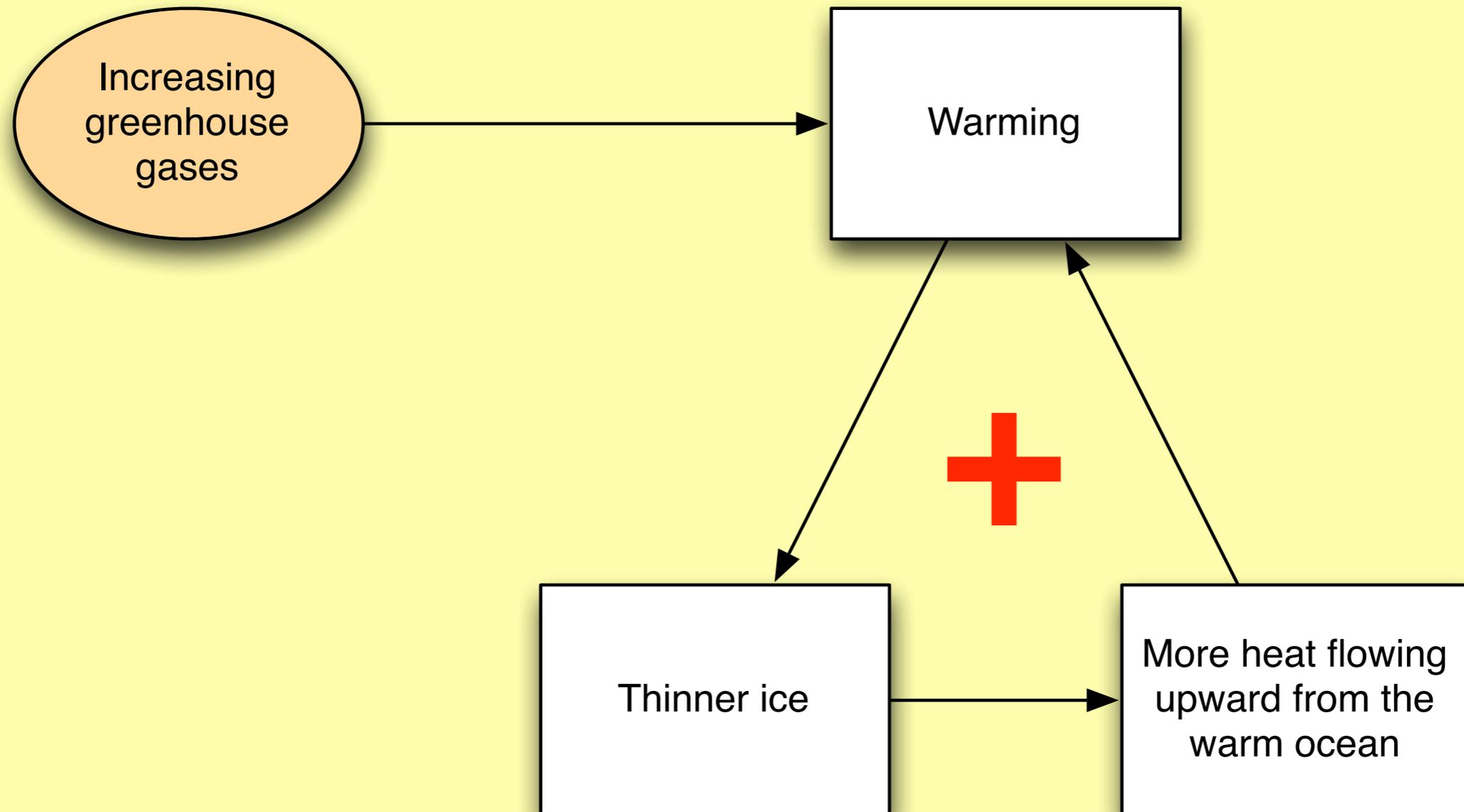
- **Reflects sunlight**
- **Blocks heat exchange between the ocean below and the air above**



