

1 **Quantifying the observability of CO₂ flux uncertainty in atmospheric CO₂ records using**
2 **products from NASA's Carbon Monitoring Flux Pilot Project**

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43 **Abstract**

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45 NASA's Carbon Monitoring System (CMS) Flux Pilot Project (FPP) was designed to better understand
46 contemporary carbon fluxes by bringing together state-of-the art models with remote sensing datasets.
47 Here we report on simulations using NASA's Goddard Earth Observing System Model, version 5
48 (GEOS-5) which was used to evaluate the consistency of two different sets of observationally
49 constrained land and ocean fluxes with atmospheric CO₂ records. Despite the strong data constraint,
50 the average difference in annual terrestrial biosphere flux between the two land (NASA Ames CASA
51 and CASA-GFED) models is 1.7 Pg C for 2009-2010. Ocean models (NOBM and ECCO2-Darwin)
52 differ by 35% in their global estimates of carbon flux with particularly strong disagreement in high
53 latitudes. Based upon combinations of terrestrial and ocean fluxes, GEOS-5 reasonably simulated the
54 seasonal cycle observed at northern hemisphere surface sites and by the Greenhouse gases Observing
55 SATellite (GOSAT) while the model struggled to simulate the seasonal cycle at southern hemisphere
56 surface locations. Though GEOS-5 was able to reasonably reproduce the patterns of XCO₂ observed by
57 GOSAT, it struggled to reproduce these aspects of AIRS observations. Despite large differences
58 between land and ocean flux estimates, resulting differences in atmospheric mixing ratio were small,
59 typically less than 5 ppmv at the surface and 3 ppmv in the XCO₂ column. A statistical analysis based
60 on the variability of observations shows that flux differences of these magnitudes are difficult to
61 distinguish from natural variability, regardless of measurement platform.

62

63 1. Introduction

64

65 Major weaknesses still exist in our understanding of the processes that control atmospheric carbon
66 dioxide (CO₂) concentrations and as a result, our ability to simulate and predict changes in the Earth
67 system. While the magnitude of the global growth rate of atmospheric CO₂ is well constrained by
68 surface observations (e.g. Conway et al. [1994]), attributing its changes to specific processes remains a
69 challenge given current CO₂ observing capabilities. The Atmospheric Carbon Cycle Inversion
70 Intercomparison (TransCom 3) compared estimates of the carbon budget produced by inverse models
71 using a variety of atmospheric transport fields and CO₂ observations from a global network of surface
72 stations for the period 1992-1996; their results indicated the presence of a larger northern hemisphere
73 (NH) land carbon sink than represented in the a priori flux distribution assumed for the experiment,
74 though the results were strongly influenced by differences in transport fields [Gurney et al., 2002]. The
75 precise location, cause of, and variability of this missing carbon sink remain poorly understood despite
76 the insights provided by intercomparison studies. In addition to uncertainty in the processes governing
77 the contemporary carbon budget, there is evidence that natural land and ocean carbon sinks have
78 decreased over the course of the twentieth century resulting in an increase in the fraction of
79 anthropogenic emissions remaining in the atmosphere, but this is still controversial [Le Quéré et al.,
80 2009; Knorr, 2009].

81

82 Model estimates of land and ocean carbon flux are important because they are the only way to
83 understand the underlying processes governing carbon storage and exchange, facilitating an enhanced
84 understanding that may one day contribute to improved abilities to predict changes in the global carbon
85 budget. A number of attempts have been made to compare estimates of carbon flux from different land
86 models. Randerson et al. [2009] compared two biogeochemistry models using a common modeling
87 framework and demonstrated that global carbon sinks differed by a factor of 2 during the 1990s.

88 Schaefer et al. [2012] and Raczka et al [2013] compared estimates of gross primary production (GPP)
89 and carbon balance among terrestrial ecosystem models over North America and found large
90 differences. Schwalm et al. [2010] compared output from 22 terrestrial biosphere models with data
91 from flux towers in North America and found that models' ability to reproduce observed monthly net
92 ecosystem exchange was poor, though performance was better at forested sites than non-forested sites.
93 Most recently, Huntzinger et al [2012] compared 19 terrestrial biosphere models over North America
94 and found large differences in estimates of GPP (between 12.2 and 32.9 PgC yr⁻¹) and heterotrophic
95 respiration (5.6 to 13.2 PgC yr⁻¹) among models with smaller differences in net ecosystem production (-
96 0.7 to 2.2 PgC yr⁻¹), underscoring the continued uncertainty in land flux processes. Observational
97 estimates of net ecosystem exchange collected worldwide at tower stations provide valuable
98 information to constrain carbon fluxes at local scales (e.g. Lafleur et al. [2003]; Hollinger et al. [2004];
99 Winderlich et al. [2013]), but are difficult to translate to realistic global estimates because of the spatial
100 heterogeneity of vegetation and limited flux sampling locations.

101

102 While ocean carbon fluxes are considered to be reasonably well constrained by the observational
103 database compiled by Takahashi et al. [2002, 2009], certain regions are poorly sampled and models are
104 still needed to understand air-sea exchange processes and predict how they might evolve in the future.
105 As is the case among land models, wide disparity exists among model-derived ocean flux estimates.
106 Doney et al. [2004] showed that errors in model physical processes complicated efforts to compare
107 ocean biogeochemical fields among models while Najjar et al. [2007] documented large circulation
108 differences among 12 global ocean models which influenced export of dissolved organic matter,
109 particularly to the Southern Ocean. Popova et al. [2012] compared five ocean models in the arctic and
110 found that, while primary production was generally consistent among models, the models disagreed
111 over the relative importance of the processes governing production. Uncertainties in the ocean carbon
112 uptake are especially important because they are directly propagated into the inference of the global

113 terrestrial sink as used by the Global Carbon Project [Le Quere et al, 2013].

114

115 It is important to design an Earth observing system that will enable reduction of uncertainties in carbon
116 fluxes and enhance our ability to predict changes in the carbon-climate system. A number of
117 components of the carbon cycle are currently constrained by different types of observations. The
118 Moderate Resolution Imaging Spectroradiometer (MODIS) instruments aboard NASA's Terra and
119 Aqua satellites provide information about vegetation characteristics and fire. Information from the
120 Advanced Very High Resolution Radiometer (AVHRR) instruments aboard NOAA satellites provide a
121 longer term record of vegetation evolution. The Sea-viewing Wide Field-of-view Sensor (SeaWiFS),
122 Suomi National Polar-orbiting Partnership (Suomi NPP), and MODIS both provide information on
123 ocean color and productivity while MODIS and geostationary weather satellites observe the physical
124 ocean state. Weather satellites also enhance our understanding of the carbon cycle by providing
125 information on atmospheric circulation that are necessary for interpreting atmospheric CO₂
126 observations. Though these remote sensing products inform a number of model-based land and ocean
127 carbon flux estimates, substantial uncertainty remains as documented in the numerous studies cited
128 above.

129

130 Atmospheric CO₂ observations provide an important constraint on carbon fluxes, but are limited in
131 space and time. NOAA maintains a network of surface sampling site in remote locations worldwide
132 where trace gas measurements are conducted several times per week as well as a smaller number of
133 observatories and tower sites that provide continuous CO₂ data. Ground-based column CO₂
134 observations are currently collected at a small number of stations as part of the Total Carbon Column
135 Observing Network (TCCON). Chevallier et al. (2011) demonstrated the utility of column CO₂ data for
136 inferring regional carbon budgets for the first time using data from this sparse network. Several satellite
137 datasets are available from the Atmospheric Infrared Sounder (AIRS) and the Japanese Greenhouses

138 gases Observing SATellite (GOSAT), and, prior to May, 2010, the Scanning Imaging Absorption
139 Spectrometer for Atmospheric Cartography (SCIAMACHY). Satellite CO₂ observations have
140 provided valuable information for constraining surface fluxes, but their impact is limited by their
141 inability to observe in the presence of clouds and aerosols (GOSAT) and low sensitivity to near surface
142 CO₂ mixing ratios (AIRS).

143

144 NASA's Carbon Monitoring System (CMS) Flux Pilot Project (FPP) was designed to better understand
145 the observational constraints placed on carbon fluxes by bringing together state of the art models with a
146 wealth of remote sensing resources. "Bottom up" surface flux estimates were computed by two land
147 and two ocean models for the 2009-2010 period using a consistent set of meteorology input from
148 NASA's Modern-Era Retrospective Analysis for Research and Applications (MERRA; Rienecker et al.
149 [2011]). These results were propagated forward in the atmosphere using NASA's Goddard Earth
150 Observing System Model, version 5 (GEOS-5) constrained by MERRA analyzed meteorological fields.
151 Unlike other model intercomparison projects which span a much larger range of model variables, our
152 intention is not to assess the full range of flux uncertainty and variability; instead, this work brings
153 together a small group of model estimates of flux and atmospheric transport that are tightly constrained
154 by remote sensing datasets and draws upon the expertise of the creators of those datasets to assess flux
155 differences and their manifestation in atmospheric CO₂ records. Here, we report on the results of these
156 simulations with a focus on understanding how uncertainty in observationally constrained land and
157 ocean flux estimates propagates into atmospheric CO₂ and characterizing how differences in flux
158 processes might be observed in the atmosphere by existing platforms. This work is complimentary to
159 the application of "top-down" approaches used to infer the surface fluxes given satellite observations
160 [Liu et al, 2014]. Section 2 of this paper describes the models used in this work while section 3 of this
161 paper describes the comparison with different CO₂ datasets. Section 4 provides a summary and
162 conclusions drawn from this work.

163

164 **2. Model and Data Background**

165 **2.1. Land Biosphere Flux Estimates**

166 **2.1.1. CASA-GFED3**

167 The Carnegie-Ames-Stanford-Approach – Global Fire Emissions Database version 3 (CASA-GFED3)
168 derives from Potter et al. [1993], diverging in development since Randerson et al. [1996]. CASA is a
169 light use efficiency type model: net primary production (NPP) is expressed as the product of
170 photosynthetically active solar radiation, a light use efficiency parameter, scalars that capture
171 temperature and moisture limitations, and fractional absorption of solar radiation by the vegetation
172 canopy (FPAR). This latter variable is derived from satellite data.

173

174 Fire parameterization was incorporated into the model by van der Werf et al. [2004], producing CASA-
175 GFED, and the model has undergone several revisions (van der Werf et al. [2006, 2010]) leading to its
176 most recent version CASA-GFED3. Input data sets include air temperature, precipitation, incident solar
177 radiation, a soil classification map, and a number of satellite derived products including MODIS
178 vegetation classification, MODIS based burned area, and AVHRR FPAR.

179

180 CASA-GFED3 is run at monthly time steps with 0.5° spatial resolution. As part of the CMS FPP,
181 fluxes were computed using MERRA meteorology and FPAR derived from AVHRR NDVI (Tucker et
182 al. [2005]) according to the procedure of Los et al. [2000]. The original 8-km, biweekly AVHRR NDVI
183 was aggregated up to the monthly, $0.5^\circ \times 0.5^\circ$ grid by averaging. Model output includes NPP,
184 heterotrophic respiration (Rh), and fire emissions from forest, savanna, deforestation, peat, and
185 agriculture. For this project, fire emissions were disaggregated from monthly to daily using the MODIS
186 active fire products as described in Mu et al. [2011]. Monthly NPP and heterotrophic respiration were
187 disaggregated to 3-hour time intervals following Olsen and Randerson [2004].

188

189 **2.1.2. NASA Ames CASA**

190 The NASA Ames CASA model (hereafter referred to as Ames CASA) relies on satellite observations
191 of vegetation cover from MODIS as time-series inputs to estimate monthly carbon fluxes from
192 terrestrial ecosystems worldwide. All model algorithms, parameter settings, and land cover data sets
193 used in Ames CASA for CMS flux computations have been documented by Potter et al. [2007].

194

195 For the CMS FPP computations of net biosphere fluxes of carbon to the atmosphere, the Ames CASA
196 version documented in Potter et al. [2007] has been modified to use global 0.5° (latitude/longitude)
197 MODIS Enhanced Vegetation Index (EVI) input data (for the years 2000-2010) generated by
198 aggregating monthly 0.05° (~6 km) values. In addition, in cropland areas, 40% of annual NPP carbon
199 is removed each year from the litter decomposition flux pathways and diverted into harvested food
200 products. This is assumed to be re-emitted as a consistent monthly flux (1/12 of the annual cropland
201 harvest carbon total) with a weighted spatial distribution corresponding to the maps of cropland harvest
202 CO_2 emissions developed by Ciais et al. [2007]. Net biosphere fluxes for both CASA-GFED3 and
203 Ames CASA were disaggregated to 3-hour time intervals following Olsen and Randerson [2004]. In
204 contrast to the CASA-GFED3 fluxes, biomass burning is not included in the Ames CASA fluxes
205 computed as part of the CMS FPP. CASA-GFED3 does not include the crop redistribution incorporated
206 in Ames CASA flux estimates.

207

208 The Ames CASA fluxes produced as part of the CMS FPP differ from a more recent version of Ames
209 CASA described in Potter et al. [2012] both in the use of different meteorological driver data and in the
210 methodology of process representation.

211

212 **2.1.3. Differences between Land Flux Estimates**

213 Figure 1 shows time series of monthly global flux totals computed by the two land models. Ames
214 CASA estimates a weaker land sink (driven by NH summer) and a stronger source during NH winter
215 compared to CASA-GFED3. These differences result in substantially different annual flux estimates
216 for both years; during 2009 (2010), Ames CASA estimates that the terrestrial biosphere is a 2.5 (2.1)
217 Pg C source while CASA-GFED3 calculates a smaller 0.34 (1.2) Pg C source. The average difference
218 in annual land flux between the two models is 1.7 Pg C. For reference, the uncertainty in land flux
219 between these two models is nearly 20% of the annual mean fossil fuel emissions for the 2009 to 2010
220 period (9 Pg C; Boden et al. [2011]). GFED3 fluxes for 2009 (2010) indicate that fires are a global 2.1
221 (2.9) Pg C source of carbon to the atmosphere while Ames CASA fluxes considered here do not
222 include the effects of fire. In both models, the land biosphere is a source of carbon the atmosphere
223 rather than a sink as indicated by multi-model inversion studies (e.g. Gurney et al. [2002]). Using
224 observed atmospheric CO₂ increases, fossil fuel emission inventories, and ocean models, Le Quéré et
225 al. [2013] estimate the magnitude of the land carbon sink as 3.3 (2.55) Pg C during 2009 (2010). This
226 may be due to the fact that neither model includes several processes thought to be important in
227 explaining the global land sink including CO₂ fertilization (e.g. Bellassen et al. [2011]; Vanuytrecht et
228 al. [2011]; Piao et al. [2013]; Los [2013]), nitrogen deposition (e.g. Esser et al. [2011]; Bala et al.
229 [2013]; Fleischer et al. [2013]; Gerber et al. [2013]), and land use history (e.g. Sentman et al. [2011]).
230 In addition to differences in the magnitude of the land biosphere flux estimates, the Ames CASA and
231 CASA-GFED3 models indicate differences in the phasing of the global seasonal cycle in terrestrial
232 carbon flux to the atmosphere. Ames CASA diagnoses an earlier pattern of drawdown during NH
233 spring and an earlier transition from sink to source during NH autumn in 2009.

234

235 Figures 2 shows the geographic distribution of land fluxes computed by CASA-GFED3 and Ames
236 CASA for four months during 2009. During January, CASA-GFED3 indicates greater drawdown of
237 CO₂ over the southern hemisphere (SH) regions of South America and Africa than does Ames CASA.

238 Fire emissions in equatorial Africa and Australia are also evident in CASA-GFED3 while absent in
239 Ames CASA. In the NH, Ames CASA estimates greater release of CO₂ to the atmosphere along the
240 east coasts of the United States and China. In April, Ames CASA fluxes are generally weak in most
241 locations as the model transitions between winter and summer seasons; areas of western Europe have
242 transitioned from a source in January to a sink of CO₂ in April. In contrast, CASA-GFED3 patterns are
243 generally similar in April and January but fluxes from the land to the atmosphere are slightly larger at
244 most NH locations. During July, CASA-GFED3 estimates greater uptake of CO₂ at most NH locations,
245 particularly at high latitudes, than does Ames CASA. This contrast is also evident over equatorial
246 South America while in mid-latitude regions of the continent, CASA-GFED3 indicates greater release
247 of CO₂ to the atmosphere. The models also differ strongly in their assessment of fluxes over India and
248 Southeast Asia during July with CASA-GFED3 indicating a substantial source and Ames CASA
249 indicating a sink. During October, Ames CASA fluxes are generally much weaker than CASA-GFED3
250 fluxes. Strong fire activity over South America is evident in CASA-GFED3 during October and not
251 included in Ames CASA flux estimates. Differences between the spatial distributions of Ames CASA
252 and CASA-GFED3 fluxes for 2010 (not shown) are qualitatively similar though 2010 was
253 characterized by larger fire emissions in CASA-GFED3.