Albedo Feedback



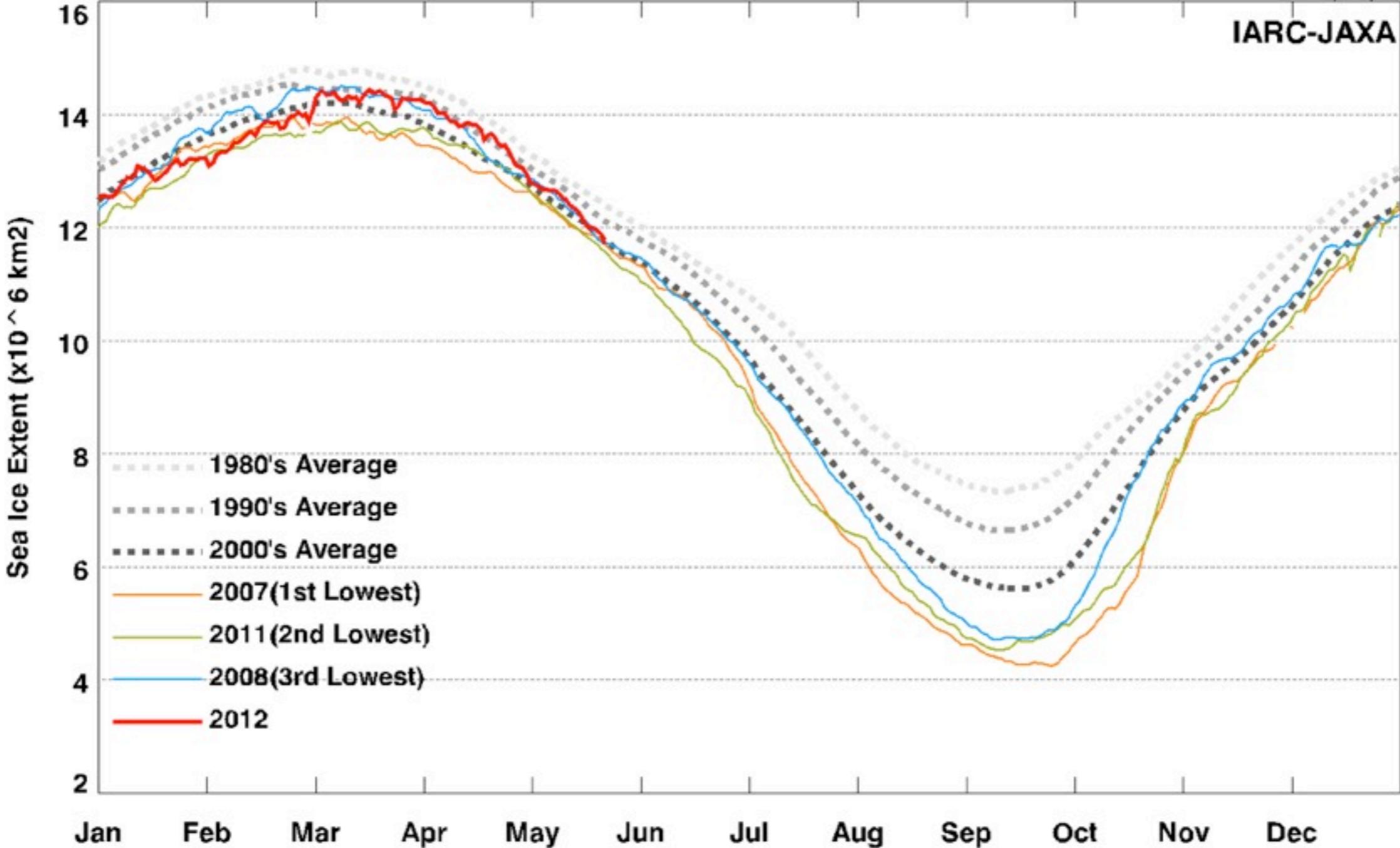
Sea Ice Insulation Feedback



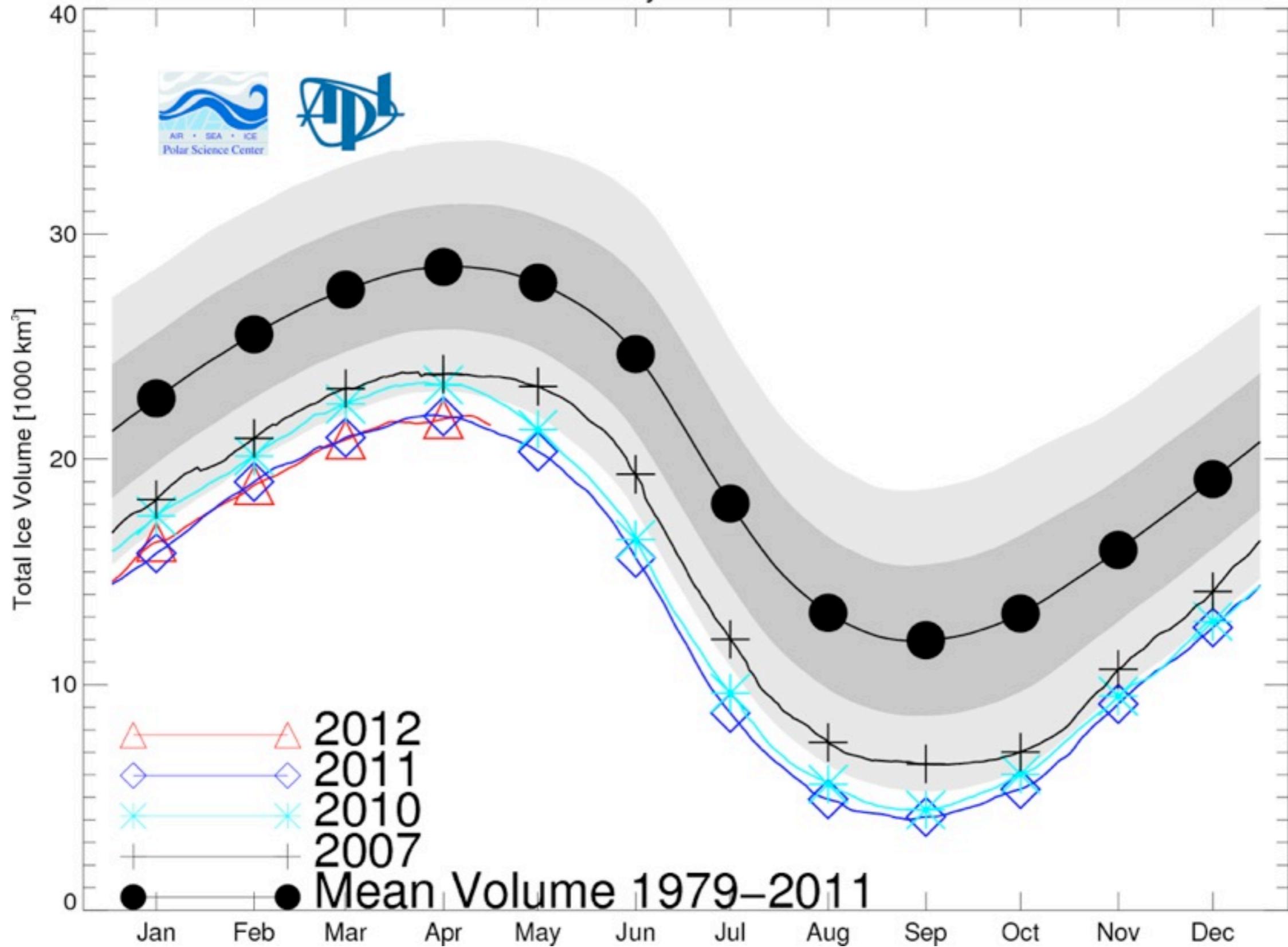
Arctic Sea Ice Extent

2012/05/22

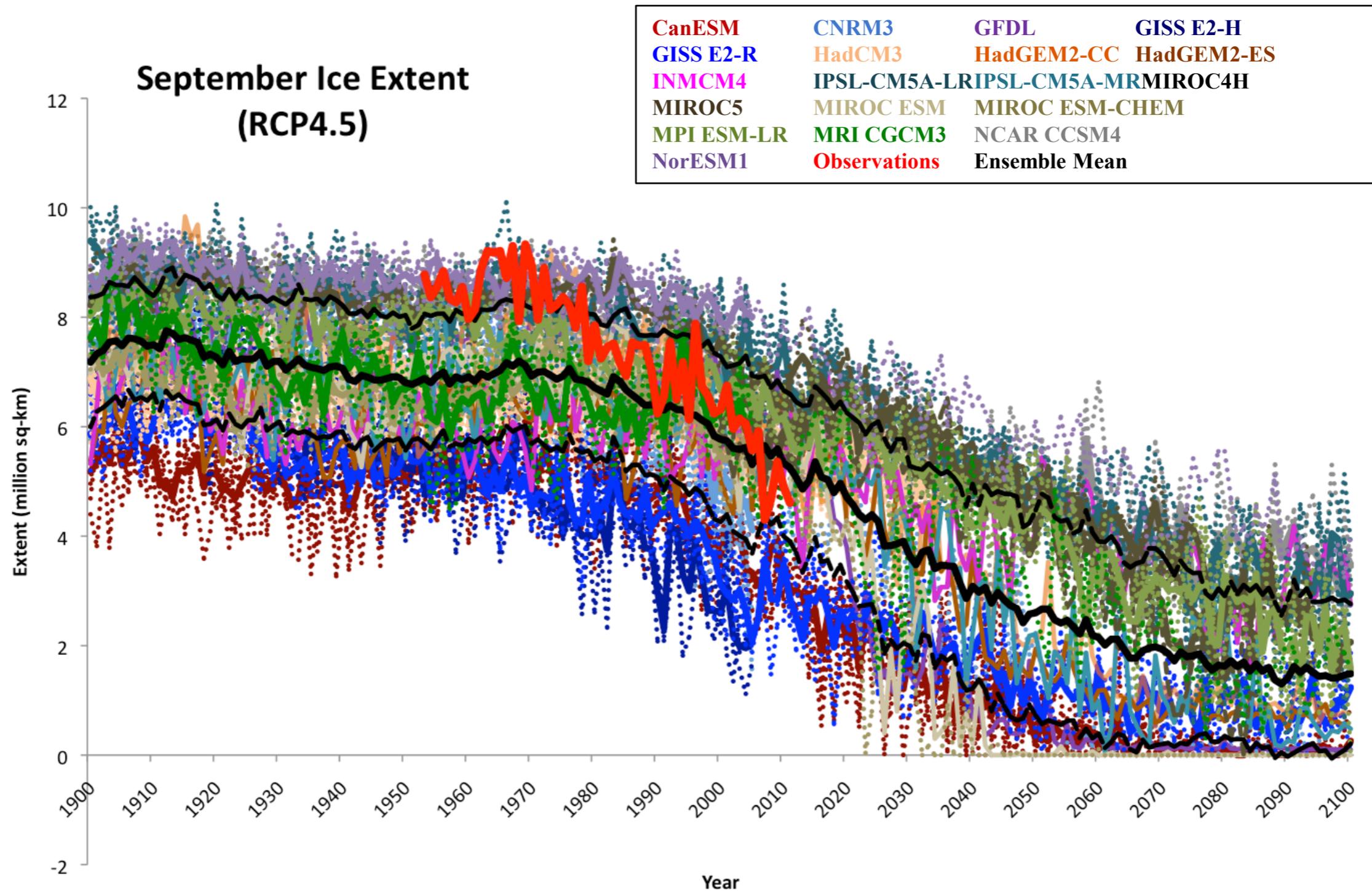
IARC-JAXA



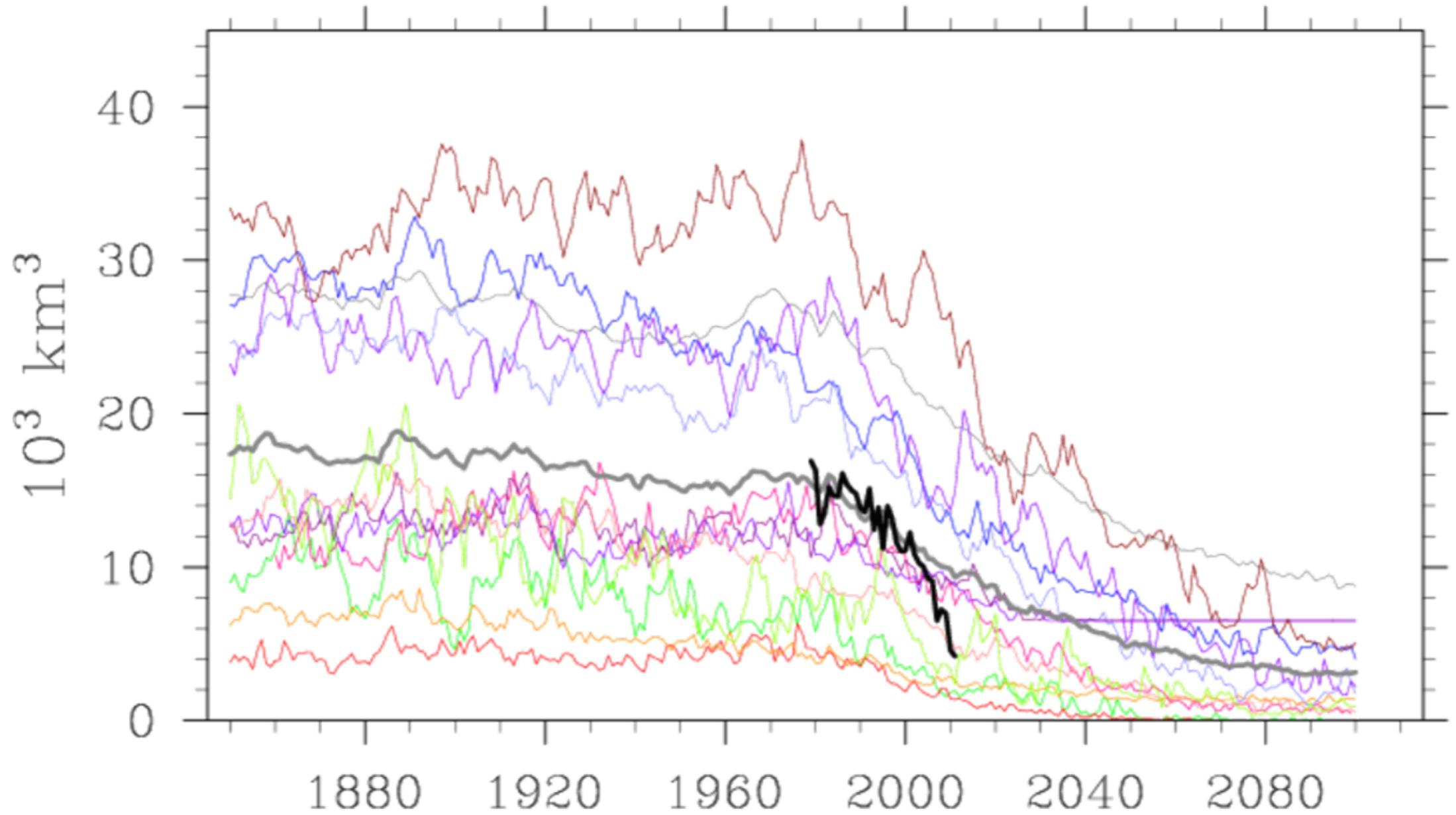
PIOMAS Daily Arctic Ice Volume



CMIP5 results

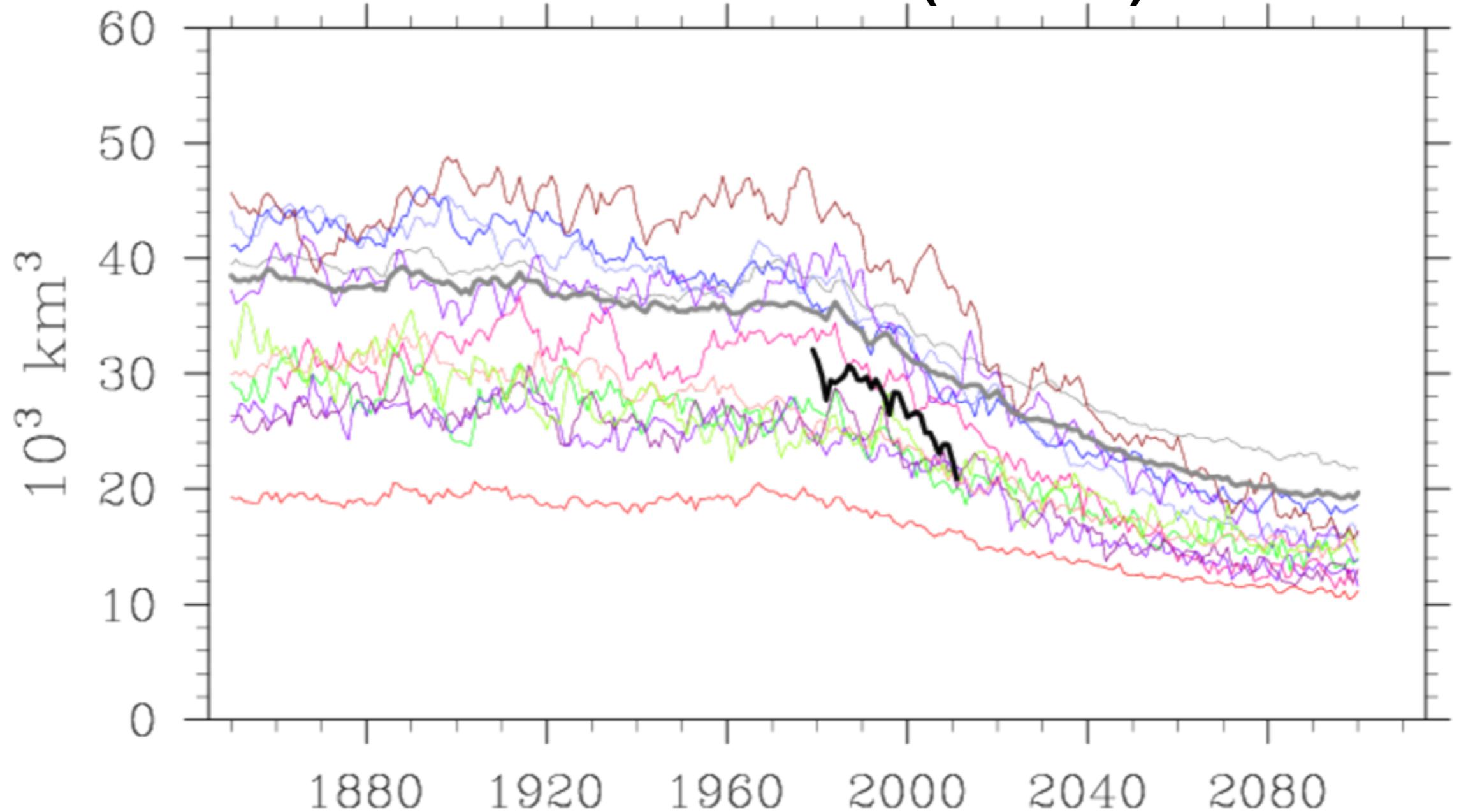


Summer Minimum (September)



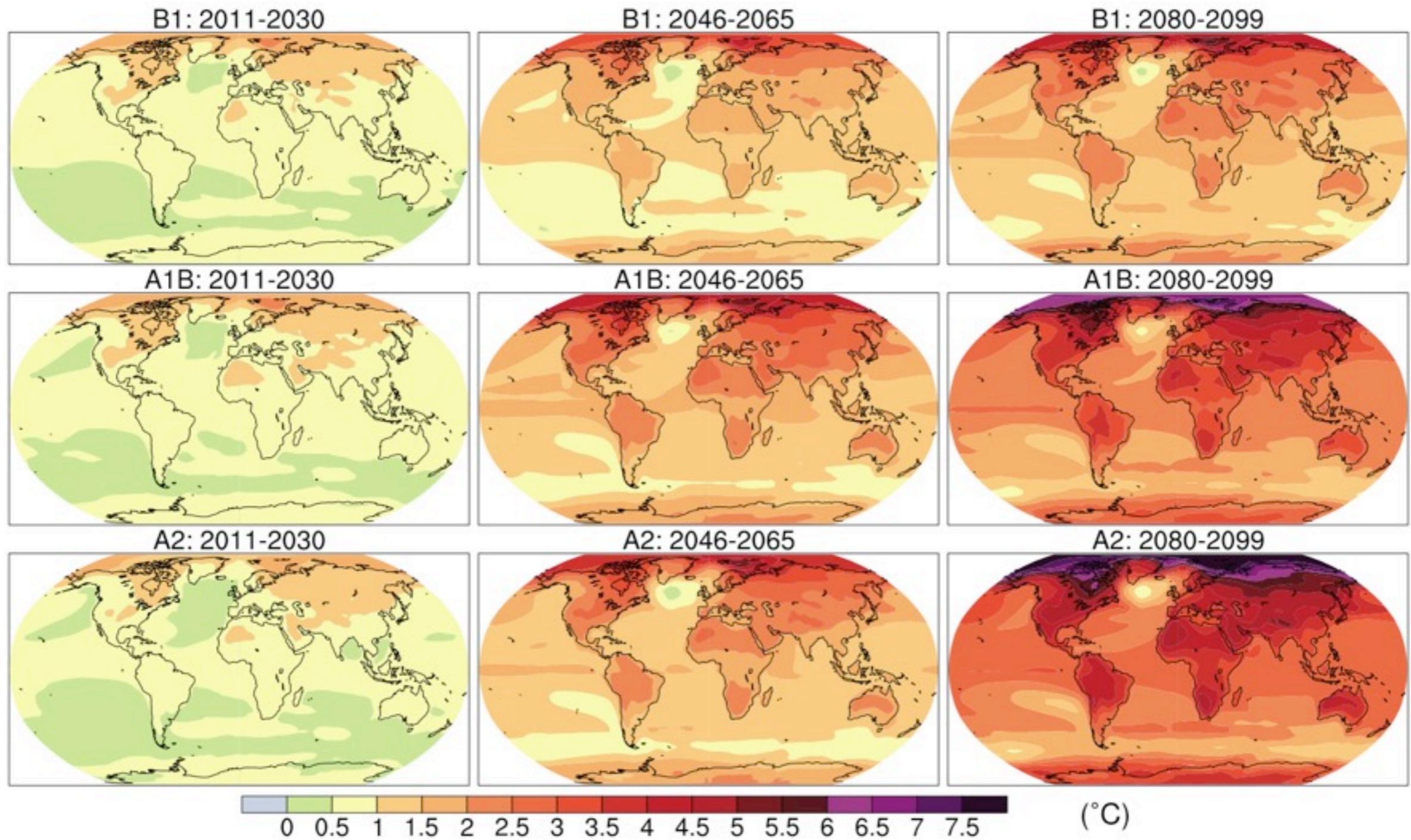
CCSM4 CNRM-CM5 CSIRO-Mk3-6-0 CanESM2 GISS-E2-R HadGEM2-ES
IPSL-CM5A-LR MIROC-ESM MIROC-ESM-CHEM MIROC5 MPI-ESM-LR MRI-CGCM3
NorESM1-M Ensemble Mean PIOMAS-simulated ice volume

Winter Maximum (March)

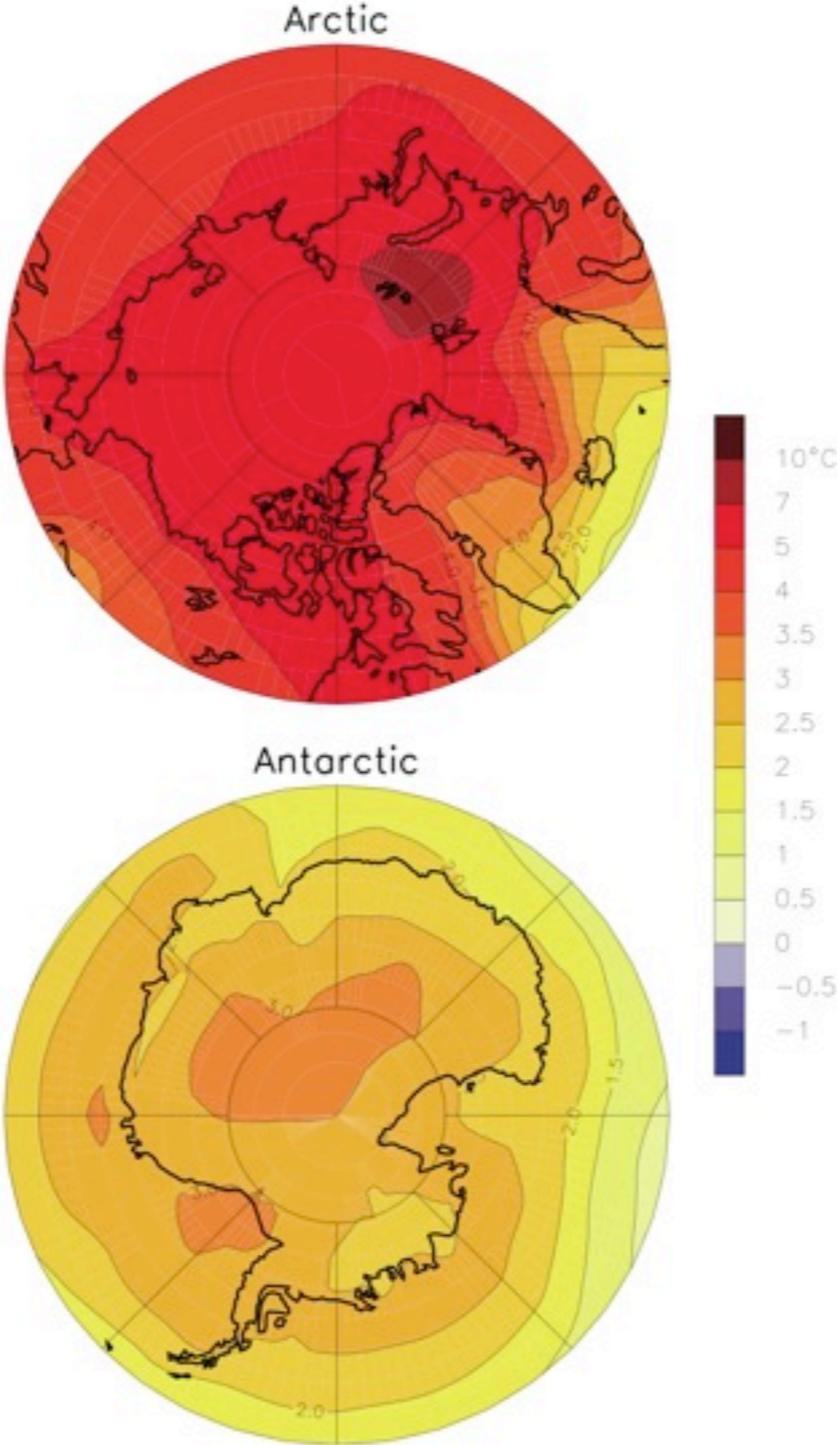


CCSM4 CNRM-CM5 CSIRO-Mk3-6-0 CanESM2 GISS-E2-R HadGEM2-ES
IPSL-CM5A-LR MIROC-ESM MIROC-ESM-CHEM MIROC5 MPI-ESM-LR MRI-CGCM3
NorESM1-M Ensemble Mean PIOMAS-simulated ice volume