254

255 **2.2. Ocean Flux Estimates**

256 **2.2.1. NOBM**

257 Global ocean carbon dynamics are simulated by the NASA Ocean Biogeochemical Model (NOBM). It
258 is a three-dimensional representation of coupled circulation, biogeochemical, and radiative processes in
259 the global oceans (Gregg et al. [2003]; Gregg and Casey [2007]). The biogeochemical processes model
260 contains 4 phytoplankton groups, 4 nutrient groups, a single herbivore group, and 3 detrital pools. The
261 phytoplankton groups differ in maximum growth rates, sinking rates, nutrient requirements, and optical
262 properties. Three detrital pools provide for storage of organic material, sinking, and eventual

263 remineralization back to usable nutrients. Radiative transfer calculations provide the underwater
264 irradiance fields necessary to drive growth of the phytoplankton groups, and interact with the heat
265 budget. Carbon cycling involves dissolved organic carbon (DOC) and dissolved inorganic carbon
266 (Gregg et al., 2013). DOC has sources from phytoplankton, herbivores, and carbon detritus, and a sink
267 to DIC. DIC has sources from phytoplankton, herbivores, carbon detritus, and DOC, and
268 communicates with the atmosphere, which can be either a source or sink. The ecosystem sink for DIC
269 is phytoplankton, through photosynthesis. This represents the biological pump portion of the carbon
270 dynamics. The solubility pump portion is represented by the interactions among temperature, alkalinity
271 (parameterized as a function of salinity), silica, and phosphate (parameterized as a function of nitrate).
272 The alkalinity/salinity parameterization utilizes the spatial variability of salinity in the model adjusted
273 to mean alkalinity from the Ocean Model Intercomparison Project (OCMIP; Orr et al. [2001]). The
274 calculations for the solubility pump follow the standards set by the OCMIP.

275

276 NOBM's domain spans from -84° to 72° latitude in increments of 1.25° longitude by $2/3^{\circ}$ latitude,
277 including only open ocean areas where bottom depth is greater than 200m. The model contains 14
278 vertical layers, in quasi-isopycnal coordinates and is driven by MERRA monthly mean wind stress, sea
279 surface temperature, and shortwave radiation fields. MODIS-Aqua chlorophyll data were assimilated to
280 represent the sum of all phytoplankton components in the model. As part of the CMS FPP, daily
281 NOBM ocean $p\text{CO}_2$ fields were used as input into GEOS-5 and ocean-atmosphere CO_2 fluxes were
282 computed within GEOS-5 using the GEOS-5 atmospheric CO_2 mixing ratios and winds.

283

284 **2.2.2. ECCO2-Darwin**

285 The ECCO2-Darwin Ocean Carbon Cycle Model is based on a global, eddying, ocean and sea ice
286 configuration of the Massachusetts Institute of Technology general circulation model (MITgcm;
287 Marshall et al. [1997a, 1997b]) and on results from two separate projects: the Estimating the

288 Circulation and Climate of the Ocean, Phase II (ECCO2) Project, which provides a data-constrained
289 estimate of the time-evolving physical ocean state, and the Darwin Project, which provides time-
290 evolving ocean ecosystem variables. Together, ECCO2 and Darwin provide a time-evolving physical
291 and biological environment for carbon biogeochemistry, which is used to compute surface fluxes of
292 carbon at high spatial and temporal resolution.

293

294 The ECCO2 model configuration is a cube-sphere grid [Adcroft et al., 2004] with 18-km horizontal
295 grid spacing and 50 vertical levels [Menemenlis et al., 2005a; 2008]. The ECCO2 model configuration
296 includes a dynamic/thermodynamic sea ice model (Losch et al. [2010]; Heimbach et al. [2010]). In a
297 first step, the ECCO2 model configuration was adjusted using a low-dimensional (Green's functions)
298 estimation approach [Menemenlis et al., 2005b]. In a second step, the method of Lagrange multipliers
299 (adjoint method) was used to adjust initial and time-evolving surface boundary conditions [Wunsch and
300 Heimbach, 2007]. Data constraints include sea level anomaly from Jason-1 and OSTM, sea surface
301 temperature from AMSR-E, and temperature and salinity profiles from Argo. This adjoint-based
302 ECCO2 solution is used to drive the Darwin ecosystem model.

303

304 The Darwin Project is an initiative to advance the development and application of novel models of
305 marine microbial communities, identifying the relationships of individuals and communities to their
306 environment, connecting cellular-scale processes to global microbial community structure (Follows et
307 al. [2007, 2011]; Dutkiewicz et al. [2009]). The particular configuration used for the CMS FPP
308 includes five phytoplankton functional types (choices based on results from previous versions of the
309 model) and two zooplankton types. The carbon cycle is explicitly included in this configuration, along
310 with those of nitrogen, phosphorus, iron, silica, oxygen, and alkalinity. The carbonate chemistry
311 follows the simplified model proposed by Follows et al. [2006] and air-sea CO₂ exchange is
312 parameterized according to Wanninkhof [1992].

313

314 ECCO2-Darwin fluxes produced as part of the CMS FPP are described in more detail in Brix et al.
315 [2014]. GEOS-5 simulations presented here use version 2.1 fluxes. Since the CMS FPP, ECCO2-
316 Darwin fluxes have undergone further development including modification of the piston velocity
317 parameterization.

318

319 **2.2.3. Differences between Ocean Flux Estimates**

320 NOBM ocean fluxes contain weaker seasonal variations than those produced by the ECCO2-Darwin
321 model (Figure 1). Both models estimate comparable global fluxes for much of the year but during June,
322 July, and August (JJA), ECCO2-Darwin fluxes indicate a weaker ocean sink (or slight source) relative
323 to NOBM; as a result, the 2009 (2010) annual ECCO2-Darwin ocean sink is 2.4 (2.6) Pg C compared
324 to 3.7 (4.0) Pg C from NOBM, a difference of 36% (35%). The Global Carbon Budget [Le Quere et al.,
325 2013] estimates a global ocean flux of 2.57 (2.55) Pg C yr⁻¹ for 2009 (2010). NOBM ocean fluxes
326 indicate a stronger ocean sink than the GCB estimate while ECCO2-Darwin estimates differ by less
327 than 10% from GCB in both years.

328

329 Figure 3 shows geographic distributions of ocean fluxes computed by NOBM and ECCO2-Darwin for
330 four months in 2009. During January, differences between model estimates of flux are largest in high
331 latitude oceans. NOBM estimates a greater Atlantic Ocean sink north of 30° while ECCO2-Darwin
332 estimates a much stronger ocean sink in the southern ocean than is evident in NOBM. April
333 distributions are generally similar to January, though a decrease in the northern NOBM sink and
334 increase in the ECCO2-Darwin sink relative to January result in spatially inhomogeneous differences
335 north of 30°. During July, ocean flux distributions differ substantially from January; the northern sink
336 in NOBM has weakened and is now evident only in the north Atlantic basin. ECCO2-Darwin's strong
337 southern ocean sink has also weakened while the NOBM southern ocean sink has become stronger.

338 During October, the NOBM distribution of fluxes is dominated by a strong sink throughout much of
339 the southern hemisphere oceans. ECCO2-Darwin indicates stronger sinks in high latitude regions of
340 both hemispheres, but with less areal extent relative to NOBM. Ocean models also differ in their
341 estimation of coastal fluxes, particularly off the Atlantic coasts of Africa and North America and the
342 Pacific coast of South America.

343

344 In regions below 30°S, ECCO2-Darwin estimates an ocean sink 27% (23%) greater than NOBM during
345 2009 (2010). In addition to differences in magnitude of the sink in this region which includes the
346 Southern Ocean, the seasonal cycle of ocean fluxes is nearly reversed; NOBM estimates peak
347 drawdown in the sink during SH winter months, while ECCO2-Darwin indicates that the sink is at its
348 weakest during these months. In SH Summer, ECCO2-Darwin simulates a maximum in CO₂
349 drawdown in the region while NOBM simulates a minimum. NOBM, whose domain only extends to
350 72°N, estimates a sink several times weaker than ECCO2 in NH high latitude oceans. The largest
351 disparities between ocean flux models occur in regions that are rarely observed by ships or are
352 observed only during certain months as shown in Takahashi et al. [2009].

353

354 **2.3. GEOS-5 Earth System Model**

355 The GEOS-5 Atmospheric General Circulation Model (AGCM) has been developed as a flexible tool to
356 represent the atmosphere on a variety of temporal and spatial scales. It is a central component of the
357 GEOS-5 atmospheric data assimilation system [Rienecker et al., 2008], where it is used with half-
358 degree spatial resolution for meteorological analysis and forecasting [Zhu and Gelaro, 2008] including
359 the production of MERRA which spans the period from 1979 to present [Reinecker et al., 2011]. It has
360 also been developed as a tool for studying atmospheric composition and climate. Ott et al. [2010] used
361 GEOS-5 to examine the impact of Indonesian biomass burning aerosols on atmospheric circulation and
362 Ott et al. [2011] examined the impact of uncertainty in GEOS-5 convection on global carbon monoxide

363 distributions. Model output has also been used by Wang et al. [2009] to compute CO:CO₂ correlations
364 and their impact on inverse modeling.

365

366 The AGCM combines the finite volume dynamical core described in Lin [2004] with the GEOS-5
367 column physics package, summarized in Rienecker et al. [2008]. The model domain extends from the
368 surface to 0.01 hPa and uses 72 hybrid layers that transition from terrain-following near the surface to
369 pure pressure levels above 180 hPa. In this study, the horizontal resolution is 1° x 1.25° (latitude by
370 longitude) and the time step is 30 minutes for physical computations, with more frequent computations
371 of resolved-scale transport in the dynamical core. Trace gases are transported on-line in GEOS-5 using
372 the Lin [2004] dynamical core for resolved scales; turbulent mixing of CO₂ is performed in the same
373 way as for moisture (using the Lock et al. [2000] boundary-layer turbulence module); and using the
374 Relaxed-Arakawa Schubert convective scheme [Moorthi and Suarez, 1992] to represent convective
375 transport. In the present simulations, transport is constrained with MERRA reanalysis fields to ensure
376 consistency with observed meteorology.

377

378 Land biosphere, biomass burning, fossil fuel, and ocean CO₂ fluxes are prescribed in GEOS-5. For the
379 CMS FPP simulations, GEOS-5 was configured to simulate the emission and transport of several
380 different CO₂ tracers representing differing combinations of land and ocean fluxes described in Table 1.
381 In addition to the land and ocean biosphere fluxes provided as part of the CMS FPP, CO₂ emissions
382 from fossil fuels are taken from the Department of Energy's Carbon Dioxide Information Analysis
383 Center (CDIAC; Boden et al. [2011]). Prior to the target 2009-2010 CMS period, CO₂ tracers were
384 spun up from 2000-2008 beginning with a uniform initial condition of 350 ppmv to ensure realistic
385 atmospheric distributions. During the spinup period, land biosphere and biomass burning fluxes from
386 an earlier version of the CASA-GFED model were used in combination with ocean and fossil fuel
387 fluxes from the TransCom Continuous Experiment [Law et al., 2008]. Simulated CO₂ mixing ratios for

388 December 2008 were calculated at the locations of NOAA Earth Science Research Laboratory (ESRL)
389 marine boundary layer stations [Novelli et al., 1992] and compared with observations; on the basis of
390 this comparison a uniform global offset was subtracted from the simulated CO₂ fields to ensure that
391 global average surface CO₂ concentration was representative of atmospheric conditions at the
392 beginning of the CMS period.

393

394 **3. Comparison with CO₂ Observational Records**

395 **3.1. Comparison with surface CO₂ in situ observations**

396 NOAA's Earth System Research Laboratory (ESRL) Carbon Cycle Greenhouse Gases Group (CCGG)
397 analyzes samples taken weekly at an international cooperative network of surface observing stations
398 (Tans et al., 1990). These data have been used to understand both long term changes in and interannual
399 variability of natural carbon sinks (e.g. Ballantyne et al. [2012]; Conway et al. [1994]). Sites are
400 typically located in remote locations (Figure 4) so that observations represent the background surface
401 CO₂ mixing ratios rather than local source and sink influences.

402

403 For comparison to surface stations, the GEOS-5 grid cell containing each station location was sampled
404 at the time that an observation was collected and simulated CO₂ mixing ratios then vertically
405 interpolated to the altitude of the observing station creating model 'pseudo data'. Observations and
406 pseudo-data are averaged over the course of a day (when multiple observations are present) and daily
407 simulated and observed CO₂ mixing ratios at Mauna Loa, Ny Alesund, and Palmer Station (indicated
408 on Figure 4) are shown in Figure 5 for the flux scenarios described in Table 1. One of the primary
409 features evident in this comparison is difference in annual atmospheric growth rate between the flux
410 scenarios. In the beginning of 2009, all scenarios in the GEOS-5 simulations begin from the same CO₂
411 mixing ratios, but over time differences in the magnitude of the combined land and ocean carbon flux
412 cause the spread in the ensemble of simulations to grow. While this spread in simulations caused by

413 differing growth rates happens at all stations, it is most evident at Mauna Loa and Palmer Station
414 because of the smaller amplitude of the seasonal cycle at these locations. We calculate growth rates for
415 each flux scenario by first calculating the monthly mean of surface CO₂ mixing ratio at all CCGG
416 marine boundary layer (MBL) sites during 2009 and 2010. For each month, the 2009 MBL monthly
417 mean is subtracted from the corresponding 2010 value and these differences are averaged over all
418 calendar months to estimate the annual mean growth rate shown in Table 1. Using the same method to
419 calculate the annual mean growth rate from observations yields a value of 2.4 ppmv. Flux scenario CG-
420 NO, which combines NOBM and CASA-GFED fluxes, provides the most favorable comparison with
421 the observed growth rate. Both the Ames CASA and ECCO-2 fluxes produce weaker natural sinks than
422 flux scenario CG-NO resulting in greater rates of accumulation of CO₂ in the atmosphere for scenarios
423 AC-NO and CG-ED relative to scenario CG-NO.

424

425 In order to examine the ability of the simulations to capture observed spatial and temporal variations in
426 CO₂, we remove the difference in growth rate by subtracting the trends from observations and
427 simulations (Figure 5, right columns). At Mauna Loa, the de-trended time series shows that land
428 biosphere flux differences have a greater impact on simulated mixing ratios than do ocean flux
429 differences. None of the flux scenarios succeeds in reproducing the magnitude of the seasonal cycle
430 observed at Mauna Loa, likely because the NH land sinks in both estimates are too weak or because of
431 deficiencies in model transport. At Ny-Alesund, all model simulations reproduce the seasonal cycle
432 reasonably well though simulations tend to overestimate CO₂, particularly during spring. Differences in
433 seasonal cycle between the Ames CASA and CASA-GFED fluxes are small, but evident at this station
434 in spring and summer months. Ocean flux differences manifest as much smaller mixing ratio
435 differences despite the fact that only one model, ECCO2-Darwin, produces fluxes at this high northern
436 latitude region. At Palmer Station Antarctica, ocean flux differences are larger than at any other station
437 due to the disparity between model estimates of Southern Ocean flux. NOBM overestimates the

438 magnitude of the observed, weak seasonal cycle by 0.5 ppmv while seasonal variations in the ECCO2-
439 Darwin simulation are much too strong resulting in a 5 ppmv overestimate of the seasonal cycle
440 amplitude. The similarity of the CG-NO and AC-NO simulations and large differences between the
441 CG-NO and CG-ED simulations suggest that the seasonal cycle of nearby ocean fluxes more strongly
442 influences the simulation of CO₂ at high latitude SH stations than does the seasonal cycle of land
443 fluxes.

444

445 Figure 6 compares observed and simulated (assuming flux combination CG-NO) monthly mean CO₂
446 mixing ratios at all NOAA ESRL stations collecting substantial amounts of data during the 2009-2010
447 period. Observations at NH stations comprise the bulk of the dataset and show a strong seasonal cycle
448 due to the influence of the land biosphere. The CG-NO simulation also shows a strong seasonal cycle
449 for most stations north of 30°N. At these locations, the model tends to overestimate CO₂, particularly
450 during NH spring and summer, because the CASA-GFED3 land sink is too weak. Comparisons with
451 the Global Carbon Budget presented by Le Quere et al. [2013] indicate an overestimate of global fluxes
452 by the GC-NO flux combination of approximately 3 Pg C which translates into an average, global
453 overestimate in the atmospheric CO₂ of 1.4 ppmv assuming that the excess CO₂ were instantly diffused
454 throughout the mass of the atmosphere. At stations north of 60°N, annual mean CO₂ is, on average,
455 overestimated by 2.3 ppmv during 2010 while stations between the equator and 60°N are overestimated
456 by an average of 1.3 ppmv. CO₂ at northern high latitude stations is overestimated by an average of 4
457 ppmv during spring months and 3 ppmv during summer while NH stations south of 60°N are
458 overestimated by 1.7 ppmv during spring and 1.9 ppmv during summer. The difference in the
459 magnitude of overestimate between NH high and mid-latitude stations may indicate that the CASA-
460 GFED3 land sink underestimate is concentrated in this area or that vertical transport errors are greater
461 at NH high latitudes, resulting in excessive concentration of CO₂ near the surface. In the SH, model
462 errors average 1.7 ppmv during 2010 and show much weaker seasonality.