Forecasts for the 21st Century

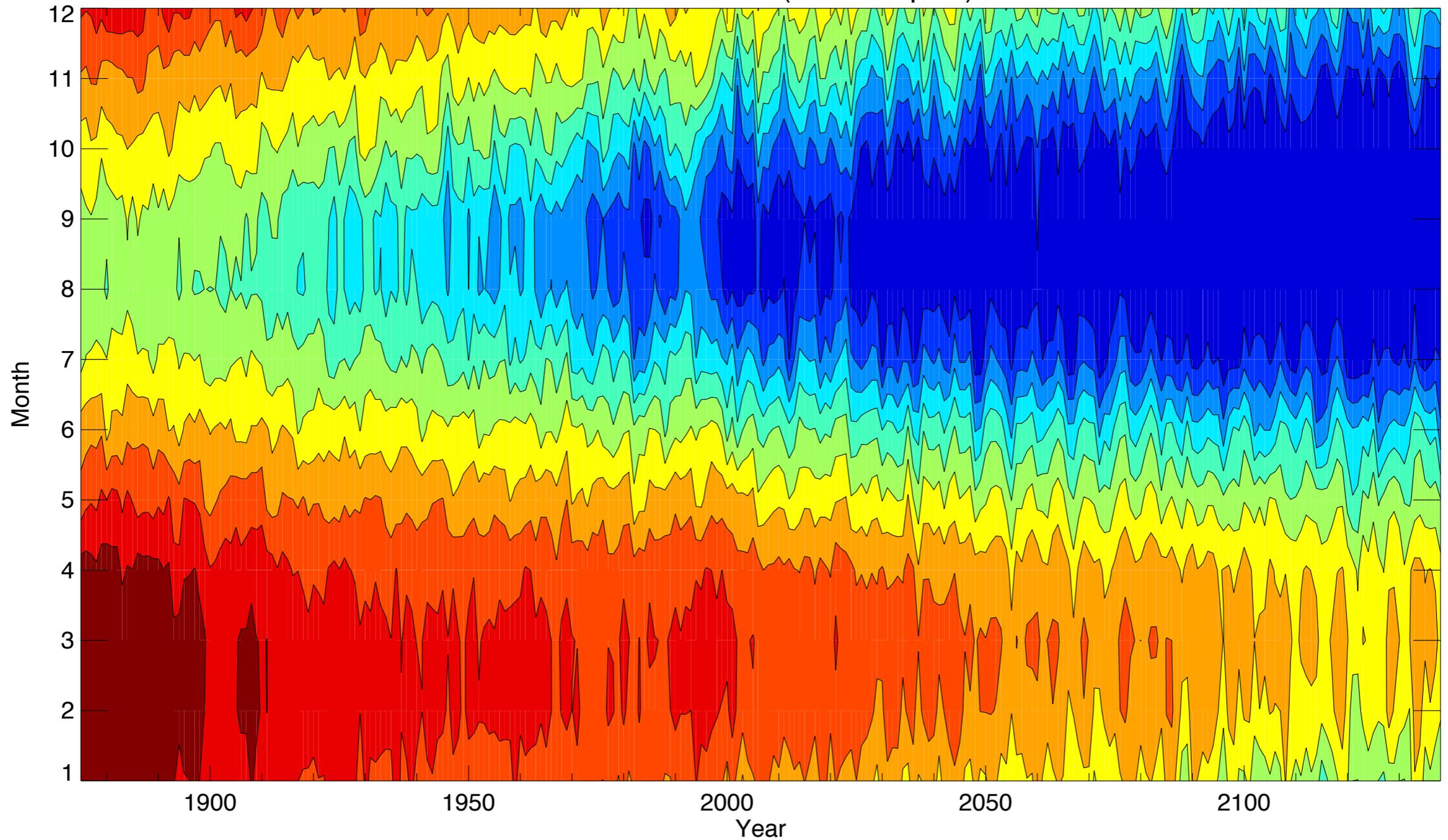


Predicted warming over the 21st century

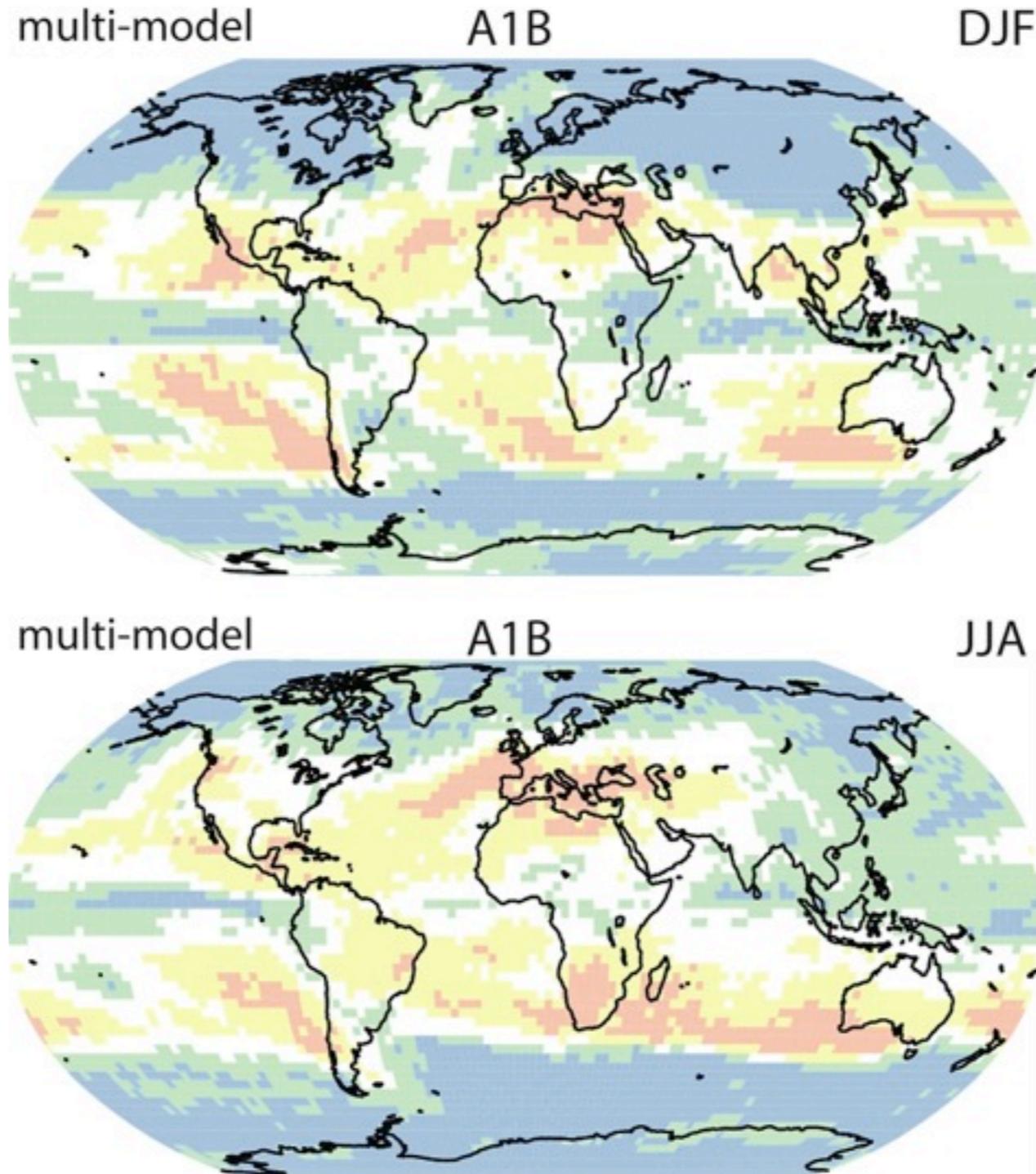


CESM 1% per year

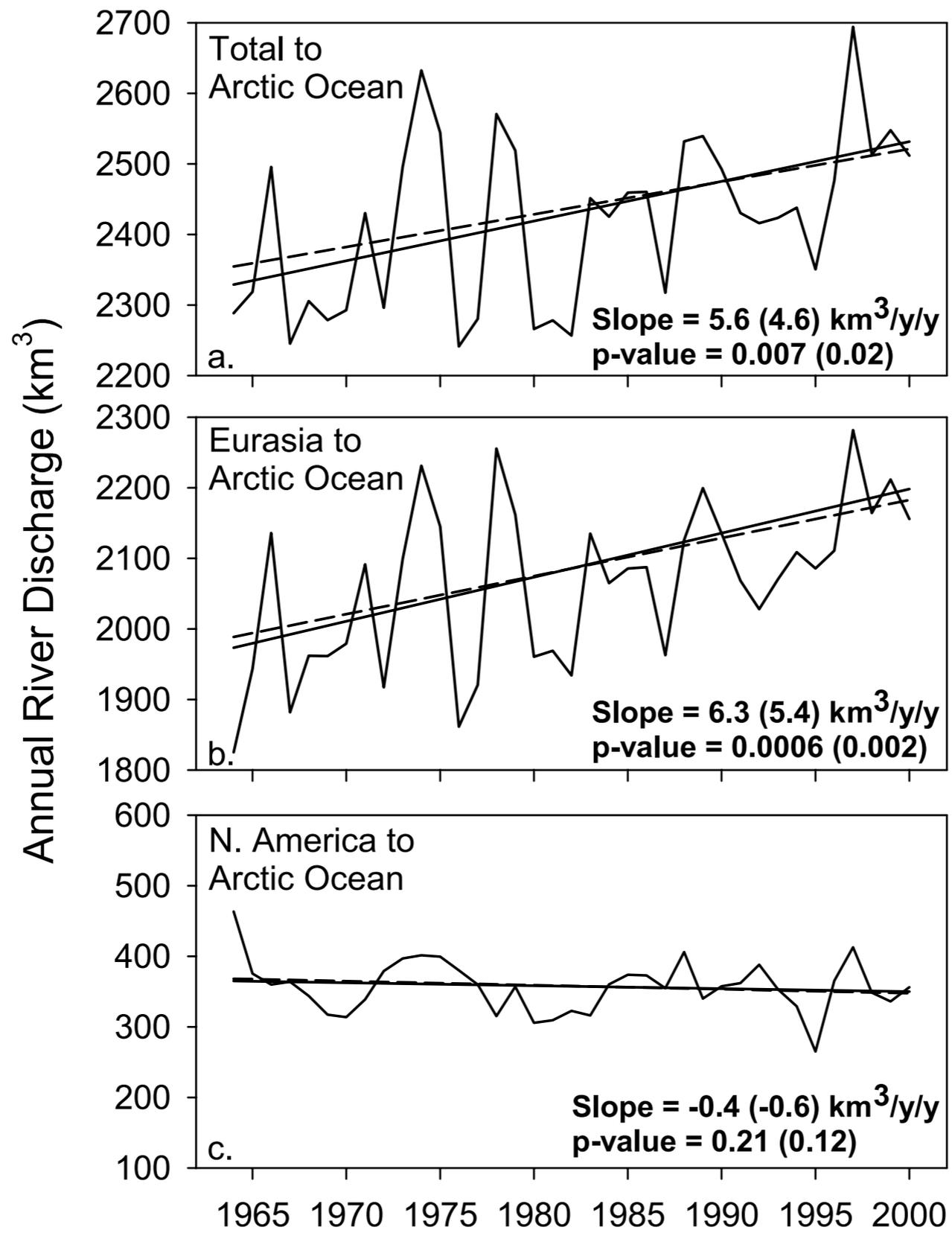
NH Sea Ice Area (million sq km)



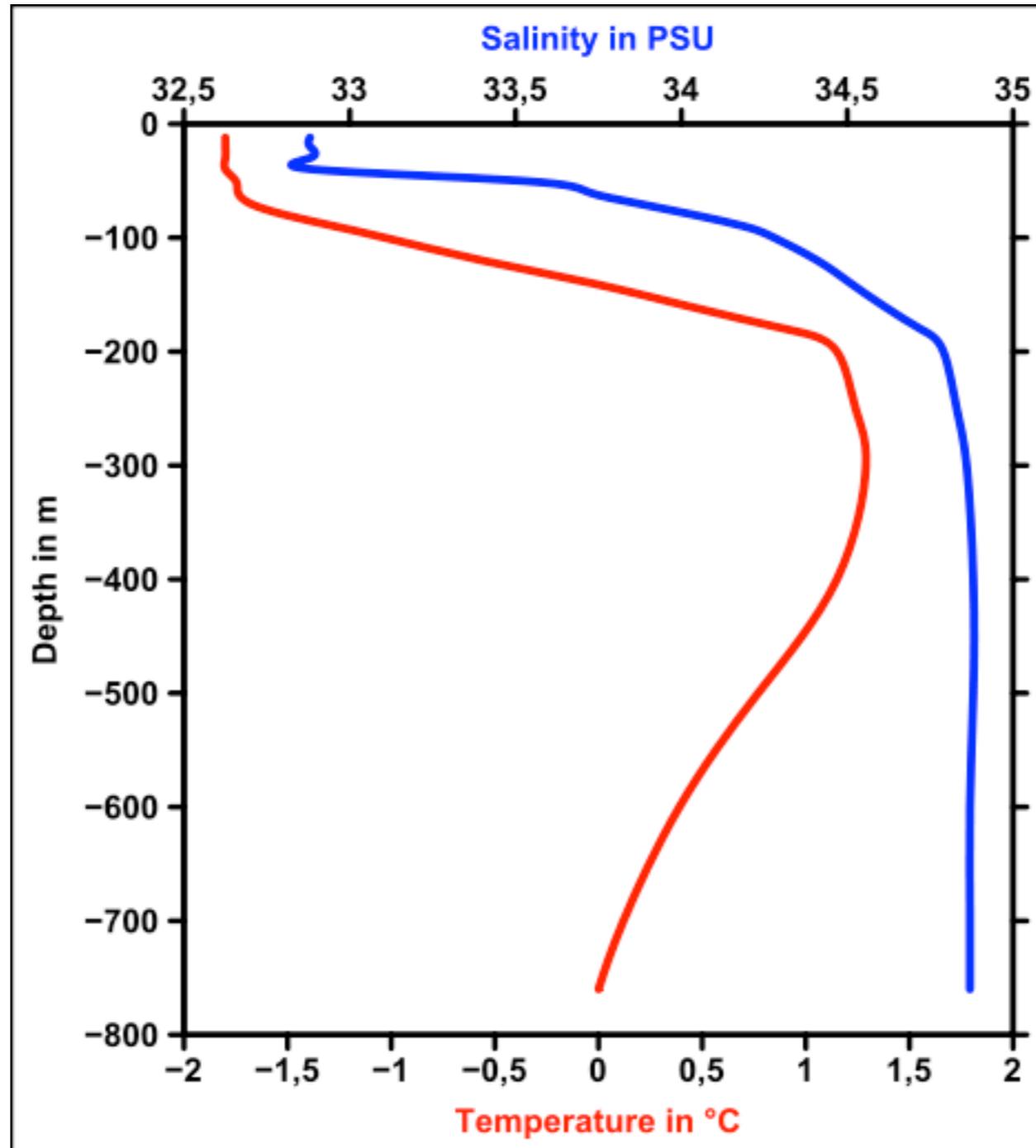
Precipitation: Late 21st century minus late 20th century



The wet get wetter and the dry get drier.