463

464 In Figure 7, we evaluate the ability of GEOS-5 simulations to reproduce the seasonal cycle in CO₂
465 observed at ESRL stations during 2010. To calculate seasonal cycle errors, monthly means were
466 calculated from observations and model output after de-trending to remove the annual growth rate. We
467 define the amplitude of the seasonal cycle as the difference between the maximum and minimum
468 monthly values. Errors in the months during which minima and maxima in the seasonal cycle occur are
469 also calculated for each station. Means of the amplitude and phasing errors are then calculated over 10
470 degree latitude bands and only include stations for which data is available in every calendar month
471 during 2010. At NH mid- and high latitude stations, performance is similar for all model flux
472 combinations. Simulations overestimate the magnitude of the seasonal cycle by a few ppmv on average,
473 generally succeed in estimating the summer month during which the minimum should occur, but have
474 more difficulty in estimating the timing of maximum CO₂ ratios in the winter. While errors in the
475 amplitude of the seasonal cycle are small at tropical stations, the model given any flux combination
476 tends to struggle reproducing the observed timing of minima and maxima at SH tropical stations. This
477 is likely due to a combination of the weak seasonal cycle in CO₂ at these locations, errors in the
478 transport of CO₂ from NH locations, and uncertainty in land fluxes in this region that contains dense
479 vegetation. All simulations considered in this study fail to adequately simulate the seasonal cycle at SH
480 high latitude stations. Amplitude errors of several ppmv are often larger than the observed seasonal
481 cycle magnitude. In these locations, ECCO2-Darwin fluxes, which indicate a stronger Southern Ocean
482 sink with greater seasonality relative to NOBM, degrade model performance in terms of both seasonal
483 cycle amplitude and the timing of the maximum in surface CO₂ mixing ratio.

484

485 Despite the substantial flux differences among the land and ocean models shown in Figures 1-3,
486 differences in atmospheric CO₂ mixing ratio at remote surface stations tend to be quite subtle as
487 evident in the time series comparisons in Figure 5. To quantitatively determine how atmospheric CO₂

488 observations might be used to discern between these realistic, observationally constrained flux
489 estimates, we follow the method of Huntzinger et al. [2011] who examined the utility of continuous
490 atmospheric flux measurements at a set of tower locations for constraining biospheric flux variability.
491 In order to test the statistical significance of the difference between simulated pseudo-observations, we
492 apply a Chi-square test of variance based on the number of observations per month, the mean squared
493 difference between pairs of pseudo-observations using different flux estimates, and the variance of
494 model-data mismatch [Huntzinger et al., 2011]. Model-data mismatch, an estimate of how closely a
495 model could reproduce an observation, is estimated in this study by estimating the variance of
496 observation residuals around a smooth curve fit to the observations at each site, a method used by
497 Bousquet et al. [1999] and Gurney et al. [2002]. It is important to note that expected model-data
498 mismatch is calculated solely based on variability in the observations. We calculate mean squared
499 difference between simulated CO₂ pairs by first removing the difference in annual growth rate as
500 shown in Figure 5. Qualitatively, this method allows us to assess how well the small atmospheric signal
501 caused by a difference in assumed flux might be distinguished from a background of strong, natural
502 variability.

503

504 Figure 8 shows the difference in monthly mean CO₂ mixing ratio at NOAA ESRL stations resulting
505 from underlying land and ocean flux differences. Differences which are not significantly larger than the
506 model-data mismatch variance at a significance level of 0.05 are indicated by diagonal lines. The
507 largest differences due to land flux are evident at stations in the NH mid- and high latitudes during
508 spring and early summer (2-4 ppmv), when Ames CASA indicates an earlier drawdown of CO₂ by the
509 land biosphere than does CASA-GFED3, and in January through March of 2010 (2-5 ppmv), when the
510 larger land to atmosphere carbon flux in Ames CASA from the latter part of 2009 is most evident. A
511 similar feature is not present in 2009 because it is too close to the initiation of the simulations, but
512 begins to be seen again in December, 2010. Land flux differences are typically less than 2 ppmv at

513 most remote SH locations. At NH locations, land flux differences manifest primarily as a difference in
514 seasonal cycle, consistent with the global flux differences shown in Figure 1. Localized differences,
515 such as those shown in Figure 2 during April and October, are generally not evident in monthly mean
516 CO₂ mixing ratios at NOAA ESRL stations because of their remote locations. The lack of evidence of
517 fire emissions, present in CASA-GFED3 and not Ames CASA, is striking, particularly at SH locations.
518 Analysis of the model's GFED3 biomass burning tracer shows that at all stations except Ascension
519 Island, deviations from the annual mean are smaller than 1 ppmv. At Ascension Island, biomass
520 burning events with greater than 1 ppmv influence only occur on a few days.

521

522 Even in locations and months where substantial differences in CO₂ mixing ratio due to land flux exist,
523 their magnitude is small enough that they are often difficult to distinguish from natural variability
524 evident in the observations. In the NH mid-latitudes, large differences in CO₂ mixing ratio attributable
525 to land flux would not be significantly larger than the expected model-data mismatch because of the
526 large variability evident in the observational records. Differences at NH high latitude stations and in the
527 SH are statistically significant, but only during certain seasons.

528

529 Differences between ocean flux models, which are nearly as large as land flux differences on a global
530 annual basis but exhibit much less seasonality, are even less evident at NOAA ESRL stations. Large
531 differences due to the difference in Southern Ocean sink exist only at stations south of 30°S but are
532 statistically significant in most months. Significant differences between the two ocean models at all
533 stations south of 30°S demonstrate the importance of ocean flux in that region. In this region, NOBM
534 fluxes are more realistic than ECCO2-Darwin fluxes which result in underestimates in surface CO₂
535 mixing ratio and errors in the gradient between tropical and SH high latitude CO₂. Observations
536 suggest a mean difference between Mauna Loa and SH high latitude stations during March, April, and
537 May (MAM) of 6.6 ppmv. While NOBM fluxes reproduce this spatial gradient fairly well, simulating a

538 mean difference of 6.1 ppmv, ECCO2-Darwin fluxes result in a gradient of 8.5 ppmv indicating that
539 Southern Ocean drawdown is too large during certain months. Large differences in ocean CO₂ flux are
540 also present in the NH (Figure 3), but the fact that they occur over smaller regions than in the SH
541 results in smaller mixing ratio differences at NH locations. The combination of smaller mixing ratio
542 differences and greater variability due to biosphere flux and fossil fuel emissions in the NH means that
543 NH ocean flux differences are difficult to distinguish from natural variability. Cold Bay and Shemya,
544 both in Alaska and strongly influenced by marine air, are two exceptions; at these locations differences
545 due to ocean flux are large enough to be distinguished.

546

547 **3.2. Comparisons with HIPPO aircraft observations**

548 Beginning in 2009, the HIAPER (High-performance Instrumented Airborne Platform for
549 Environmental Research) Pole-to-Pole Observations (HIPPO) project has measured atmospheric CO₂,
550 other trace gases, and aerosols in a series of field campaigns. HIPPO missions consist of north-south
551 transects spanning the Pacific ocean from 85°N to 67°S with profiles of atmospheric trace gases every
552 2.2° latitude [Wofsy et al., 2011]. HIPPO data have been used extensively for validation and calibration
553 of remote sensing data products (e.g. Wunch et al. [2010]; Inoue et al. [2013]), for evaluation of
554 atmospheric transport models (e.g. Keppel-Aleks et al. [2013]; Mann et al. [2012]), and to better
555 understand emission and transport processes (e.g. Keppel-Aleks et al. [2012]; Kipling et al. [2013]).

556

557 Three of the five planned HIPPO missions occurred during the 2009-2010 period. Because the first
558 deployment occurred in January, 2009, shortly after the beginning of the CMS target period, our results
559 focus on the second and third HIPPO deployments (HIPPO-2 and HIPPO-3). Both campaigns sampled
560 the Pacific Ocean between 150°E and 90°W. During the second deployment, the majority of samples
561 were collected between 180°W and 150°W while during the third campaign, sampling was focused
562 between 170°W and 140°W. HIPPO data shown are 10 second average data. GEOS-5 pseudo-data are

563 created by sampling the appropriate three-hour model output, interpolating model profiles to the
564 measurement latitude and longitude, and then interpolating resulting profiles vertically to replicate the
565 measurement conditions as well as possible.

566

567 Figure 9 shows a longitudinal cross section comparison between the observed CO₂ data and GEOS-5
568 simulations for HIPPO-2, which occurred during October and November, 2009. Relative to the HIPPO
569 data, GEOS-5 assuming flux combination CG-NO tends to overestimate CO₂ throughout the sampled
570 region of the Pacific. Errors are greater in the NH (1-5 ppmv) than the SH (less than 3 ppmv). The
571 model is able to capture the major features of the observed transect, including elevated near surface
572 CO₂ mixing ratios at NH high latitudes and enhanced CO₂ mixing ratios aloft at 30°S and 30°N.
573 Transects simulated using flux combinations AC-NO and CG-ED are generally similar to the results
574 produced by flux combination CG-NO, but with some notable exceptions. The differences between
575 observations and the CG-NO and AC-NO simulations shows that at NH locations, overestimates in
576 CO₂ are worse when using Ames CASA fluxes compared to CASA-GFED. Near 30°S, the Ames
577 CASA fluxes improve the model's underestimate of near surface CO₂ during SH spring. However,
578 throughout most of the campaign Ames CASA fluxes result in greater errors because the difference in
579 the magnitude of the global land sink is less realistic. The use of ECCO2-Darwin fluxes in flux
580 combination CG-ED results in stronger ocean drawdown at NH high latitudes during autumn. This
581 helps to reduce the overestimate in CO₂ mixing ratios at these locations, but elsewhere, model
582 performance is degraded because of the overall weaker ocean sink.

583

584 During HIPPO-3, conducted during March and April, 2010 (Figure 10), model performance is
585 generally similar to performance during the second deployment. All flux combinations result in
586 overestimates in NH CO₂ but this is worse when using Ames CASA fluxes than when using CASA-
587 GFED fluxes. During March and April, the much stronger southern ocean sink in ECCO2-Darwin is

588 evident in observations near 60°S, resulting in larger errors in this region compared to NOBM.

589

590 We also evaluate the ability of HIPPO-2 and HIPPO-3 data to discern between flux combinations in
591 different regions of the atmosphere following Huntzinger et al. (2011). Model data mismatch is
592 estimated here as the variance of CO₂ observations within 10 degree latitude bins for layers from the
593 surface to 2 km, from 2 to 4 km, from 4 to 6 km, from 6 to 8 km, and from 8 to 9 km. The Chi-squared
594 test statistic is calculated for each of these latitude-altitude regions using the number of observations,
595 the mean squared difference between simulations, and the model data mismatch. Dashed lines on
596 Figures 9 and 10 indicate regions where the test of variance indicates that flux difference induces
597 differences in atmospheric CO₂ mixing ratios that are statistically significant at the 95% confidence
598 level. During HIPPO-2, land flux differences are primarily detectable in the SH mid-latitudes. SH CO₂
599 differences are smaller than those in the NH, but data exhibit much less variability making the
600 differences caused by land flux easier to distinguish from the background variability of measurements.
601 Flux differences in the SH mid-latitudes also tend to be distinguishable through a deep layer of the
602 atmosphere, not only near the surface. Ocean fluxes are difficult to distinguish during the November
603 time frame of HIPPO-2 because the greatest disparity between ocean fluxes is in NH high latitudes
604 where background CO₂ variability is strong. Land fluxes during HIPPO-3, in contrast to HIPPO-2, are
605 detectable only below 4 km and at mid- to high latitudes of both hemispheres. Ocean flux differences
606 are strongly evident in the SH high and mid-latitude locations.

607

608 The HIPPO datasets are also extremely valuable for evaluating the model's ability to simulate vertical
609 gradients in atmospheric CO₂. Figure 11 shows the difference between mean CO₂ near the surface
610 (below 2 km) and mean CO₂ between 6 and 8 km calculated over 5 degree latitude bins for both
611 HIPPO observations and GEOS-5 pseudo-data. During HIPPO-2, GEOS-5 tends to reasonably
612 reproduce the observed interhemispheric gradient both at low levels and aloft. During HIPPO-3, all

613 flux combinations result in the model overestimating the gradient between northern and southern
614 hemisphere CO₂. However, the vertical gradient is reasonably well simulated with the exception of
615 ECCO2-Darwin which results in much stronger negative vertical gradients at mid- and high latitude SH
616 locations than either simulated by NOBM or observed. During HIPPO-3, the earlier spring drawdown
617 in the NH Ames CASA fluxes also results in a slight overestimate in the vertical gradient in CO₂ at
618 most locations. The fact that the interhemispheric gradient in CO₂ is overestimated during HIPPO-3
619 while local vertical gradients appear reasonable could indicate either errors in the assumed source
620 distributions and magnitudes or an error in large scale transport pathways.

621

622 **3.3. Comparison with TCCON column CO₂ observations**

623 The Total Carbon Column Observing Network (TCCON), established in 2004, is a network of ground-
624 based Fourier Transform Spectrometers recording near-infrared direct solar spectra for a number of
625 atmospheric trace gases including CO₂ [Wunch et al., 2011a]. Data are collected continuously during
626 daylight hours when viewing is not obscured by optically thick cloud and aerosol. Observations are
627 estimated to have a precision as high as 0.25% (~ 1 ppmv) under clear sky conditions. In contrast to the
628 NOAA ESRL network, TCCON provides total column CO₂ observations comparable to the
629 observations made by GOSAT. TCCON is also much smaller than the NOAA flask network with 16
630 stations operating during the 2009-2010 study period considered here (Figure 4). TCCON observations
631 are the primary calibration and validation dataset for GOSAT [Wunch et al., 2011b] and their use for
632 flux inference has been demonstrated by Chevallier et al. [2011]. For this analysis, TCCON
633 observations are slightly adjusted to account for laser sampling errors noted by Messerschmidt et al.
634 [2012] and Dohe et al. [2013].

635

636 In order to compare with TCCON observations, GEOS-5 CO₂ fields are sampled at the appropriate
637 observation time and convolved with the averaging kernel and a priori profile information appropriate

638 for each station as described in Wunch et al. (2010). Simulated and observed daily means of column
639 CO₂ are then calculated; the results for four stations are shown in Figure 12. As is expected, column
640 CO₂ observations exhibit smaller seasonal variations than do surface observations. At both high NH
641 mid-latitude stations shown (Garmisch and Lamont), GEOS-5 simulations are able to reasonably
642 reproduce the observed column CO₂ mixing ratios with a slight underestimation in the amplitude of the
643 seasonal cycle (average underestimate over all NH mid- and high latitude stations of 1.4 ppmv for CG-
644 NO fluxes). Differences due to the flux combination assumed are small, typically no more than a few
645 ppmv, but the difference in seasonal cycle between CASA-GFED3 and Ames CASA is evident
646 particularly during NH spring. At the Izaña, Tenerife station, GEOS-5 simulations overestimate
647 summer CO₂ mixing ratios relative to observations, a feature also found in the comparison against
648 Mauna Loa surface observations (Figure 5).

649

650 Monthly mean errors between GEOS-5 simulations and TCCON observations (Figure 13) show that
651 model errors are typically less than 3 ppmv when flux combination CG-NO is assumed. The model
652 tends to overestimate CO₂ at NH locations during spring and summer while during winter months,
653 errors are small (less than 1 ppmv at most stations). Model errors at NOAA ESRL NH stations (Figure
654 6) during winter are greater than 5 ppmv at some locations, resulting in small overestimates in seasonal
655 cycle amplitude at the surface (Figure 7). In contrast, the model slightly underestimates the seasonal
656 cycle amplitude at NH TCCON locations. This may be because of errors in simulated transport during
657 winter months resulting in too much CO₂ near the surface, or because of differences in sampling
658 locations between the ESRL and TCCON networks.

659

660 Figure 14 shows differences in column CO₂ mixing ratio at TCCON stations due to the difference in
661 land and ocean fluxes assumed. The seasonality of differences due to differences between Ames CASA
662 and CASA-GFED3 is similar to the analysis of surface observations, but the magnitude of flux

663 differences manifests as a smaller magnitude atmospheric mixing ratio difference when the column is
664 considered. Despite the smaller total magnitude of the column differences, they are distinguishable
665 from natural variability at most NH stations during late spring and early summer, and winter, 2010.
666 This is largely due to the weaker natural variability in the column data and greater data volume yielded
667 by continuous measurements. The Ames CASA fluxes result in lower CO₂ (0.5-1.5 ppmv) at NH
668 stations during MAM 2009, and higher (2-3 ppmv) CO₂ at NH mid- and high latitude locations during
669 2010 winter, degrading performance relative to the CASA-GFED3 fluxes.

670

671 Detecting ocean flux differences at TCCON locations is considerably more difficult than detecting land
672 flux differences. Data from the Lauder station shows some ability to differentiate between fluxes
673 during some months of 2010, but data are not available at this site during 2009. These results point to
674 the difficulty of detecting ocean CO₂ flux differences with column observations. The nearest surface
675 station, Baring Head (BHD), New Zealand, indicates a higher percentage of data useful for
676 differentiating between ocean fluxes throughout the year. Atmospheric mixing ratio differences caused
677 by ocean flux uncertainty are typically smaller than differences caused by land flux and, as a result,
678 more difficult to separate from natural variability. Additionally, the TCCON network includes no
679 stations further south than Lauder making it impossible to directly observe Southern Ocean flux
680 differences.