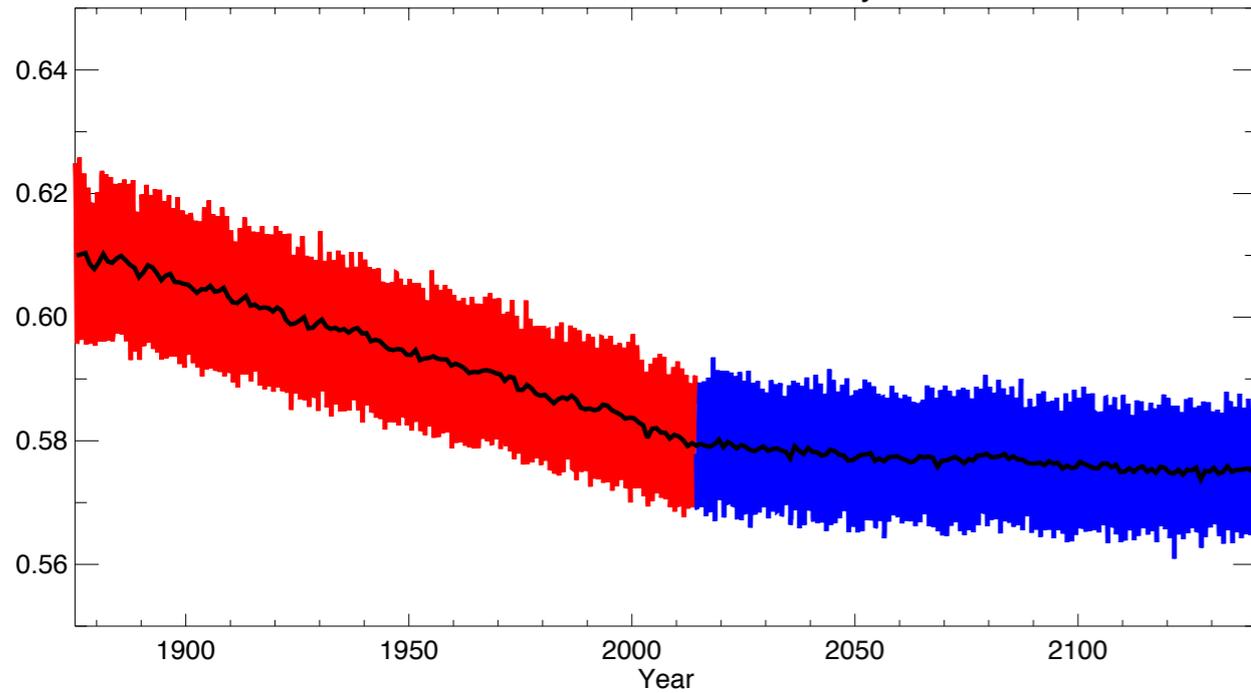


Halocline

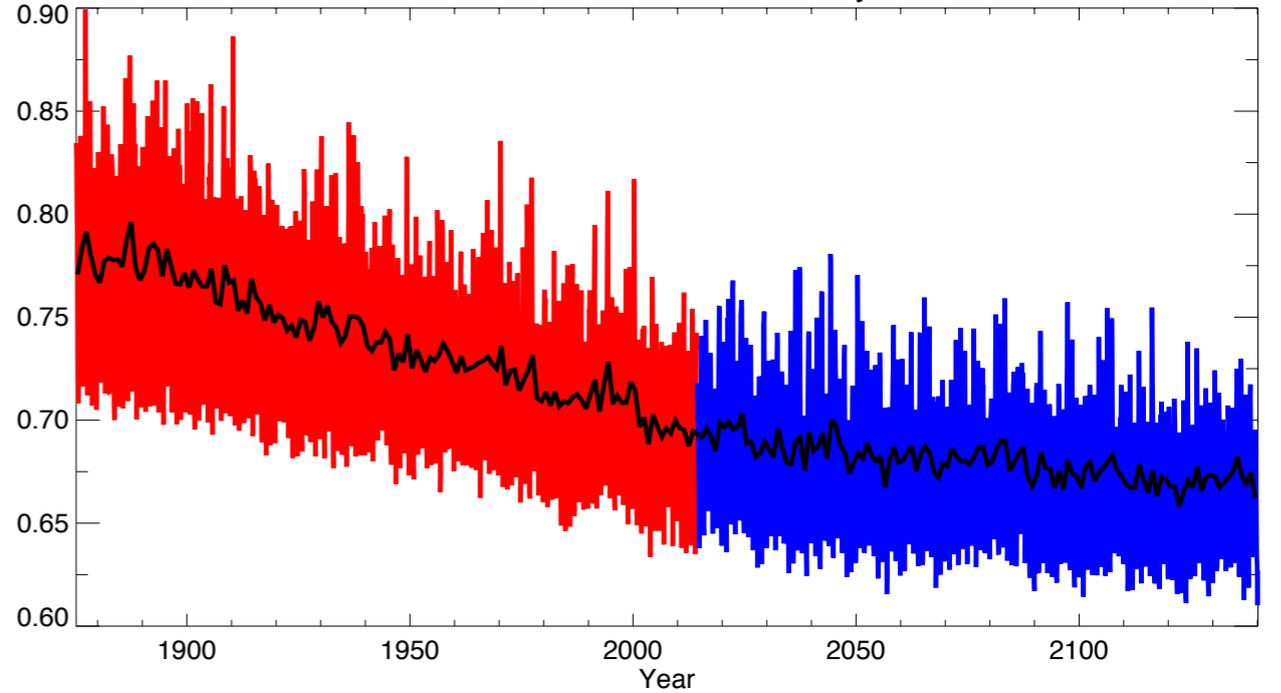


Two Greenhouse Indices

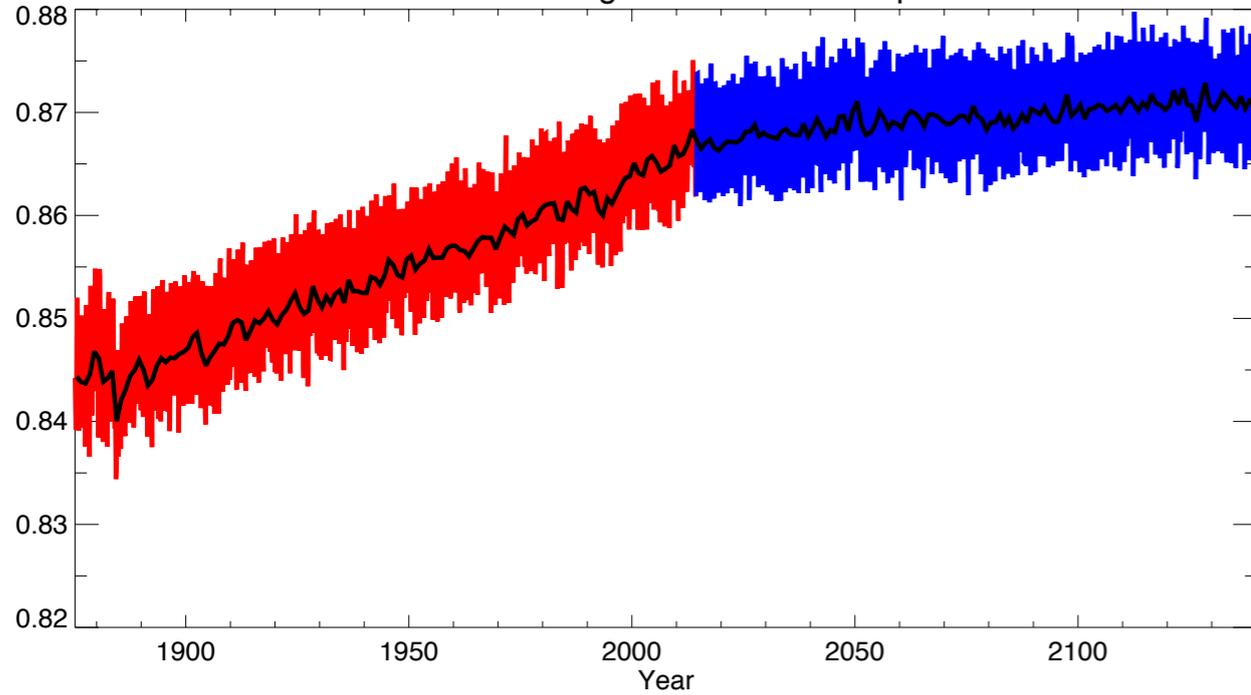
Global Mean Bulk Emissivity



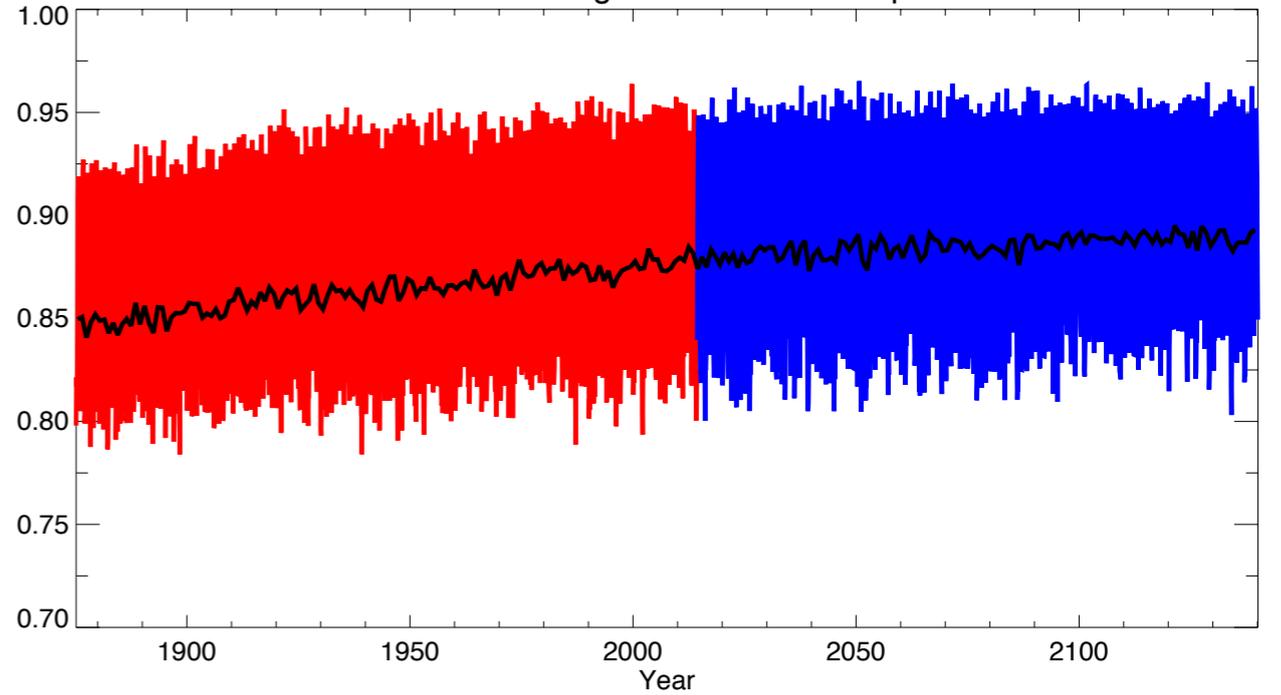
Arctic Mean Bulk Emissivity



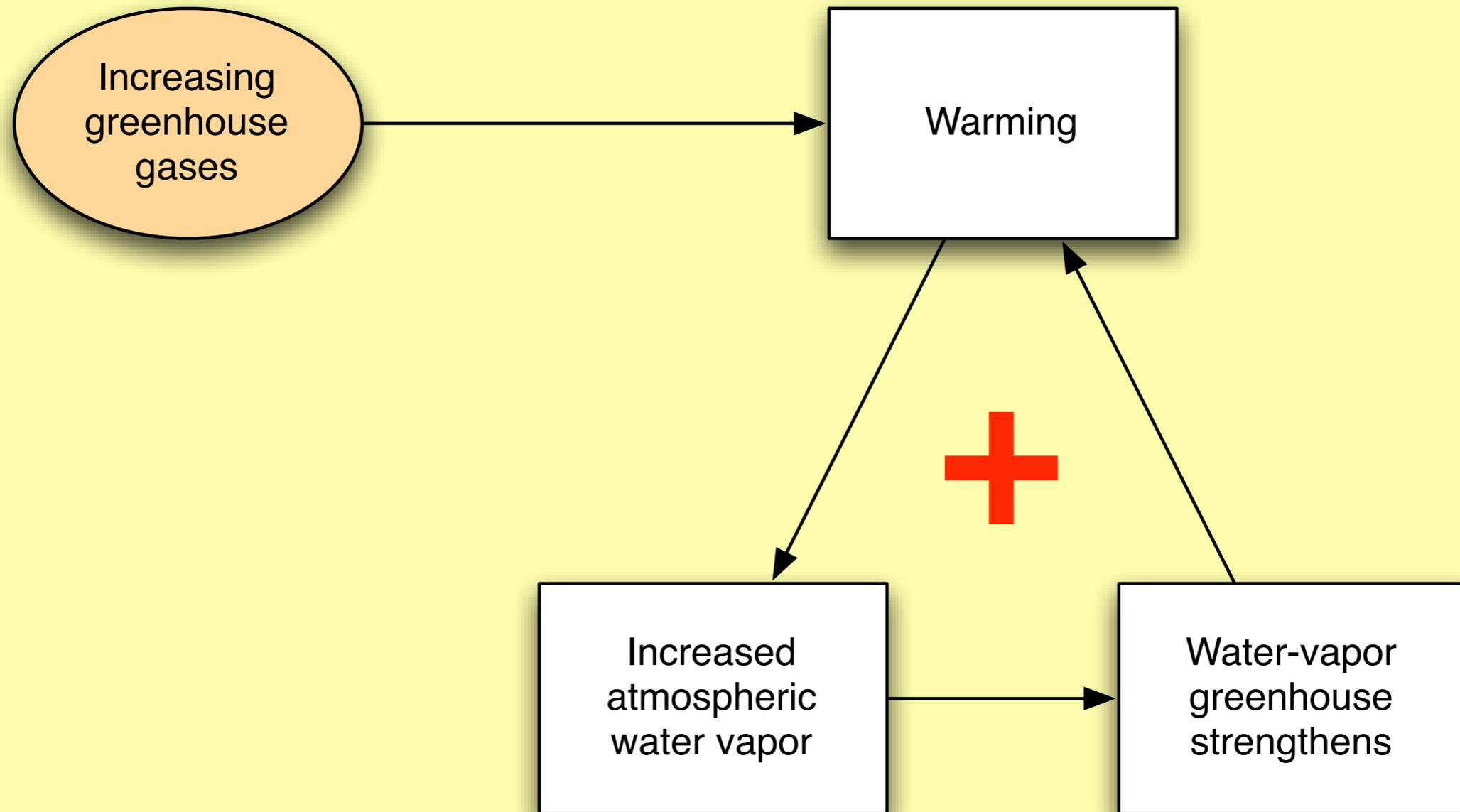
Global Mean Longwave Downward/Upward



Arctic Mean Longwave Downward/Upward

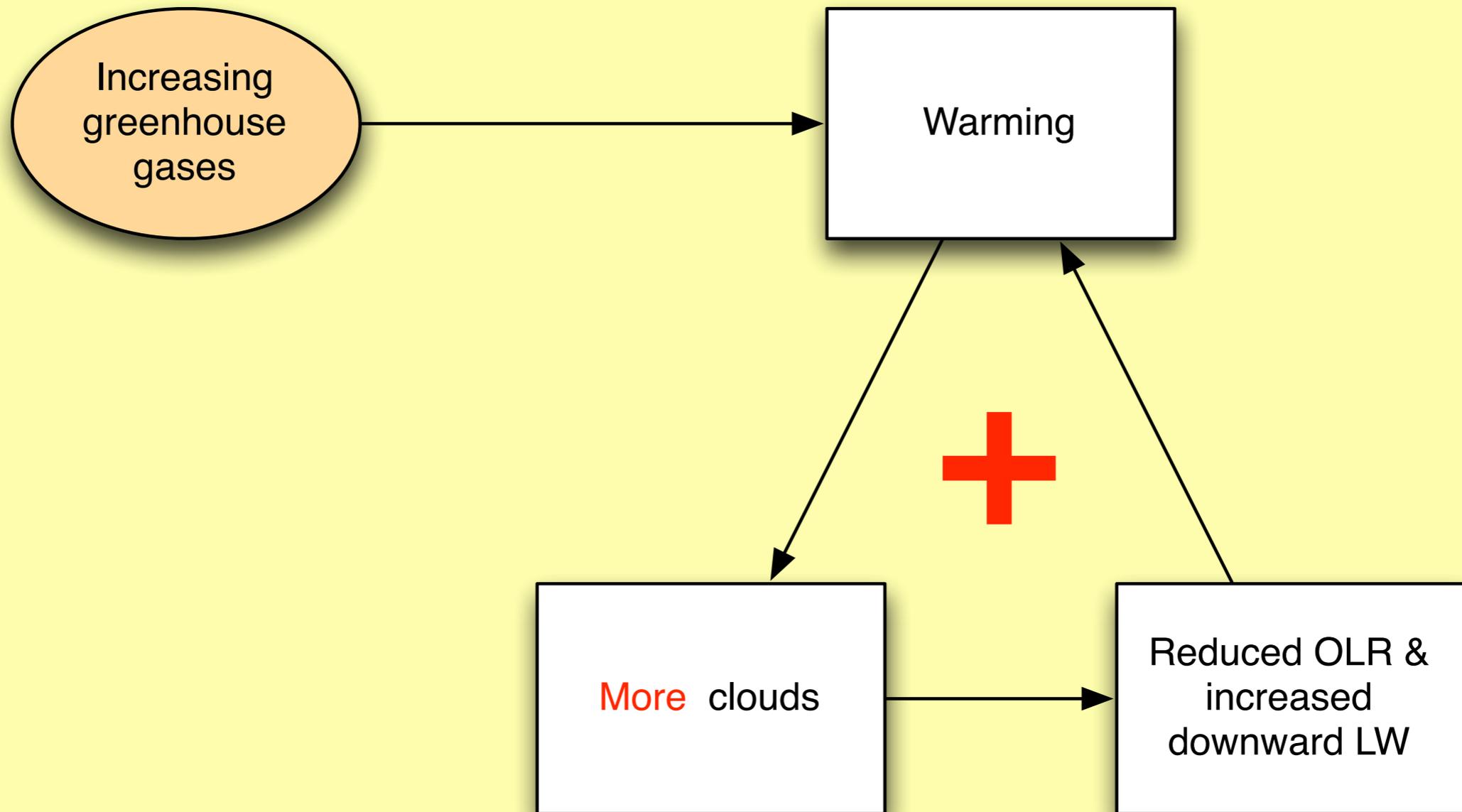


Water Vapor Feedback



As water vapor increases, precipitation and evaporation also increase.

Arctic Winter Cloud Feedback





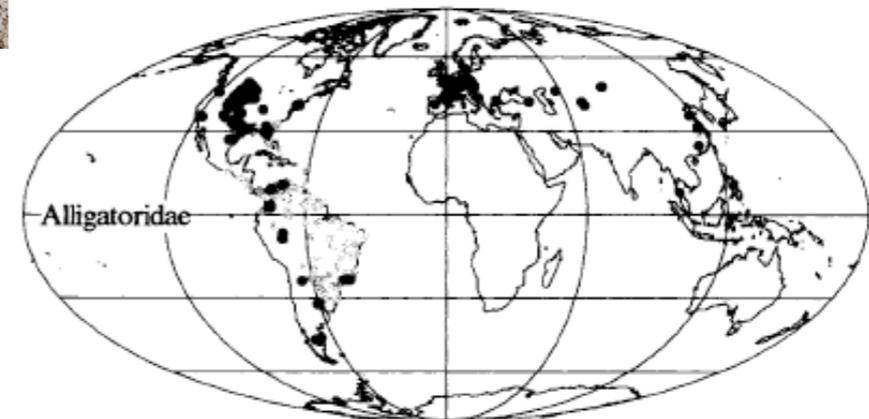
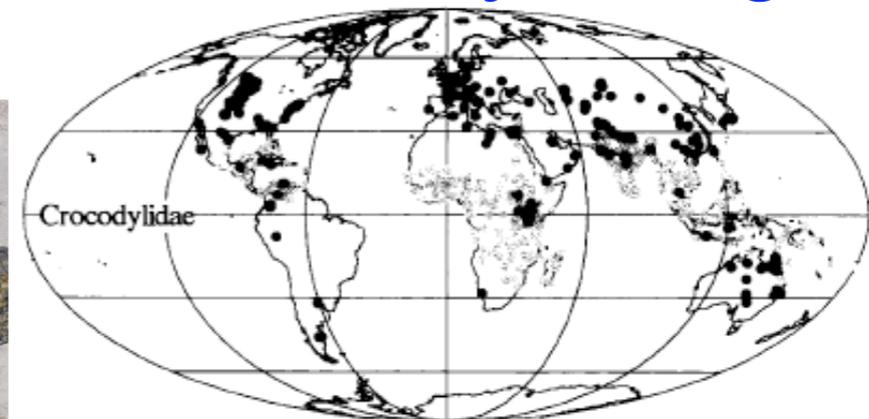
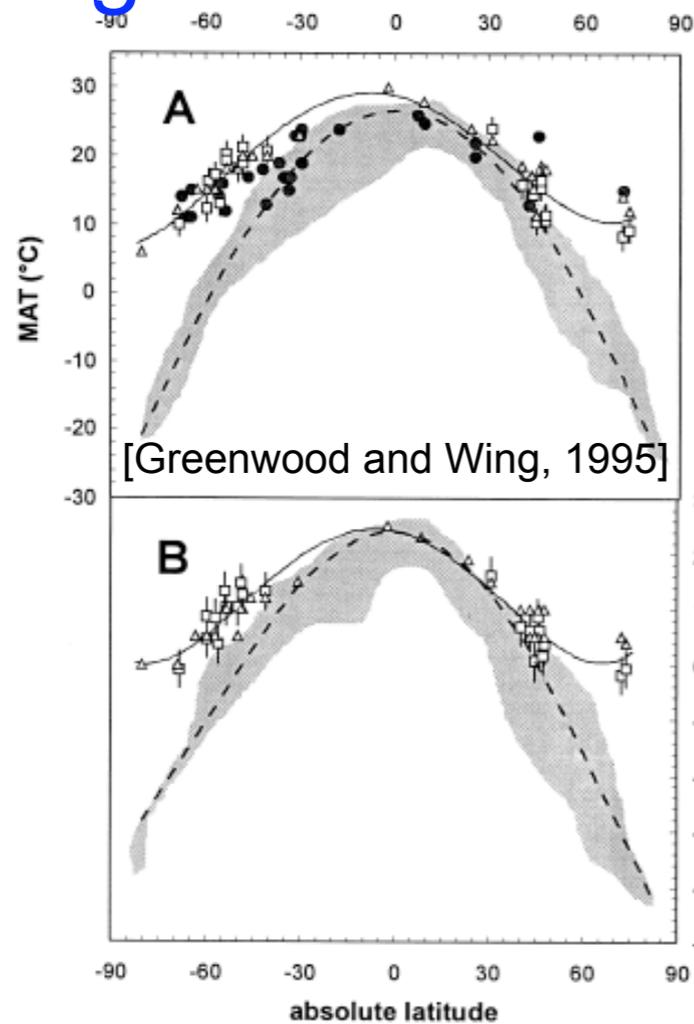
Ellesmere Island
80 °N
50 MYA
1200 ppmv CO₂



Eocene (50 MYA, ~1200 ppmv)

Cool tropics, warm high-latitudes

Crocodiles in Greenland, Palm trees in Wyoming!



- - Modern land temp.
- △ □ - Eocene SST; NLR&LMA, CLAMP

Crocodiles need: Mean annual $T > 14.2^{\circ}\text{C}$ & Cold month mean $> 5.5^{\circ}\text{C}$ [Markwick, 1998]

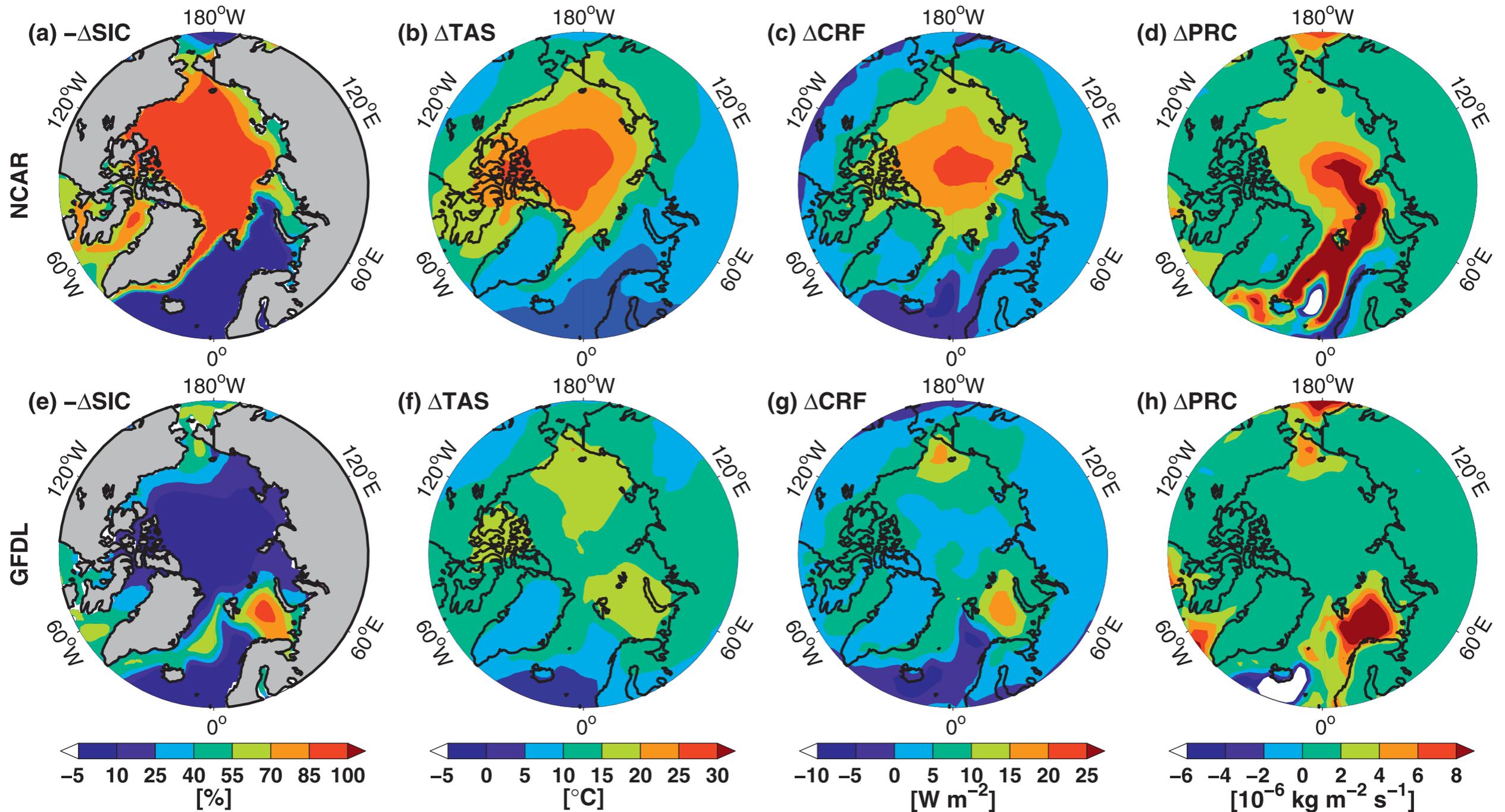
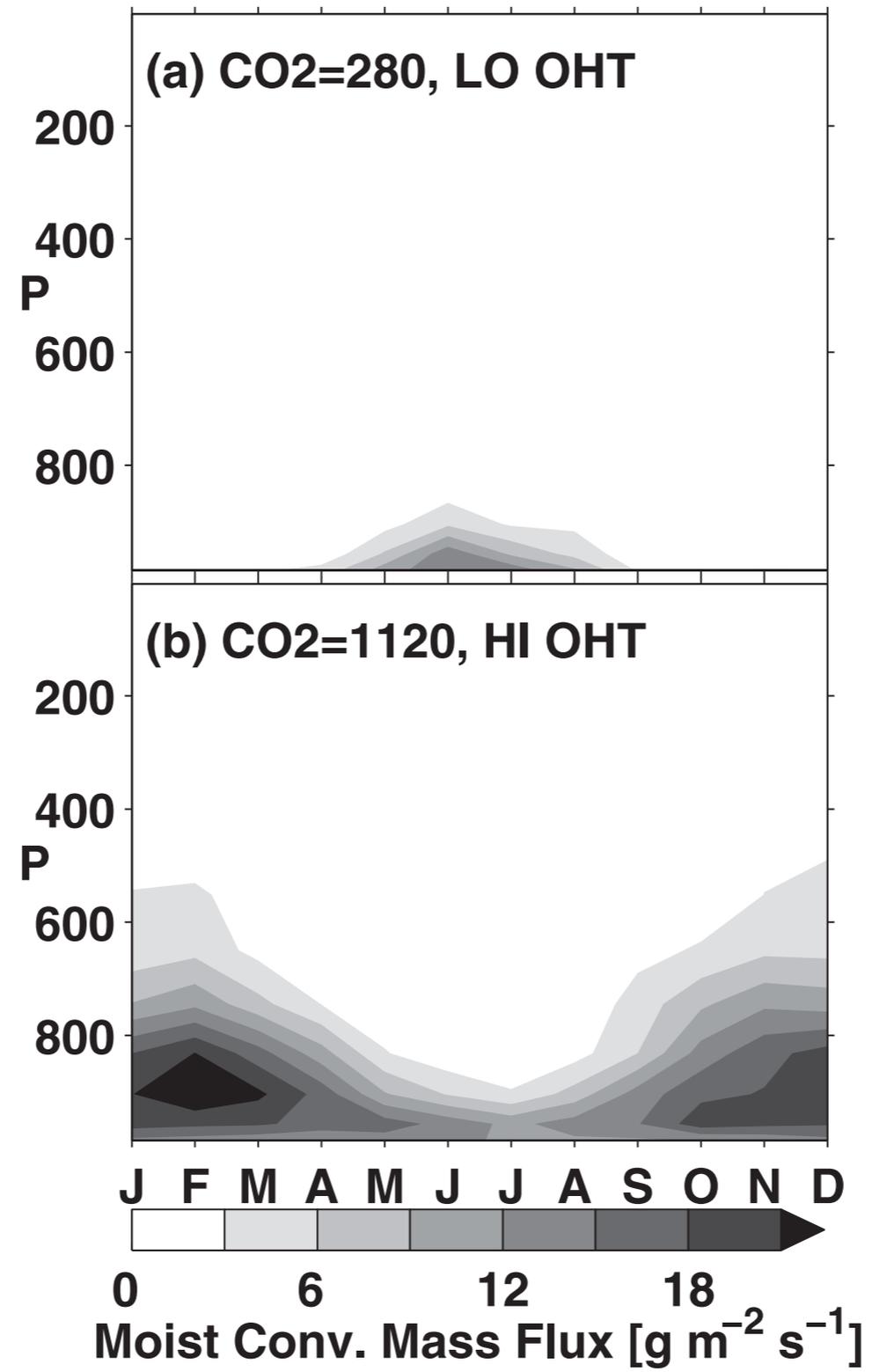