681

682 **3.4. Comparisons with AIRS satellite observations**

683 The Atmospheric Infrared Sounder (AIRS) instrument aboard the Aqua satellite provides a record of
684 mid-tropospheric CO₂ from 2002 to present. Because data are collected during both day and night and
685 retrievals are performed in the presence of clouds, AIRS data provide unprecedented global coverage.
686 Accuracies are reported to be 2 ppmv or better by Chahine et al. [2005] with a nadir footprint of 90 by
687 90 km² [Chahine et al., 2008]. AIRS CO₂ observations have been used to study temporal (e.g. Jiang et

688 al. [2010]; Li et al. [2010]) and spatial (e.g. Ruzmaikin et al. [2012]; Bai et al. [2010]) variability in
689 mid-tropospheric CO₂, to evaluate atmospheric CO₂ simulations (e.g. Feng et al. [2011]), and for data
690 assimilation studies (e.g. Liu et al. [2012]; Engelen et al. [2009]).

691

692 GEOS-5 is sampled according to AIRS version 5 CO₂ observations using level 2 swath data. Simulated
693 CO₂ profiles for the appropriate 3-hour period are interpolated to the location of each observation that
694 passes AIRS quality assurance procedures. AIRS CO₂ weighting functions specific to the observation
695 latitude and background CO₂ mixing ratio are applied as described in Chahine et al. [2008]. Monthly
696 mean simulated and observed CO₂ mixing ratios are then calculated over 5 degree latitude by 5 degree
697 longitude regions.

698

699 Figure 15 presents a comparison between AIRS observations and GEOS-5 simulated CO₂ for October,
700 2009, a month that coincides with the beginning of the HIPPO-2 campaign. There is little agreement
701 between the AIRS observations and the CG-NO GEOS-5 simulation. AIRS indicates peak CO₂ mixing
702 ratios in SH mid-latitudes with minimum values in tropics and sub-tropics of both hemispheres. In
703 contrast, GEOS-5 indicates enhanced mid-tropospheric CO₂ over South America and SH Africa, likely
704 because of the lofting of fire emissions in these regions (Figure 2). GEOS-5 does not indicate the
705 presence of enhanced CO₂ mixing ratios in the mid-latitudes of either hemisphere resulting in a 2-5
706 ppmv underestimate in these regions relative to AIRS. In biomass burning regions, however, GEOS-5
707 overestimates mid-tropospheric CO₂ by 1-3 ppmv relative to AIRS. These results are in sharp contrast
708 to the comparison between HIPPO-2 CO₂ observations and GEOS-5 presented in Figure 9; that
709 comparison indicates that GEOS-5 tends to overestimate mid-tropospheric CO₂ in the 30-60°S band
710 over the Pacific Ocean from 0 to 8 km. The discrepancy between AIRS and GEOS-5 in this region
711 cannot be easily explained by errors in GEOS-5 vertical transport because Figure 11 indicates that
712 GEOS-5 vertical CO₂ gradients throughout the SH differ from observed vertical gradients by less than

713 1 ppmv. NH underestimates of GEOS-5 relative to AIRS also coincide with regions where HIPPO-2
714 data indicate that GEOS-5 is overestimating CO₂ by 2-4 ppmv in the mid-troposphere.

715

716 AIRS data show a significant amount of temporal variability evident in the model data mismatch
717 calculated for each 5 by 5 degree grid box (Figure 15). Model data mismatch is calculated for each grid
718 cell by computing daily means of available data during the 2009-2010 period, fitting a smooth curve
719 through the daily data, and calculating the standard deviation of the residuals, similar to the calculation
720 of model data mismatch for NOAA ESRL and TCCON data. Using this method, expected model data
721 mismatch is typically between 1.5 and 4 ppmv with the largest observed variability in the mid- and
722 high latitudes of both hemispheres. Differences between GEOS-5 simulations assuming different flux
723 combinations are typically small in the mid- and upper troposphere region represented by AIRS
724 observations. The background difference between flux combinations CG-NO and AC-NO, due to the
725 difference in annual growth rate, is 1 ppmv during October. In biomass burning regions in SH Africa
726 and South America, Ames CASA fluxes result in CO₂ mixing ratios 1 ppmv less than CASA-GFED
727 fluxes, or nearly 2 ppmv if the difference in annual growth rate is removed. CASA-GFED results in 2
728 ppmv more CO₂ over equatorial Africa and SE Asia, but if the difference in growth rate is removed,
729 these differences are only 1 ppmv. In tropical regions, where vertical transport of surface fluxes is most
730 rapidly communicated to the mid- and upper troposphere, spatial patterns of atmospheric mixing ratio
731 differences resemble underlying surface flux differences shown in Figure 2. However, the magnitude of
732 these differences (1-2 ppmv) is typically smaller than the expected model data mismatch leading to a
733 lack of ability to discern between flux scenarios with any statistical significance. Ocean flux
734 differences (shown in Figure 15f), which are smaller in magnitude but more consistent throughout the
735 year, are not evident when model simulations are sampled with AIRS pressure weighting functions
736 except as a very slight modification to the north-south CO₂ gradient that is not statistically significant.
737 AIRS is not able to observe the large ocean flux differences in high latitude regions; data are not

738 available south of 60°S and in NH high latitudes, slow vertical transport times produce no spatial
739 signature of the underlying flux differences in the mid- and upper troposphere where AIRS
740 measurements are most sensitive.

741

742 In all months during 2009-2010, AIRS data would have little utility for differentiating between spatial
743 patterns of flux differences. Figure 16 shows the percentage of 5 degree grid boxes per latitude bin that
744 contain statistically significant land and ocean flux differences for each month during the study period.
745 Land flux differences would seldom be detectable and only in a small number of grid cells in the
746 tropics and subtropics. Ocean flux differences are not detectable in any month or location because the
747 magnitude of the mixing ratio differences is always smaller than the variability inherent in the
748 measurements.

749

750 Figure 17 shows a comparison between simulated and observed monthly zonal mean mixing ratios
751 derived from AIRS data and model pseudo-data. GEOS-5 fails to capture the seasonal cycle of zonal
752 mean CO₂ observed by AIRS. GEOS-5 mid-tropospheric CO₂ exhibits a much stronger minimum
753 during the NH growing season (July-October) than does AIRS. AIRS indicates much stronger maxima
754 in the NH high latitudes during spring than indicated by GEOS-5, and a secondary maxima during
755 September that is not simulated by GEOS-5. GEOS-5 tends to produce larger than observed CO₂
756 mixing ratios in the tropics and subtropics throughout the year. In the SH mid-latitudes, GEOS-5
757 underestimates AIRS zonal mean CO₂ with the largest differences in spring and autumn.

758

759 **3.5. Comparisons with GOSAT satellite observations**

760 The Greenhouse gases Observing SATellite (GOSAT), launched in 2009 by the Japanese Aerospace
761 Exploration Agency (JAXA), monitors CO₂ and methane from space by analyzing high resolution
762 spectra of reflected sunlight within several near infrared bands (Kuze et al. [2009]; Yokota et al.,

763 [2009]). The near infrared measurement technique allows for greater sensitivity to CO₂ near the surface
764 than the thermal infrared measurements of AIRS that are sensitive primarily to the mid- and upper
765 troposphere. Retrievals are only performed under cloud-free conditions resulting in a much smaller
766 yield of usable soundings than AIRS. GOSAT observations have been used to examine CO₂ mixing
767 ratios in megacities [Kort et al., 2012] and to estimate regional carbon fluxes (e.g. Takagi et al. [2011];
768 Basu et al. [2013]; Maksyutov et al. [2013]). Several different retrievals of column averaged CO₂ dry
769 air mole fraction (XCO₂) based on GOSAT observations are currently being produced. In this work,
770 we use data produced by NASA's Atmospheric CO₂ Observations from Space (ACOS) effort, a
771 collaboration between the original JAXA GOSAT team and NASA's Orbiting Carbon Observatory
772 (OCO) science team [Crisp et al., 2012]. These data are produced using a modified version of the OCO
773 retrieval algorithm applied to GOSAT observed spectra as described in O'Dell et al. [2012] and have
774 been validated extensively using TCCON measurements (e.g. Wunch et al. [2010, 2011a, 2011b]).
775 Based on retrievals of realistic simulated observations, O'Dell et al. [2012] estimate that observations
776 contain RMS XCO₂ errors of ~1 ppmv and a positive bias of 0.3 ppmv. Dated used here are ACOS
777 version 3.4.

778

779 GEOS-5 is sampled at GOSAT measurement times and locations. Model profiles are interpolated to the
780 twenty atmospheric pressure levels used in the retrieval process and averaging kernels provided as part
781 of the ACOS data product are applied to convolve GEOS-5 mixing ratios with the a priori CO₂ mixing
782 ratio profile. XCO₂ is calculated from this simulated pseudo data profile. Monthly mean simulated and
783 observed CO₂ mixing ratios are then calculated over 5 degree latitude by 5 degree longitude regions as
784 was done for the AIRS data comparison.

785

786 During October, 2009, the spatial distribution of GEOS-5 assuming flux CG-NO compares much more
787 favorably with GOSAT observations (Figure 18) than with AIRS (Figure 15). The distribution of

788 GOSAT XCO₂ shows fairly uniform mixing ratios globally with slightly larger XCO₂ over South
789 America and Africa. The GEOS-5 distribution is similar though XCO₂ is overestimated by 1-3 ppmv in
790 most locations. This is consistent with HIPPO-2 aircraft observations (Figure 9) which show similar
791 overestimates between 0 and 8 km. Model data mismatch was calculated as described above for AIRS
792 data; observations were binned into 5 degree grid cells, daily means calculated and then the standard
793 deviation of residuals around a smooth curve considered the model data mismatch. Using this
794 technique, estimated model data mismatch is typically between 0.5 and 1 ppmv. However, despite the
795 smaller model data mismatch values, differentiating between land and ocean fluxes remains difficult
796 because of the small number of observations collected. The difference in XCO₂ due to the underlying
797 difference in land flux would be primarily detectable over Africa. Ocean flux differences are too small
798 to be detectable at any location.

799

800 A comparison between GOSAT data and GEOS-5 simulations during July 2009 (Figure 19) shows
801 similar features in both models and observations. The model is able to reasonably reproduce the
802 observed spatial distribution of XCO₂ but tends to overestimate XCO₂ by up to 5 ppmv in most mid-
803 and high latitude locations. This is consistent with model overestimates of 3-5 ppmv at surface (Figure
804 6) and TCCON stations (Figure 13) during NH summer months. Model overestimates are larger during
805 July than during October because the land sink, at its peak during NH summer, is likely weaker in both
806 CASA-GFED and Ames CASA than in reality. The Ames CASA land fluxes, characterized by earlier
807 drawdown of CO₂ in the NH spring, results in smaller CO₂ mixing ratios during July in the NH than
808 CASA-GFED despite the fact that the annual total Ames CASA land sink is smaller. Though the
809 largest differences in land flux during July are in the NH high latitudes (Figure 3), the resultant
810 atmospheric mixing ratio difference is not generally detectable at these locations. Instead, land flux
811 differences are most readily observed over NH ocean locations and over North Africa because the
812 observations exhibit much less variability in these locations. As in October, ocean flux differences are

813 not large enough to be detectable at any locations.

814

815 Figure 16 also shows the percentage of 5 degree grid boxes per latitude bin that contain statistically
816 significant ($p < 0.05$) land and ocean flux differences for each month when GOSAT data are considered.

817 In contrast to AIRS and in spite of very sparse sampling, GOSAT observations would be able to
818 distinguish spatial pattern differences in atmospheric CO₂ due to differing land fluxes in both
819 hemispheres primarily during their spring and summer months. Ocean flux differences are statistically
820 significant in only a small number of grid cells in the SH tropics.

821

822 Simulated zonal mean XCO₂ from GEOS-5 agrees much more favorably with GOSAT data (Figure 20)
823 than with AIRS (Figure 17) as was the case when spatial distributions of CO₂ during October 2009
824 were considered. GEOS-5 tends to overestimate XCO₂ throughout the year with the largest degree of
825 overestimation during NH spring and summer. The use of Ames CASA instead of CASA GFED land
826 fluxes help to improve the overestimation during spring, but degrades the comparison with
827 observations during NH winter. Atmospheric CO₂ differences caused by differing land fluxes are
828 detectable when zonal means are considered primarily during spring and summer months in both
829 hemispheres. Ocean flux differences are too small to be detected by GOSAT observations in all months
830 and in all locations.

831

832 **4. Summary and Conclusions**

833 As part of NASA's CMS Flux Pilot Project, the GEOS-5 GCM was used to simulate atmospheric CO₂
834 mixing ratios during 2009 and 2010 using two sets of model-based land and ocean flux estimates. All
835 land and ocean flux estimates were informed by multiple satellite datasets and compared with
836 numerous observations with the goal of better understanding the constraint on carbon flux provided by
837 current observing systems. Despite the strong data constraint, the two land models used in this work

838 differed by an average of 1.7 Pg C in their estimate of the global land carbon sink during 2009 and
839 2010. Both land flux models estimate the land biosphere as a net source of carbon to the atmosphere
840 rather than a ~ 3 Pg C sink as indicated by the Global Carbon Budget [Le Quere et al., 2013]. While the
841 presence of a missing land sink was indicated by TRANSCOM models over 10 years ago (e.g. Gurney
842 et al. [2002]) and much progress has since been made in understanding the processes which may
843 account for the disparity between forward and inverse model flux estimates, the CMS FPP results show
844 that estimating realistic net land carbon fluxes at both global and regional scales remains a challenge
845 for models, even those informed by remote sensing observations. Ocean models differ by 1.4 Pg C per
846 year in global carbon flux. In ocean waters south of 30°S, model flux differences are 25% as large as
847 the mean regional flux and the phasing of the seasonal cycle also differs substantially. The magnitude
848 of these differences underscores the continued uncertainty surrounding the ability of natural carbon
849 reservoirs to compensate for increasing fossil fuel emissions.

850

851 Despite considerable differences in global flux, the performance of GEOS-5 simulations is generally
852 similar because of the strong role played by meteorological transport. GEOS-5 simulations assuming
853 all flux combinations tended to overestimate surface CO₂ mixing ratios in the NH, particularly during
854 spring and summer seasons when errors are 3-5 ppmv. The amplitude and phasing of the seasonal cycle
855 at most NH locations is reasonably well simulated though the model tends to slightly overestimate the
856 amplitude of the seasonal cycle at the surface. All flux combinations examined in this study struggle to
857 adequately reproduce the timing of the observed seasonal cycle at SH stations and at mid- and high
858 latitude SH locations and strongly overestimate the magnitude of seasonal variability by as much as 5
859 ppmv in some locations. Because the seasonal cycle is much smaller at SH mid- and high latitude
860 compared to NH locations, errors of this magnitude are greater than the observed seasonal cycle
861 amplitude.

862

863 Comparisons with HIPPO observations provide the opportunity to evaluate the realism of simulated
864 vertical mixing processes. As expected from flux and surface CO₂ comparisons, GEOS-5 tends to
865 overestimate CO₂ mixing ratios relative to aircraft observations, particularly in the NH mid- and high
866 latitudes. Differences between mean near surface CO₂ and CO₂ observed between 6 and 8 km show the
867 model succeeds in simulating realistic vertical gradients in CO₂ during October-November, 2009 and
868 March-April, 2010. The HIPPO data comparisons indicate that during these months and over the
869 Pacific, model errors in surface and column CO₂ are not likely to be attributable to vertical transport
870 errors.

871
872 GEOS-5 simulations succeed in capturing the major features of column CO₂ observations from
873 TCCON and GOSAT though the model tends to overestimate CO₂ by 3-4 ppmv during NH summer
874 regardless of the combination of fluxes used. GEOS-5 is also able to reproduce the spatial patterns of
875 XCO₂ observed by GOSAT. When compared with AIRS mid-tropospheric CO₂ mixing ratios,
876 however, the model fails to represent either observed spatial distributions or seasonal cycle information.

877
878 Comparisons between GEOS-5 and surface, aircraft, TCCON, and GOSAT data reveal a consistent
879 picture of the capabilities of contemporary models to reproduce atmospheric CO₂ observations. Models
880 tend to overestimate CO₂ because the combined land and ocean flux assumed is too weak.
881 Overestimates at the surface are largest during the NH summer months, consistent with an
882 underestimate in the strength of land biosphere sink that is most active in this region and during this
883 season. HIPPO observations show that vertical gradients are reasonably well simulated, meaning that
884 excess CO₂ in the model atmosphere is distributed through the deep layer (typically between the
885 surface and 8 km) sampled by the aircraft. Total column CO₂ measurements (TCCON and GOSAT)
886 are overestimated by GEOS-5 in the same locations and seasons as indicated by comparisons with in
887 situ measurements, but by a slightly smaller amount. It is more difficult to reconcile AIRS mid-

888 tropospheric CO₂ observations with other observational records and GEOS-5. While AIRS vertical
889 weighting functions peak in the mid-troposphere, they are strongly sensitive to the upper
890 troposphere/lower stratosphere (UTLS). Depending on the latitude of the observation, 60-71% of the
891 observed signal comes from pressures lower than 500 hPa. In order for GEOS-5 simulations to both
892 agree with TCCON and GOSAT and disagree with AIRS CO₂ observations, the model would have to
893 have large errors in the UTLS including overestimates in the tropics and alternating under- and
894 overestimates in the NH high latitudes. While there is no evidence of this type of model bias in the
895 comparisons with HIPPO presented here, sampling above 8 km is fairly sparse. More work is needed to
896 thoroughly evaluate the consistency of AIRS and other CO₂ data products, especially if they are to be
897 used together in joint assimilation and inversion frameworks.