FIG. 1. Change in winter (November–February) Arctic climate over the course of the IPCCAR4 $4 \times \text{CO}_2$ experiment. The models are (a)–(d) NCAR CCSM3.0, which loses most Arctic winter sea ice and (e)–(h) GFDL CM2.0, which loses minimal winter Arctic winter sea ice. For each variable, the difference between the mean over the last 10 yr and the mean over the first 10 yr is plotted. (a),(e) $-\Delta\text{SIC}$, the negative of the change in sea ice concentration (100% means a complete loss of sea ice); (b),(f) ΔTAS , the change in surface air temperature; (c),(g) ΔCRF , the change in cloud radiative forcing; and (d),(h) ΔPRC , the change in convective precipitation rate.



“OHT” = ocean heat transport

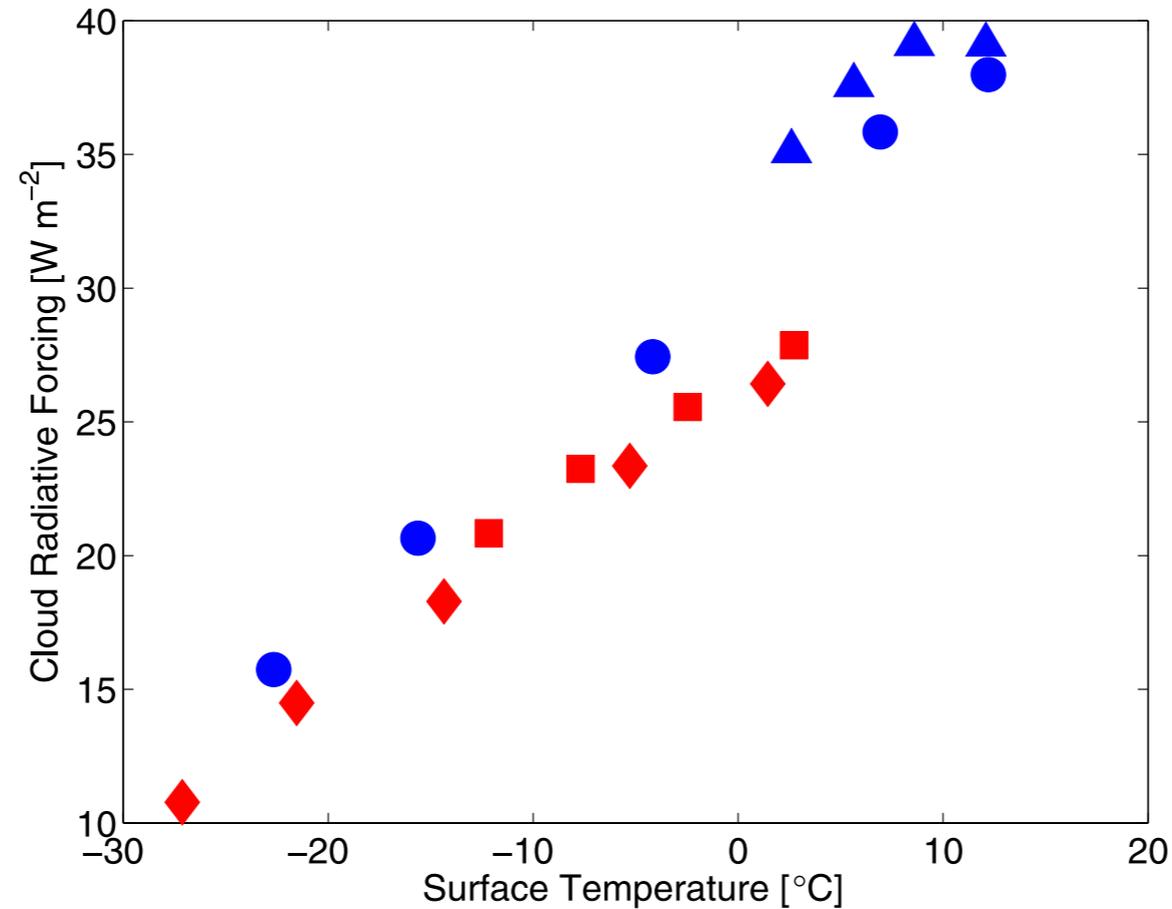
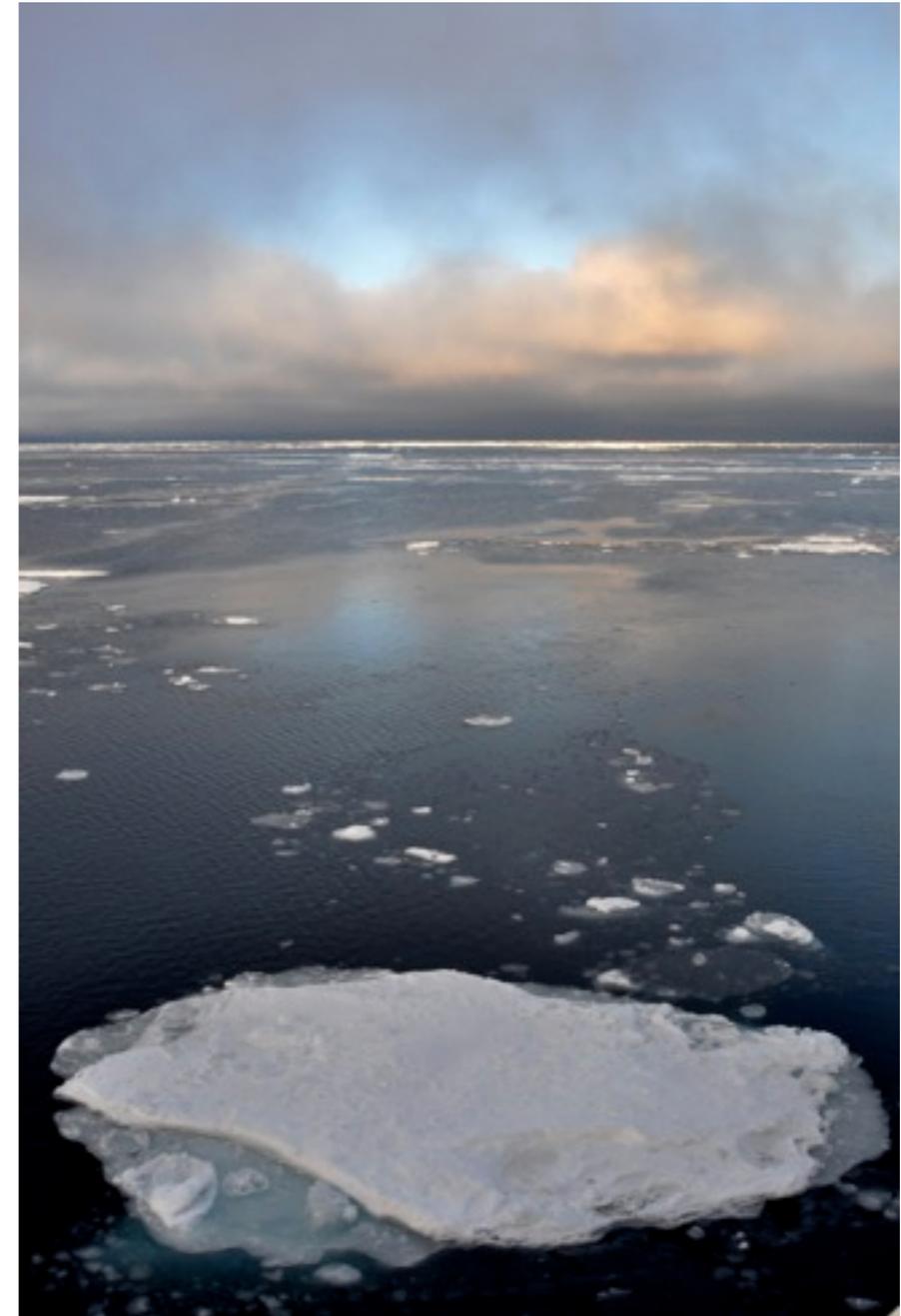


Figure 2. Winter (DJF) cloud radiative forcing averaged north of 60°N as a function of surface temperature averaged north of 60°N, which changes due to either changes in boundary conditions or CO₂. For each of the following cases, one datapoint represents one model run at a different CO₂ concentration: modern configuration land (red diamonds), Eocene configuration land (red squares), modern configuration ocean (blue circles), and Eocene configuration ocean (blue triangles).

Conclusions

- **Current models do not account for the observed rapid melt-back of the Arctic sea ice.**
- **The Arctic shows us climate change in “fast forward.”**
- **The positive surface albedo feedback is only one of several contributing factors.**
- **Cloud and water vapor feedbacks may also be very important.**



Schematic of accelerated warming in the Arctic-Boreal zone from effects in the physical climate system.