898

899 Despite differences between land and ocean flux estimates greater than 1 Pg C, resulting differences in
900 atmospheric mixing ratio at remote surface sites are small, typically less than 5 ppmv at the surface and
901 3 ppmv in the column, and difficult to distinguish from natural observed variability. At NOAA ESRL
902 surface stations, the difference between the Ames CASA and CASA-GFED land fluxes manifests most
903 clearly in the atmosphere during NH spring and winter months. Outside of NH high latitude locations,
904 CO₂ mixing ratio differences are rarely statistically significant when the natural variability of
905 observations is considered. TCCON observations succeed in detecting seasonal differences between
906 land flux estimates in NH winter and spring months. Though flux differences result in smaller mixing
907 ratio differences in the column, the continuous monitoring strategy of TCCON produces larger
908 quantities of data, which facilitates separation of flux differences from natural variability. AIRS
909 satellite observations are unable to discern between land flux models because the variability of
910 observations is larger than the magnitude of the resulting simulated mixing ratio differences in the mid-
911 and upper troposphere. GOSAT observations are better suited to observing flux differences because of
912 greater sensitivity near the surface and they show an ability to distinguish between land flux estimates

913 during summer months. However, GOSAT's ability to differentiate between land flux estimates
914 remains limited by sparse coverage.

915

916 Ocean flux differences, characterized by large disparities at high latitudes of both hemispheres, are
917 primarily distinguishable from natural variability at SH mid- and high latitude surface stations. HIPPO
918 aircraft observations also succeed in differentiating Southern Ocean flux differences from natural
919 variability. While in situ observations show an ability to discern between ocean flux estimates, remote
920 sensing techniques fail for several reasons. The sparse TCCON network contains no stations below
921 45°S, the region where flux uncertainty is greatest and where surface observations show the greatest
922 ability to differentiate between ocean flux estimates. As was the case with land flux, ocean flux
923 estimates produce mixing ratio differences that are smaller than the temporal variability inherent in
924 AIRS data. GOSAT observations, which have smaller single observation errors than AIRS and greater
925 sensitivity near surface, do not observe ocean regions south of 40°S. The difference between the ocean
926 flux models included in the CMS FPP highlights the larger uncertainty in how high latitude ocean
927 carbon storage has changed in recent decades and how it may continue to change in response to future
928 climate change. The inability of current remote sensing observations to detect the large differences in
929 ocean flux presented here highlights the need for continued in situ observations and the development of
930 remote sensing techniques which have the potential to increase data yield over high latitude regions.

931

932 Small mixing ratio differences resulting from flux differences make it difficult to assess the
933 performance of the individual flux estimates. While CASA-GFED compares more favorably than
934 Ames CASA to the total land flux estimate from the Global Carbon Budget [Le Quere et al., 2013], it
935 still diagnoses the land biosphere as a net source of CO₂ to the atmosphere rather than a sink. NOBM
936 overestimates the magnitude of the ocean sink relative to the Global Carbon Budget and ECCO2-
937 Darwin, but because both land flux estimates underestimate the magnitude of the global sink, the CG-

938 NO simulation produces the most realistic atmospheric growth rate when compared with NOAA
939 surface observations. The main land flux difference that manifests in atmospheric CO₂ mixing ratios is
940 the difference in seasonal cycle between CASA-GFED and Ames CASA. While the earlier spring
941 drawdown in Ames CASA results in slightly better comparisons with surface observations during the
942 spring, earlier release of CO₂ from the land to the atmosphere in autumn degrades comparisons during
943 winter. Small-scale differences between CASA-GFED and Ames CASA, including the presence of fire
944 emissions (CASA-GFED) or redistribution of crop CO₂ (Ames CASA), are difficult to evaluate
945 because these differences are not readily evident in the available observational CO₂ records. While
946 ECCO2-Darwin produces a global ocean sink magnitude nearly equal to the Global Carbon Budget
947 estimate, comparisons with surface and aircraft data show that it is less realistic than NOBM over the
948 Southern Ocean where it diagnoses a larger sink with stronger seasonal variations.

949

950 While all flux combinations perform reasonably well in reproducing observed seasonal cycles and
951 spatial gradients in CO₂, the troubling implication of this agreement between simulations is that even
952 differences between flux models on the order of Pg C are difficult to disentangle using current
953 atmospheric CO₂ observations. Ocean flux differences are particularly difficult to discern because they
954 are smaller and tend to occur in high latitude regions that are poorly observed by current remote
955 sensing platforms. The use of aircraft campaigns, such as HIPPO, provide a valuable complement to
956 existing long term carbon monitoring strategies and may become more important in the future as
957 natural carbon reservoirs in remote high latitude locations respond to a changing climate.

958

959 Observations from OCO-2 are expected to greatly improve data yield over much of the globe, which
960 would allow for a greater ability to use space-based observations for flux discrimination. However,
961 OCO-2 will also be limited in its ability to observe high latitude locations meaning that it will not be
962 able to directly observe many locations associated with large flux uncertainty throughout the year. The

963 ASCENDS mission, which focuses on active rather than passive observations of CO₂, would improve
964 the ability of GOSAT and OCO-2 to observe high latitude locations, but is not likely to launch until the
965 2020s. These observational challenges underscore the need for a variety of types of CO₂ observations
966 to help fill the gaps left by satellite observations and to provide additional, complementary information
967 to maximize the impact of satellite observations in regions where they are available.

968

969 The small differences in atmospheric CO₂ mixing ratio due to flux also underscore the importance of
970 quality meteorological analyses and models. The ability to reasonably simulate small gradients in
971 atmospheric CO₂ mixing ratio and to successfully track the transport of air parcels from the surface to
972 the locations and altitudes at which measurements occur is key to the success of atmospheric inversion
973 studies which seek to reduce uncertainty in natural carbon sinks using atmospheric CO₂ observations.
974 Work is ongoing as part of the CMS project to better understand and quantify transport model errors.

975

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985

986 **References**

987

988 Adcroft, A., J. Campin, C. Hill, and J. Marshall (2004), Implementation of an atmosphere-ocean
989 general circulation model on the expanded spherical cube, *Monthly Weather Review*, *132*, 2845–2863.

990

991 Bai W., X. Zhang, and P. Zhang (2010), Temporal and spatial distribution of tropospheric CO₂ over
992 China based on satellite observations, *Chinese Science Bulletin*, *55*(31), 3612-3618.

993

994 Bala, G., N. Devaraju, R. K. Chaturvedi, K. Caldeira, and R. Nemani (2013), Nitrogen deposition: how
995 important is it for global terrestrial carbon uptake?, *Biogeosciences*, *10*, 7147-7160, doi:10.5194/bg-10-
996 7147-2013.

997

998 Ballantyne, A. P., C. B. Alden, J. B. Miller, P. P. Tans, and J. W. C. White (2012), Increase in observed
999 net carbon dioxide uptake by land and oceans during the last 50 years, *Nature*, *488*, 70-72.

1000

1001 Basu, S., S. Guerlet, A. Butz, S. Houweling, O. Hasekamp, I. Aben, P. Krummel, P. Steele, R.
1002 Langenfelds, M. Torn, S. Biraud, B. Stephens, A. Andrews, and D. Worthy (2013), Global CO₂ fluxes
1003 estimated from GOSAT retrievals of total column CO₂, *Atmospheric Chemistry and Physics*
1004 *Discussions*, *13*, 4535-4600, doi:10.5194/acpd-13-4535-2013.

1005

1006 Bellassen, V., N. Viovy, S. Luysaert, G. Le Maire, M.-J. Schelhaas, and P. Ciais (2011),
1007 Reconstruction and attribution of the carbon sink of European forests between 1950 and 2000, *Global*
1008 *Change Biology*, *17*, 3274–3292. doi: 10.1111/j.1365-2486.2011.02476.x.

1009

1010 Boden, T. A., G. Marland, and R. J. Andres (2011), Global, Regional, and National Fossil-Fuel CO₂
1011 Emissions. Carbon Dioxide Information Analysis Center, Oak Ridge National Laboratory, U.S.
1012 Department of Energy, Oak Ridge, Tenn., U.S.A. doi 10.3334/CDIAC/00001_V2011.

1013

1014 Bousquet, P., P. Ciais, P. Peylin, M. Ramonet, and P. Monfray (1999), Inverse modeling of annual
1015 atmospheric CO₂ sources and sinks: 1. Method and control inversion, *Journal of Geophysical Research*,
1016 *104*(D21), 26161–26178, doi:10.1029/1999JD900342.

1017

1018 Brix, H., D. Menemenlis, C. Hill, S. Dutkiewicz, O. Jahn, D. Wang, K. Bowman, and H. Zhang (2014),
1019 Using Green's Functions to initialize and adjust a global, eddying ocean biogeochemistry general
1020 circulation model, *Ocean Modelling*, submitted.

1021

1022 Chahine, M., C. Barnet, E. T. Olsen, L. Chen, and E. Maddy (2005), On the determination of
1023 atmospheric minor gases by the method of vanishing partial derivatives with application to CO₂,
1024 *Geophysical Research Letters*, *32*, L22803, doi:10.1029/2005GL024165.

1025

1026 Chahine, M. T., L. Chen, P. Dimotakis, X. Jiang, Q. Li, E. T. Olsen, T. Pagano, J. Randerson, and Y. L.
1027 Yung (2008), Satellite remote sounding of mid-tropospheric CO₂, *Geophysical Research Letters*, *35*,
1028 L17807, doi:10.1029/2008GL035022.

1029

1030 Chevallier, F., et al. (2011), Global CO₂ fluxes inferred from surface air-sample measurements and
1031 from TCCON retrievals of the CO₂ total column, *Geophysical Research Letters*, *38*, L24810,
1032 doi:10.1029/2011GL049899.

1033

1034
1035 Ciais, P., P. Bousquet, A. Freibauer, and T. Naegler (2007), Horizontal displacement of carbon
1036 associated with agriculture and its impacts on atmospheric CO₂, *Global Biogeochemical Cycles*,
1037 doi:10.1029/2006GB002741.
1038
1039 Conway, J., P. P. Tans, L. S. Waterman, K. W. Thoning, D. R. Kitzis, K. A. Masarie, and N. Zhang,
1040 (1994), Evidence of interannual variability of the carbon cycle from the NOAA/CMDL global air
1041 sampling network, *Journal of Geophysical Research*, *99*, 22831-22855.
1042
1043 Crisp, D., et al. (2012), The ACOS CO₂ retrieval algorithm – Part II: Global XCO₂ data
1044 characterization, *Atmos. Meas. Tech.*, *5*, 687-707, doi:10.5194/amt-5-687-2012.
1045
1046 Doney, S. C., et al. (2004), Evaluating global ocean carbon models: The importance of realistic physics,
1047 *Global Biogeochemical Cycles*, *18*, GB3017, doi:10.1029/2003GB002150.
1048
1049 Dutkiewicz, S., M. Follows, and J. Bragg (2009), Modeling the coupling of ocean ecology and
1050 biogeochemistry, *Global Biogeochem.Cycles*, *23*, GB4017.
1051
1052 Engelen, R. J., S. Serrar, and F. Chevallier (2009), Four-dimensional data assimilation of atmospheric
1053 CO₂ using AIRS observations, *Journal of Geophysical Research*, *114*, D03303,
1054 doi:10.1029/2008JD010739.
1055
1056 Esser, G., J. Kattge, and A. Sakalli (2011), Feedback of carbon and nitrogen cycles enhances carbon
1057 sequestration in the terrestrial biosphere, *Global Change Biology*, *17*, 819–842, doi: 10.1111/j.1365-
1058 2486.2010.02261.x.
1059
1060 Feng, L., P. I. Palmer, Y. Yang, R. M. Yantosca, S. R. Kawa, J.-D. Paris, H. Matsueda, and T. Machida
1061 (2011), Evaluating a 3-D transport model of atmospheric CO₂ using ground-based, aircraft, and space-
1062 borne data, *Atmospheric Chemistry and Physics*, *11*, 2789-2803, doi:10.5194/acp-11-2789-2011
1063
1064 Fleischer, K., K. T. Rebel, M. K. van der Molen, J. W. Erisman, M. J. Wassen, E. E. van Loon, L.
1065 Montagnani, C. M. Gough, M. Herbst, I. A. Janssens, D. Gianelle, and A. J. Dolman (2013), The
1066 contribution of nitrogen deposition to the photosynthetic capacity of forests, *Global Biogeochemical*
1067 *Cycles*, *27*, 187–199, doi:10.1002/gbc.20026.
1068
1069 Follows, M., T. Ito, and S. Dutkiewicz (2006), On the solution of the carbonate chemistry system in
1070 ocean biogeochemistry models, *Ocean Modelling*, *12*, 290–301.
1071
1072 Follows, M., S. Dutkiewicz, S. Grant, and S. Chisholm (2007), Emergent biogeography of microbial
1073 communities in a model ocean, *Science*, *315*, 1843–1846.
1074
1075 Follows, M. and S. Dutkiewicz (2011), Modeling diverse communities of marine microbes, *Annu. Rev.*
1076 *Marine Science*, *3*, 427–451.
1077
1078 Gerber, S., L. O. Hedin, S. G. Keel, S. W. Pacala, and E. Shevliakova (2013), Land use change and
1079 nitrogen feedbacks constrain the trajectory of the land carbon sink, *Geophysical Research*
1080 *Letters*, *40*, 5218–5222, doi:10.1002/grl.50957.
1081
1082

1083 Gregg, W. W., P. Ginoux, P. S. Schopf, and N. W. Casey (2003), Phytoplankton and Iron: Validation
1084 of a global three-dimensional ocean biogeochemical model, *Deep-Sea Research II*, 50, 3143-3169.
1085

1086 Gregg, W. W. and N. W. Casey (2007), Modeling coccolithophores in the global oceans, *Deep-Sea*
1087 *Research II*, 54, 447-477.
1088

1089 Gregg, W. W., N. W. Casey, and C. S. Rousseaux (2013), Global surface ocean estimates in a model
1090 forced by MERRA, NASA Global Modeling and Assimilation Series, M. Suarez, ed., NASA
1091 Technical Memorandum 2012-104606, Vol. 31, 32 pp.

1092 Gurney, K. R., et al. (2002), Towards robust regional estimates of CO₂ sources and sinks using
1093 atmospheric transport models, *Nature*, 415, 626-630.
1094

1095 Heimbach, P., D. Menemenlis, M. Losch, J. Campin, and C. Hill (2010), On the formulation of sea-ice
1096 models. Part 2: Lessons from multi-year adjoint sea ice export sensitivities through the Canadian Arctic
1097 Archipelago, *Ocean Modelling*, 33, 145–158.
1098

1099 Hollinger, D. Y., et al. (2004), Spatial and temporal variability in forest–atmosphere CO₂ exchange,
1100 *Global Change Biology*, 10: 1689–1706. doi: 10.1111/j.1365-2486.2004.00847.x
1101

1102 Huntzinger, D. N., S. M. Gourdj, K. L. Mueller, and A. M. Michalak (2011), The utility of continuous
1103 atmospheric measurements for identifying biospheric CO₂ flux variability, *Journal of Geophysical*
1104 *Research*, 116, D06110, doi:10.1029/2010JD015048.
1105

1106 Huntzinger, D. N., et al. (2012), North American Carbon Program (NACP) regional interim synthesis:
1107 Terrestrial biospheric model intercomparison, *Ecological Modelling*, 232, 144-157.
1108

1109 Inoue, M., et al. (2013), Validation of XCO₂ derived from SWIR spectra of GOSAT TANSO-FTS with
1110 aircraft measurement data, *Atmospheric Chemistry and Physics Discussions*, 13, 3203-3246,
1111 doi:10.5194/acpd-13-3203-2013.
1112

1113 Jiang, X., M. T. Chahine, E. T. Olsen, L. L. Chen, and Y. L. Yung (2010), Interannual variability of
1114 midtropospheric CO₂ from Atmospheric Infrared Sounder, *Geophysical Research Letters*, 37, L13801,
1115 doi:10.1029/2010GL042823.
1116

1117 Keppel-Aleks, G., et al. (2013), Atmospheric carbon dioxide variability in the Community Earth
1118 System Model: Evaluation and transient dynamic during the 20th and 21st centuries, *Journal of*
1119 *Climate*, 26, 4447-4475, doi:10.1175/JCLI-D-12-00589.1.

1120 Keppel-Aleks, G., (2012), The imprint of surface fluxes and transport on variations in total column
1121 carbon dioxide, *Biogeosciences*, 9, 875-891, doi:10.5194/bg-9-875-2012.
1122

1123 Kipling, Z., P. Stier, J. P. Schwarz, A. E. Perring, J. R. Spackman, G. W. Mann, C. E. Johnson, C. E.,
1124 and P. J. Telford (2013), Constraints on aerosol processes in climate models from vertically-resolved
1125 aircraft observations of black carbon, *Atmospheric Chemistry and Physics*, 13, 5969-5986,
1126 doi:10.5194/acp-13-5969-2013.
1127

1128 Kort, A. E., C. Frankenberg, C. E. Miller, and T. Oda (2012), Space-based observations of megacity
1129 carbon dioxide, *Geophysical Research Letters*, 39, L17806, doi:10.1029/2012GL052738.

1130
1131 Kuze, A., S. Hiroshi, N. Masakatsu, and H. Takashi (2009), Thermal and near infrared sensor for
1132 carbon observation Fourier-transform spectrometer on the Greenhouse Gases Observing Satellite for
1133 greenhouse gases monitoring, *Applied Optics*, 48, 6716-6733, doi:10.1364/AO.48.006716.
1134
1135 Lafleur, P. M., N. T. Roulet, J. L. Bubier, S. Frohking, and T. R. Moore (2003), Interannual variability
1136 in the peatland-atmosphere carbon dioxide exchange at an ombrotrophic bog, *Global Biogeochem.*
1137 *Cycles*, 17, 1036, doi:10.1029/2002GB001983, 2.
1138
1139 Law, R. M., et al. (2008), TransCom model simulations of hourly atmospheric CO₂: Experimental
1140 overview and diurnal cycle results for 2002, *Global Biogeochem. Cycles*, 22, GB3009,
1141 doi:10.1029/2007GB003050.
1142
1143 Li, K. F., B. Tian, D. E. Waliser, and Y. L. Yung (2010), Tropical mid-tropospheric CO₂ variability
1144 driven by the Madden-Julian oscillation, *PNAS*, 107(45), 19171- 19175, doi:10.1073/pnas.1008222107.
1145
1146 Le Quéré, C. et al. (2013), The global carbon budget 1959–2011, *Earth Syst. Sci. Data*, 5, 165-185,
1147 doi:10.5194/essd-5-165-2013.
1148
1149 Lin, S.-J. (2004), A “vertically Lagrangian” finite-volume dynamical core for global models, *Monthly*
1150 *Weather Review*, 132(10):2293-2307.
1151
1152 Liu, J., I. Fung, E. Kalnay, J.-S. Kang, E. T. Olsen, and L. Chen (2012), Simultaneous assimilation of
1153 AIRS Xco₂ and meteorological observations in a carbon climate model with an ensemble Kalman filter,
1154 *Journal of Geophysical Research*, 117, D05309, doi:10.1029/2011JD016642.
1155
1156 Lock, A. P., A. R. Brown, M. R. Bush, G. M. Martin, and R. N. B. Smith (2000), A New Boundary
1157 Layer Mixing Scheme. Part I: Scheme Description and Single-Column Model Tests, *Monthly Weather*
1158 *Review*, 128, 3187–3199.
1159
1160 Los, S. O, G. J. Collatz, P. J. Sellers, C. M. Malmstrom, N. H. Pflack, R. S. DeFries L. Bounoua, M. T.
1161 Parris, C. J. Tucker, and D. A. Dazlich (2000), A global 9-yr biophysical land surface dataset from
1162 NOAA AVHRR data, *Journal of Hydrometeorology*, 1, 183-199.
1163
1164 Los, S. O. (2013), Analysis of trends in fused AVHRR and MODIS NDVI data for 1982–2006:
1165 Indication for a CO₂ fertilization effect in global vegetation, *Global Biogeochemical Cycles*, 27, 318–
1166 330, doi:10.1002/gbc.20027.
1167
1168 Losch, M., D. Menemenlis, P. Heimbach, J. Campin, and C. Hill (2010), On the formulation of sea-ice
1169 models. Part 1: Effects of different solver implementations and parameterizations, *Ocean Modelling*, 33,
1170 129–144.
1171
1172 Maksyutov, S., et al. (2013), Regional CO₂ flux estimates for 2009–2010 based on GOSAT and
1173 ground-based CO₂ observations, *Atmospheric Chemistry and Physics*, 13, 9351-9373, doi:10.5194/acp-
1174 13-9351-2013.
1175
1176 Mann, G. W., et al. (2012), Intercomparison of modal and sectional aerosol microphysics
1177 representations within the same 3-D global chemical transport model, *Atmospheric Chemistry and*
1178 *Physics*, 12, 4449-4476, doi:10.5194/acp-12-4449-2012.

1179
1180 Marshall, J., A. Adcroft, C. Hill, L. Perelman, and C. Heisey (1997a), A finite-volume, incompressible
1181 Navier-Stokes model for studies of the ocean on parallel computers, *J. Geophys. Res.*, *102*, 5753–5766.
1182
1183 Marshall, J., C. Hill, L. Perelman, and A. Adcroft (1997b), Hydrostatic, quasi-hydrostatic and non-
1184 hydrostatic ocean modeling, *Journal of Geophysical Research*, *102*, 5733–5752.
1185
1186 Menemenlis, D., C. Hill, A. Adcroft, J. Campin, B. Cheng, B. Ciotti, I. Fukumori, P. Heimbach, C.
1187 Henze, A. Koehl, T. Lee, D. Stammer, J. Taft, and J. Zhang (2005a), NASA supercomputer improves
1188 prospects for ocean climate research, *Eos Trans. AGU*, *86*, 89, 95–96.
1189
1190 Menemenlis, D., I. Fukumori, and T. Lee (2005b), Using Green's functions to calibrate an ocean
1191 general circulation model, *Monthly Weather Review*, *133*, 1224–1240.
1192
1193 Menemenlis, D., J. Campin, P. Heimbach, C. Hill, T. Lee, A. Nguyen, M. Schodlock, and H. Zhang
1194 (2008), ECCO2: High resolution global ocean and sea ice data synthesis, *Mercator Ocean Quarterly*
1195 *Newsletter*, *31*, 13–21.
1196
1197 Messerschmidt, J., et al. (2012), Automated ground-based remote sensing measurements of greenhouse
1198 gases at the Białystok site in comparison with collocated in situ measurements and model data,
1199 *Atmospheric Chemistry and Physics*, *12*(15), 6741–6755, doi:10.5194/acp-12-6741-2012.
1200
1201 Moorthi S., and M. J. Suarez (1992), Relaxed Arakawa–Schubert: A parameterization of moist
1202 convection for general circulation models, *Monthly Weather Review*, *120*, 978–1002.
1203
1204 Mu, M., et al. (2011), Daily and 3-hourly variability in global fire emissions and consequences for
1205 atmospheric model predictions of carbon monoxide, *Journal of Geophysical Research*, *116*, D24303,
1206 doi:10.1029/2011JD016245.
1207
1208 Najjar, R. G., et al. (2007), Impact of circulation on export production, dissolved organic matter, and
1209 dissolved oxygen in the ocean: Results from Phase II of the Ocean Carbon-cycle Model
1210 Intercomparison Project (OCMIP-2), *Global Biogeochemical Cycles*, *21*, GB3007,
1211 doi:10.1029/2006GB002857.
1212
1213 Novelli, P. C., L. P. Steele, and P. P. Tans (1992), Mixing ratios of carbon monoxide in the troposphere,
1214 *Journal of Geophysical Research*, *97*, 20,731–20,750.
1215
1216 O'Dell, C. W. et al. (2012), The ACOS CO₂ retrieval algorithm – Part 1: Description and validation
1217 against synthetic observations, *Atmos. Meas. Tech.*, *5*, 99–121, doi:10.5194/amt-5-99-2012.
1218
1219 Olsen, S. C., and J. T. Randerson (2004), Differences between surface and column atmospheric CO₂
1220 and implications for carbon cycle research, *Journal of Geophysical Research*, *109*, D02301,
1221 doi:10.1029/2003JD003968.
1222
1223 Orr, J. C., et al. (2001), Estimates of anthropogenic carbon uptake from four three-dimensional global
1224 ocean models, *Global Biogeochem. Cycles*, *15*(1), 43–60, doi:10.1029/2000GB001273.
1225
1226 Ott, L., B. Duncan, S. Pawson, P. Colarco, M. Chin, C. Randles, T. Diehl, and J. E. Nielsen (2010),
1227 Influence of the 2006 Indonesian Biomass Burning Aerosols on Tropical Dynamics Studied with the

1228 GEOS-5 AGCM, *Journal of Geophysical Research*, *115*. doi:10.1029/2009JD013181.
1229
1230 Ott, L., S. Pawson, and J. Bacmeister (2011), An analysis of the impact of convective parameter
1231 sensitivity on simulated global atmospheric CO distributions, *Journal of Geophysical Research*, *116*,
1232 D21310. doi: 10.1029/2011JD016077.
1233
1234 Piao, S. et al. (2013), Evaluation of terrestrial carbon cycle models for their response to climate
1235 variability and to CO₂ trends, *Global Change Biology*, *19*, 2117–2132. doi: 10.1111/gcb.12187.
1236
1237 Popova, E. E., A. Yool, A. C. Coward, F. Dupont, C. Deal, S. Elliott, E. Hunke, M. Jin, M. Steele, and
1238 J. Zhang (2012), What controls primary production in the Arctic Ocean? Results from an
1239 intercomparison of five general circulation models with biogeochemistry, *Journal of Geophysical
1240 Research*, *117*, C00D12, doi:10.1029/2011JC007112.
1241
1242 Potter, C. S., J. T. Randerson, C. B. Field, P. A. Matson, P. M. Vitousek, H. A. Mooney, and S. A.
1243 Klooster (1993), Terrestrial ecosystem production – A process model based on global satellite
1244 and surface data, *Global Biogeochemical Cycles*, *7*, 811–841.
1245
1246 Potter, C., S. Klooster, A. Huete, and V. Genovese (2007), Terrestrial carbon sinks for the United
1247 States predicted from MODIS satellite data and ecosystem modeling, *Earth Interactions*, *11*, 1-21.
1248
1249 Potter, C., S. Klooster, V. Genovese, C. Hiatt, S. Boriah, V. Kumar, V. Mithal, and A. Garg (2012),
1250 Terrestrial ecosystem carbon fluxes predicted from MODIS satellite data and large-scale disturbance
1251 modeling, *International Journal of Geosciences*, doi:10.4236/ijg.2012.
1252
1253 Raczka, B. M. et al. (2013), Evaluation of continental carbon cycle simulations with North American
1254 flux tower observations, *Ecological Monographs*, *83*:4, 531-556.

1255 Randerson J. T., M. V. Thompson, and C. M. Malmstrom (1996), Substrate limitations for heterotrophs:
1256 Implications for models that estimate the seasonal cycle of atmospheric CO₂, *Global Biogeochemical
1257 Cycles*, *10*, 585-602.
1258
1259 Randerson, J. T. G. J. Collatz, J. E. Fessenden, A. D. Munoz, C. J. Still, J. A. Berry, I. Y. Fung, N.
1260 Suits, and A. S. Denning (2002), A possible global covariance between terrestrial gross primary
1261 production and ¹³C discrimination: Consequences for the atmospheric ¹³C budget and its response to
1262 ENSO, *Global Biogeochemical Cycles*, *16*, doi: 10.1029/2001GB001845.
1263
1264 Randerson, J. T. et al. (2009), Systematic assessment of terrestrial biogeochemistry in coupled climate–
1265 carbon models, *Global Change Biology*, *15*, 2462–2484. doi: 10.1111/j.1365-2486.2009.01912.
1266
1267 Reinecker, M. M., Suarez, M. J., Todling, R., et al. (2008), The GEOS-5 Data Assimilation System–
1268 Documentation of Versions 5.0.1, 5.1.0, and 5.2.0, Tech. Rep. 104606 V27, NASA.
1269
1270 Rienecker, M. M., et al. (2011), MERRA: NASA’s Modern-Era Retrospective Analysis for Research
1271 and Applications, *Journal of Climate*, *24*, 3624–3648.
1272
1273 Ruzmaikin, A., H. H. Aumann, and T. S. Pagano (2012), Patterns of CO₂ Variability from Global
1274 Satellite Data, *Journal of Climate*, doi: 10.1175/JCLI-D-11-00223.1
1275

1276 Schaefer, K., et al. (2012), A model-data comparison of gross primary productivity: Results from the
1277 North American Carbon Program site synthesis, *Journal of Geophysical Research*, *117*, G03010, doi:
1278 10.1029/2012JG001960.

1279

1280 Schwalm, C. R. et al. (2010), A model-data intercomparison of CO₂ exchange across North America:
1281 Results from the North American Carbon Program site synthesis, *Journal of Geophysical Research*,
1282 *115*, G00H05, doi:10.1029/2009JG001229.

1283

1284 Sentman, L. T., E. Shevliakova, R. J. Stouffer, and S. Malyshev (2011), Time Scales of Terrestrial
1285 Carbon Response Related to Land-Use Application: Implications for Initializing an Earth System
1286 Model. *Earth Interact.*, *15*, 1–16. doi: <http://dx.doi.org/10.1175/2011EI401.1>.

1287 Takagi, H., et al. (2011), On the Benefit of GOSAT Observations to the Estimation of Regional CO₂
1288 Fluxes, *SOLA*, 2011, Vol. 7, 161-164, doi:10.2151/sola.2011-041.

1289

1290 Takahashi, T., et al. (2002), Global sea-air CO₂ flux based on climatological surface ocean pCO₂, and
1291 seasonal biological and temperature effects, *Deep-Sea Res. II*, *49*, 1601-1622.

1292

1293 Takahashi, T., et al. (2009). Climatological mean and decadal changes in surface ocean pCO₂, and net
1294 sea-air CO₂ flux over the global oceans, *Deep-Sea Res. II*, *56*, 554-577.

1295

1296 Tans, P. P., I. Y. Fung, and T. Takahashi (1990), Observational constraints on the global atmospheric
1297 CO₂ budget, *Science*, *247*, 1431-1438.

1298

1299 Tucker, C. J., J. E. Pinzon, M. E. Brown, D. A. Slayback, E. W. Pak, R. Mahoney, E. F. Vermote, and
1300 N. El Saleous (2005), An extended AVHRR 8-km NDVI dataset compatible with MODIS and SPOT
1301 vegetation NDVI data, *International Journal of Remote Sensing*, *26*(20), 4485-4498.

1302

1303 van der Werf, G. R., J. T. Randerson, G. J. Collatz, L. Giglio, P. S. Kasibhatla, A. F. Arellano, S. C.
1304 Olsen, and E. S. Kasischke (2004), Continental-scale partitioning of fire emissions during the 1997 to
1305 2001 El Nino/La Nina period, *Science*, *303*, 73-76.

1306

1307 van der Werf G. R., J. T. Randerson, L. Giglio, G. J. Collatz, P. S. Kasibhatla, and A. F. Arellano
1308 (2006), Interannual variability of global biomass burning emissions from 1997 to 2004, *Atmospheric
1309 Chemistry and Physics*, *6*, 3423-3441.

1310

1311 van der Werf G. R., J. T. Randerson, L. Giglio, G. J. Collatz, M. Mu, P. S. Kasibhatla, D. C. Morton, R.
1312 S. DeFries, Y. Jin, and T. T. van Leeuwen (2010), Global fire emissions and the contribution of
1313 deforestation, agriculture, and peat fires (1997-2009), *Atmospheric Chemistry and Physics*, *10*, 11707-
1314 11735.

1315

1316 Vanuytrecht, E., D. Raes, and P. Willems (2011), Considering sink strength to model crop production
1317 under elevated atmospheric CO₂, *Agricultural and Forest Meteorology*, *151*, 1753-1762, ISSN 0168-
1318 1923, <http://dx.doi.org/10.1016/j.agrformet.2011.07.011>.

1319

1320 Wang, H., D. J. Jacob, M. Kopacz, D. B. A. Jones, P. Suntharalingam, J. A. Fisher, R. Nassar, and S.
1321 Pawson (2009), Error Correlation Between CO₂ and CO as a Constraint for CO₂ Flux Inversions Using
1322 Satellite Data, *Atmos. Chem. Phys.*, *9*, 7313-7323. doi:10.5194/acp-9-7313-2009.

1323

1324 Wanninkhof, R. (1992), Relationship between wind speed and gas exchange
1325 over the ocean, *J. Geophys. Res.*, 97(C5), 7373–7382.
1326

1327 Winderlich, J., C. Gerbig, O. Kolle, and M. Heimann (2014), Inferences from CO₂ and CH₄
1328 concentration profiles at the Zotino Tall Tower Observatory (ZOTTO) on regional summertime
1329 ecosystem fluxes, *Biogeosciences*, 11, 2055-2068, doi:10.5194/bg-11-2055-2014.
1330

1331 Wofsy, S. C., et al. (2011), HIAPER Pole-to-Pole Observations (HIPPO): fine-grained, global-scale
1332 measurements of climatically important atmospheric gases and aerosols, *Proc. Roy. Soc. A*, 369, 2073-
1333 2086, doi:10.1098/rsta.2010.0313.
1334

1335 Wunch, D., et al. (2010), Calibration of the Total Carbon Column Observing Network using aircraft
1336 profile data, *Atmospheric Measurement Techniques*, 3(5), 1351-1362, doi:10.5194/amt-3-1351-2010.
1337

1338 Wunch, D., G. C. Toon, J.-F. L. Blavier, R. A. Washenfelder, J. Notholt, B. J. Connor, D. W. T.
1339 Griffith, V. Sherlock, and P. O. Wennberg (2011a), The total carbon column observing network,
1340 *Philosophical Transactions of the Royal Society - Series A: Mathematical, Physical and Engineering*
1341 *Sciences*, 369(1943), 2087-2112, doi:10.1098/rsta.2010.0240.

1342 Wunch, D., et al. (2011b), A method for evaluating bias in global measurements of CO₂ total columns
1343 from space, *Atmospheric Chemistry and Physics*, 11(23), 12317-12337, doi:10.5194/acp-11-12317-
1344 2011.
1345

1346 Wunsch, C. and P. Heimbach (2007), Practical global ocean state estimation, *Physica D*, 230, 197–208.
1347

1348 Yokota, T., Y. Yoshida, N. Eguchi, Y. Ota, T. Tanaka, H. Watanabe, and S. Maksyutov (2009), Global
1349 Concentrations of CO₂ and CH₄ Retrieved from GOSAT: First Preliminary Results, *SOLA*, 5, 160-163,
1350 2009.
1351

1352 Zhu, Y. and R. Gelaro (2008), Observation Sensitivity Calculations Using the Adjoint of the Gridpoint
1353 Statistical Interpolation (GSI) Analysis System, *Mon. Wea. Rev.*, 136, 335-351.
1354

1355 **Tables**

1356

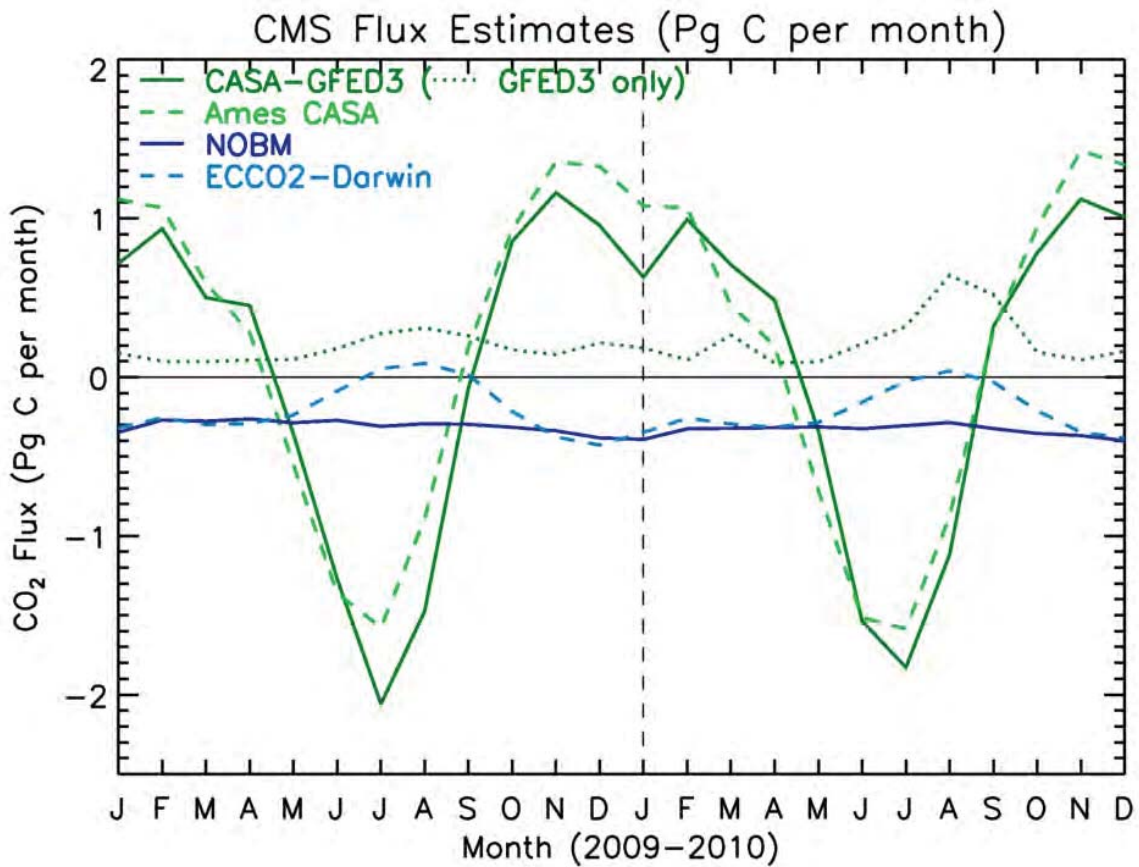
1357 **Table 1.** Combinations of surface (land biosphere and oceanic) fluxes used in the model computations
1358 in the Flux-Pilot Project. Fossil fuel fluxes were identical in all four combinations.

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Flux Combination	Land Biosphere	Ocean	Total Land and Ocean Flux (Pg C yr⁻¹)	Annual Mean Growth Rate at MBL sites (ppmv yr⁻¹)
CG-NO	CASA/GFED-3	NOBM	-3.4 (2009), -2.8 (2010)	2.7
AC-NO	Ames CASA	NOBM	-1.2 (2009), -1.9 (2010)	3.5
CG-ED	CASA/GFED-3	ECCO2-Darwin	-2.1 (2009), -1.4 (2010)	3.3
AC-ED	Ames CASA	ECCO2-Darwin	0.1 (2009), -0.5 (2010)	4.1

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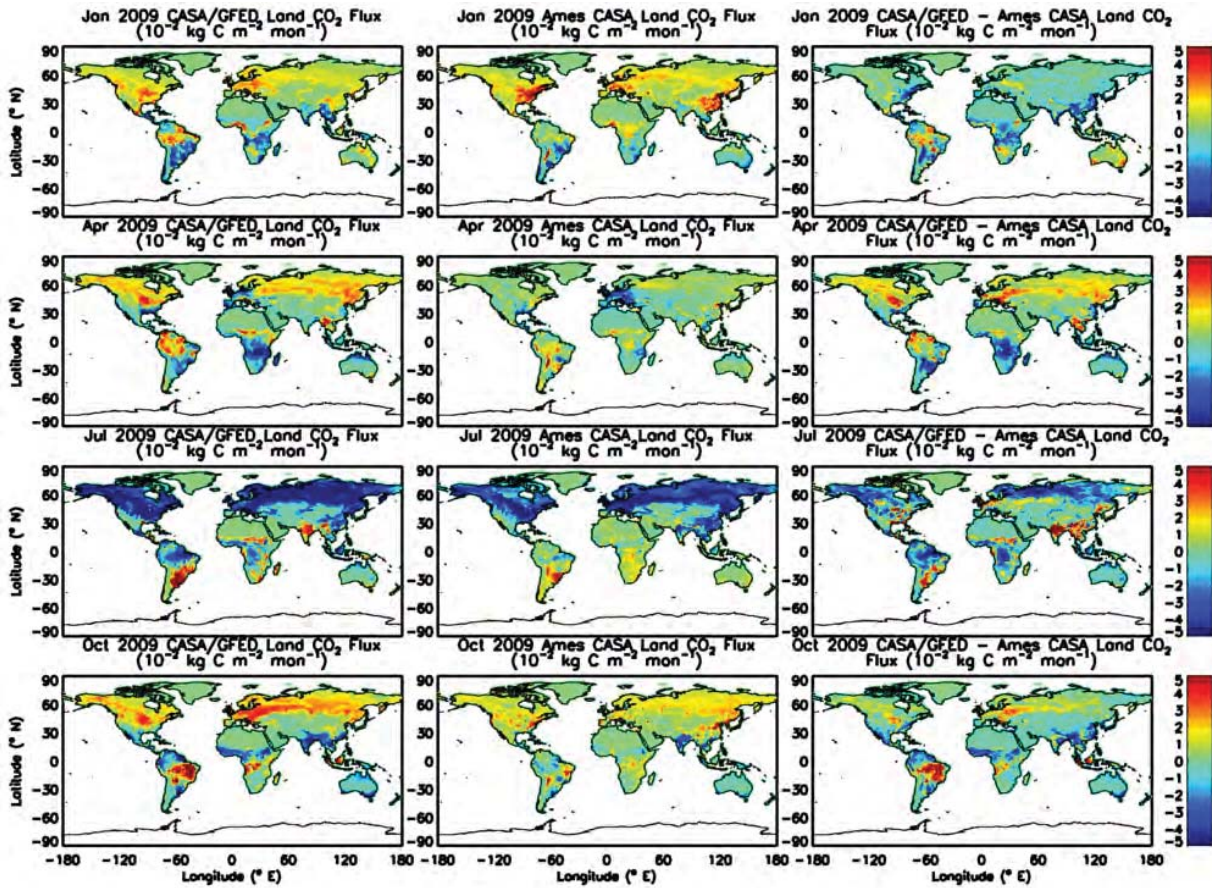
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1363 **Figure 1.** Global land and ocean flux estimates produced as part of NASA's CMS FPP for 2009 and
 1364 2010 in Pg carbon per month. Dark green solid line indicates combined NEP and biomass burning
 1365 fluxes computed from by the CASA-GFED3 model (dotted dark green line shows biomass burning
 1366 contribution only). Light green dashed line indicates NEP from the Ames CASA model. Blue lines
 1367 show ocean flux estimates from NOBM (dark blue solid) and ECCO2-Darwin (light blue dashed).

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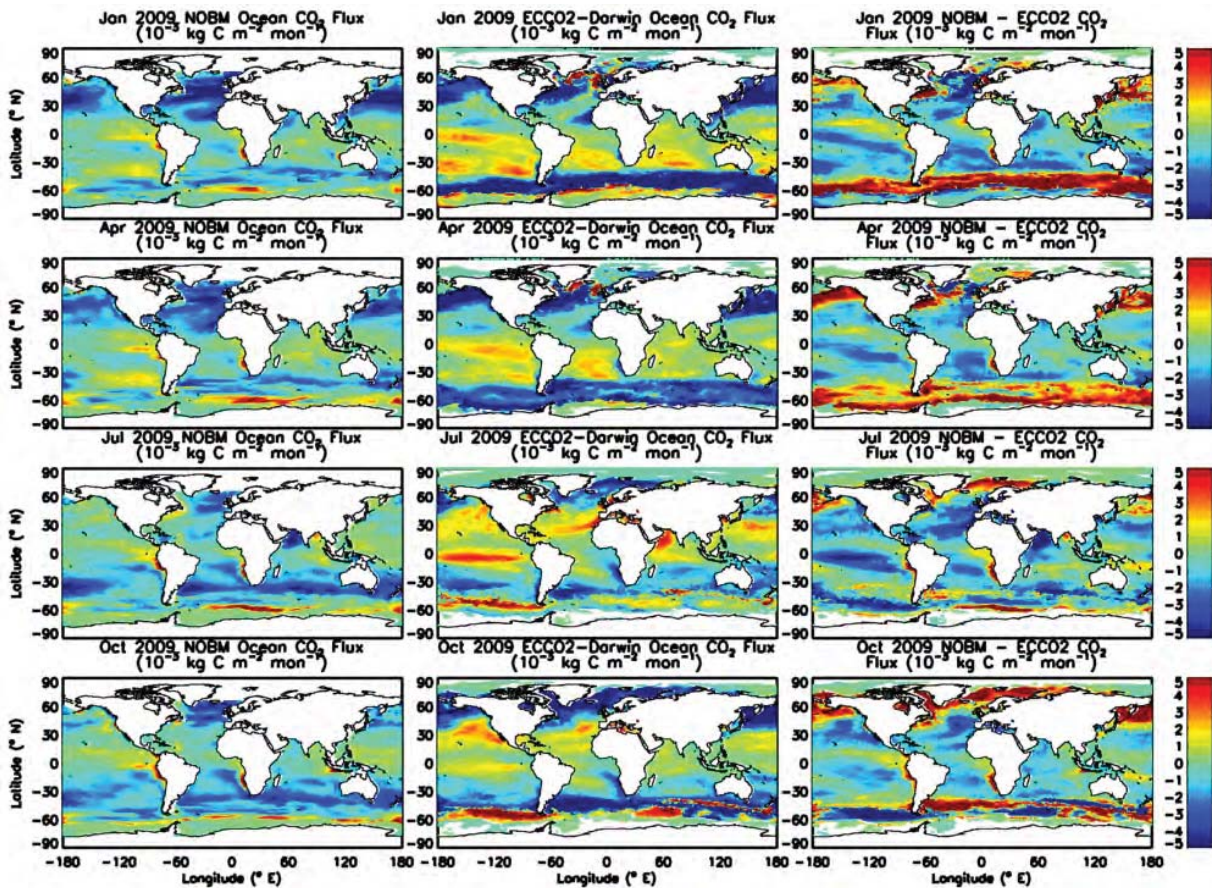
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1372 **Figure 2.** Monthly land flux estimates from the CASA-GFED3 (left) and Ames CASA (middle)
 1373 models and their difference (CASA-GFED3 minus Ames CASA; right) for January, April, July and
 1374 October of 2009. Units are 10⁻² kg Carbon per m² per month.

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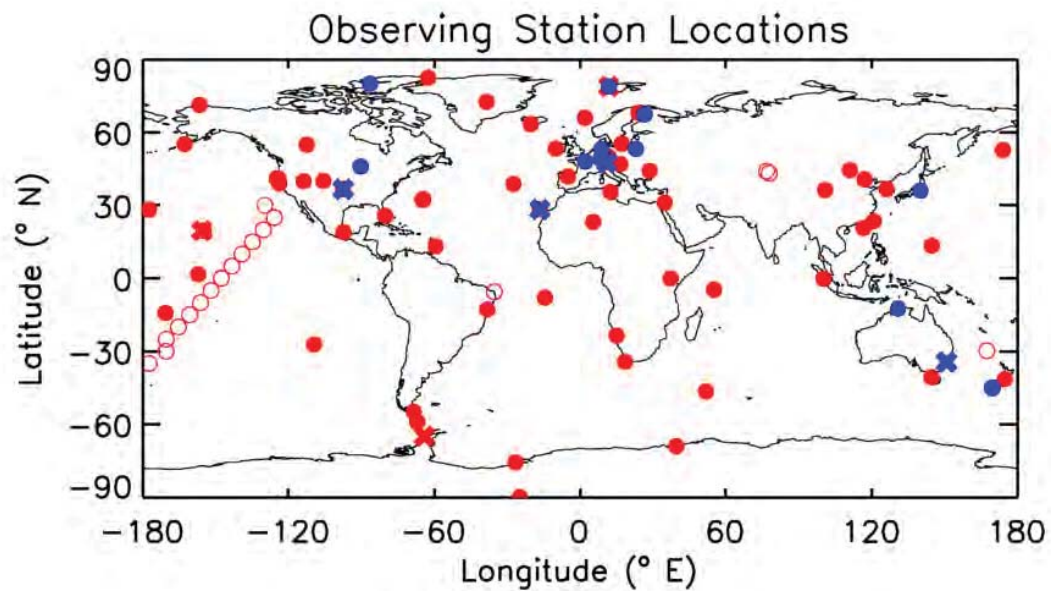
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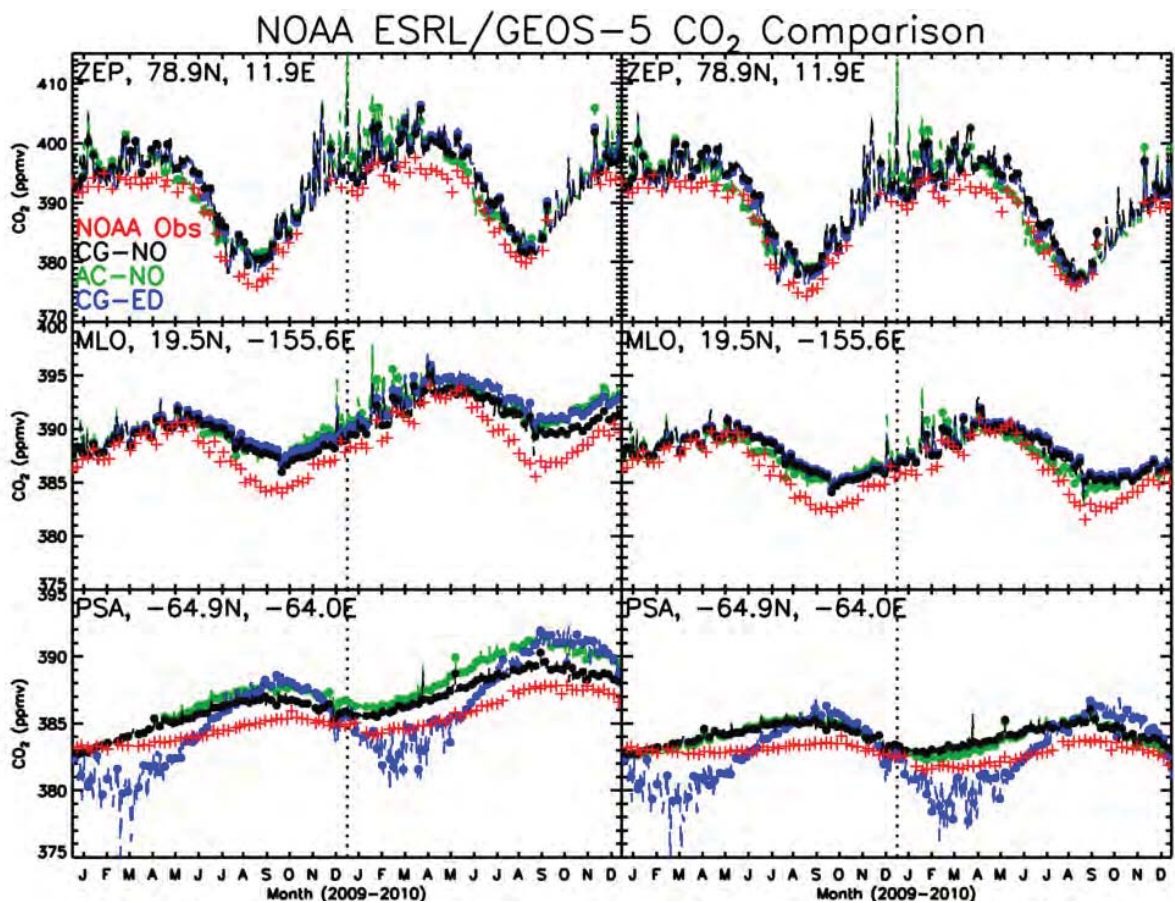
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 1379 **Figure 3.** Monthly ocean flux estimates from the NOBM (left) and ECCO2-Darwin (middle) models
 1380 and their difference (NOBM minus ECCO2-Darwin; right) for January, April, July and October of
 1381 2009. Units are 10⁻³ kg Carbon per m² per month (note that the ocean fluxes shown here are an order of
 1382 magnitude smaller than the land fluxes shown in Figure 2).

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1391 **Figure 4.** Locations of NOAA (red) and TCCON (blue) observing stations in operation during the
 1392 study period. Open red circles indicate stations that collected an insufficient amount of data in 2009-
 1393 2010 and thus were not included in Figures 6-8. Red (blue) 'x' marks indicate NOAA (TCCON)
 1394 stations shown in Figure 5 (12).



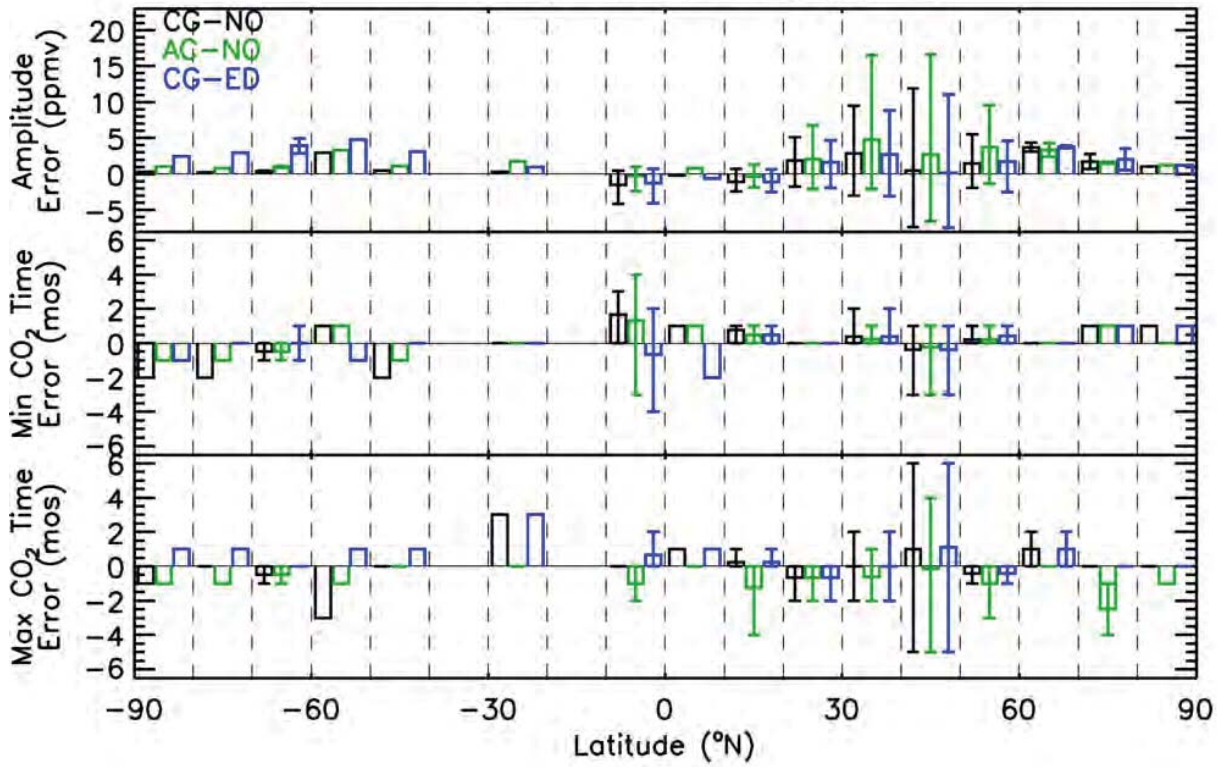
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1396 **Figure 5.** Left panels show simulated and observed CO₂ at the Mauna Loa, Ny-Alesund, and Palmer
 1397 Station. Right panels show simulated and observed CO₂ at the same locations when differences in
 1398 atmospheric growth rate are removed. Red indicates NOAA ESRL observations while black, green, and
 1399 blue indicates GEOS-5 simulated mixing ratios assuming flux scenarios CG-NO, AC-NO, and CG-ED,
 1400 respectively (flux scenarios are described in detail in Table 1).

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Error in the 2010 Seasonal Cycle
of Monthly Surface CO₂ (ppmv)



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1410 **Figure 7.** Evaluation of the simulations' ability to represent the seasonal cycle of CO₂ observed at
 1411 NOAA surface stations during 2010. The top plot shows the mean error in the magnitude of the
 1412 seasonal cycle calculated over 10 degree latitude bins and for flux combinations CG-NO, AC-NO, and
 1413 CG-ED. The middle (bottom) plot shows the mean error in the month during which the minimum
 1414 (maximum) in the seasonal cycle occurs. Boxes identify mean values while vertical lines indicate
 1415 maximum and minimum values within each 10 degree latitude range.

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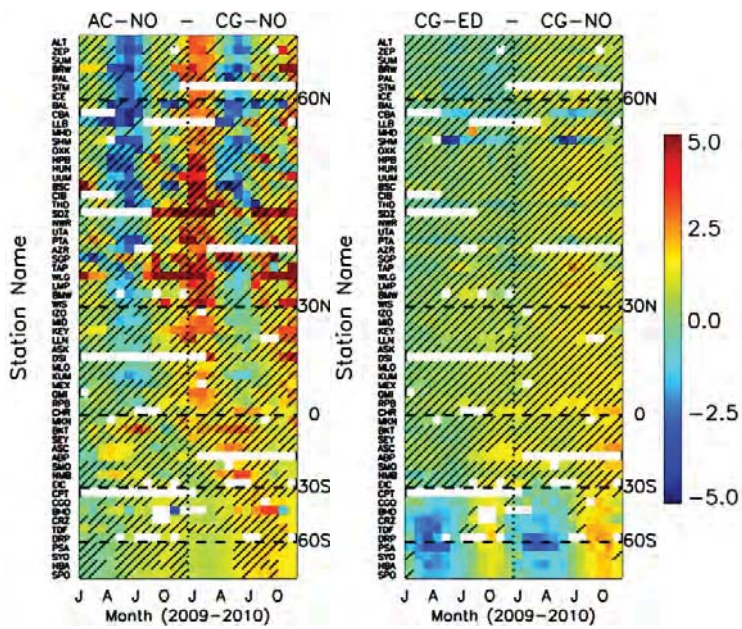
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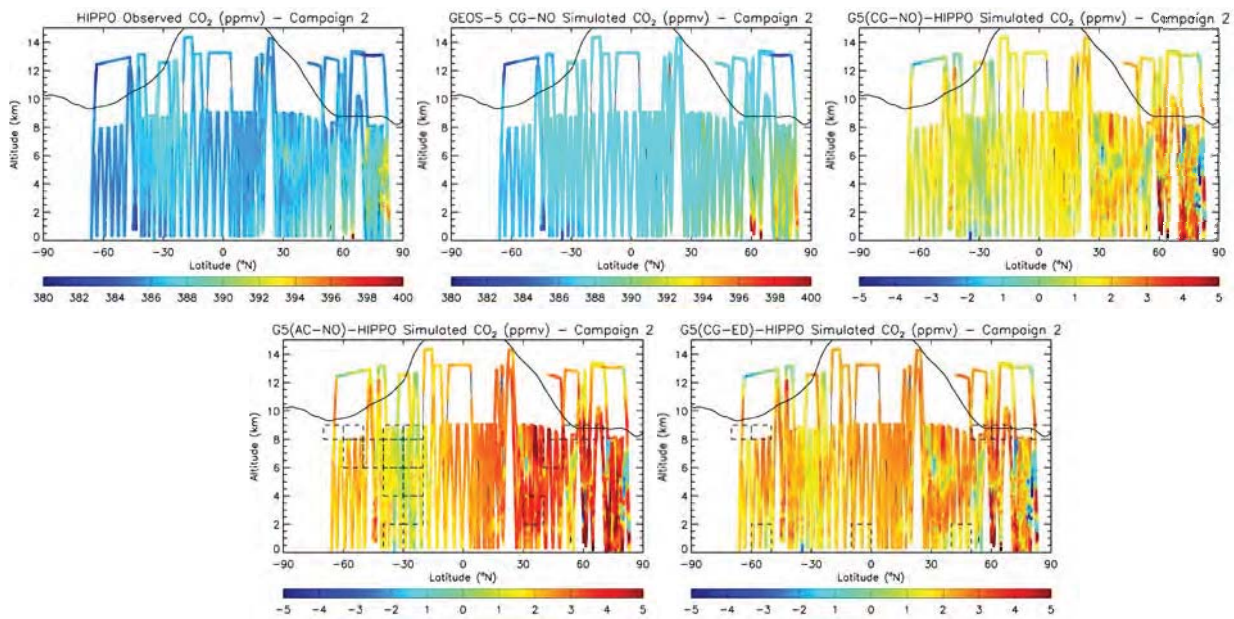
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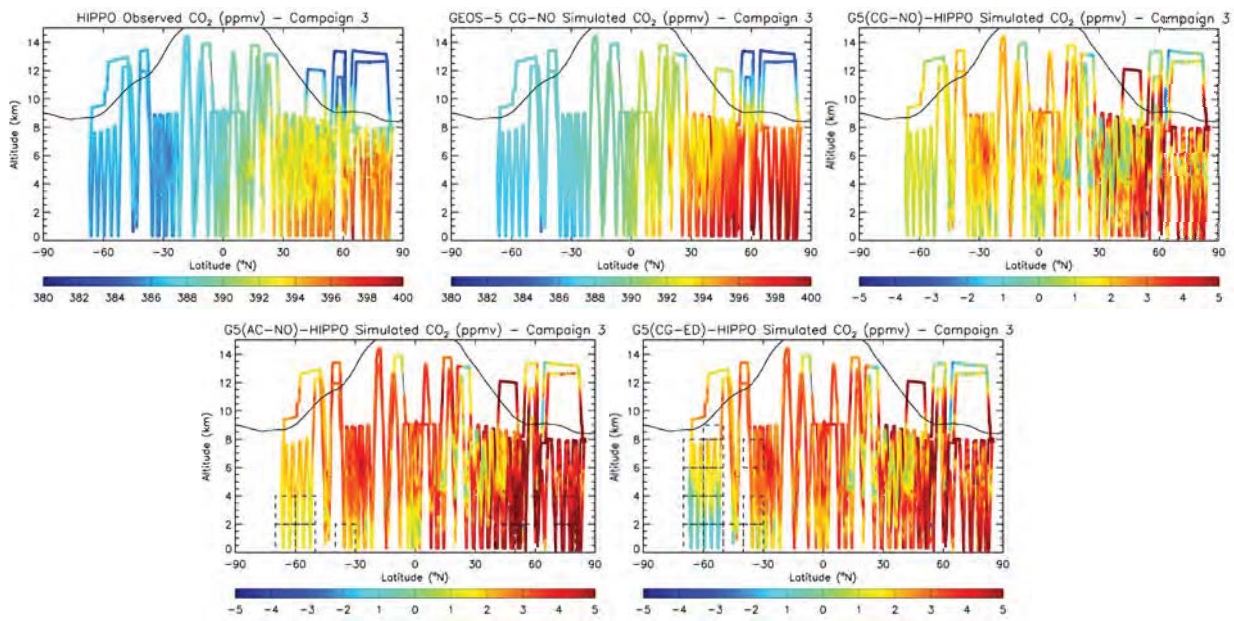
1424 **Figure 8.** Differences in monthly mean CO₂ mixing ratio at NOAA surface stations due to differing
 1425 land flux estimates (left) and ocean flux estimates (right). Diagonal bars indicate instances in which
 1426 flux differences would not be statistically significant from observation variability.

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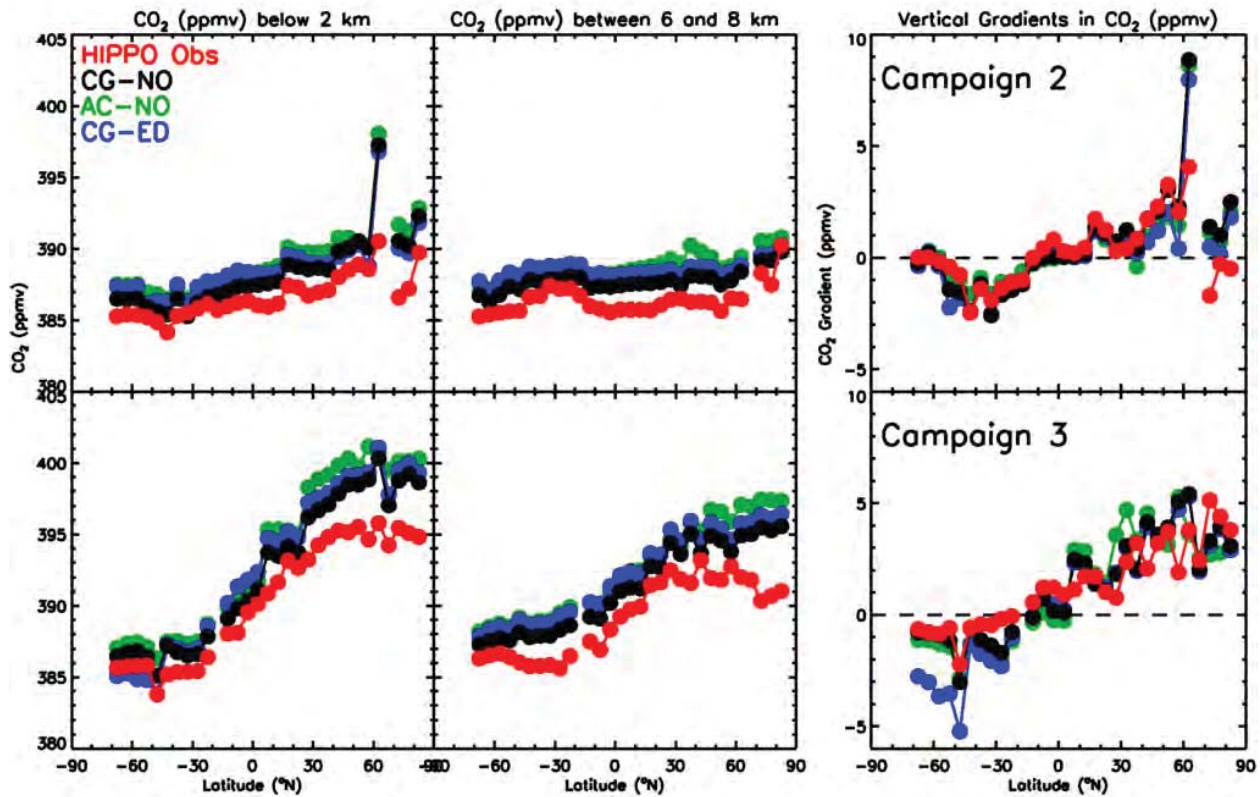
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Figure 9. Comparison of observed and simulated CO₂ mixing ratios (top) during HIPPO-2 in October and November, 2009 show HIPPO observations (left), GEOS-5 using flux combination CG-NO (middle) and the simulated minus observed difference (right). Bottom plots show the difference between GEOS-5 and observations using alternate flux scenarios AC-NO (left) and CG-ED (right). Dashed boxes indicate regions where flux differences are significantly larger than the variance of observations.



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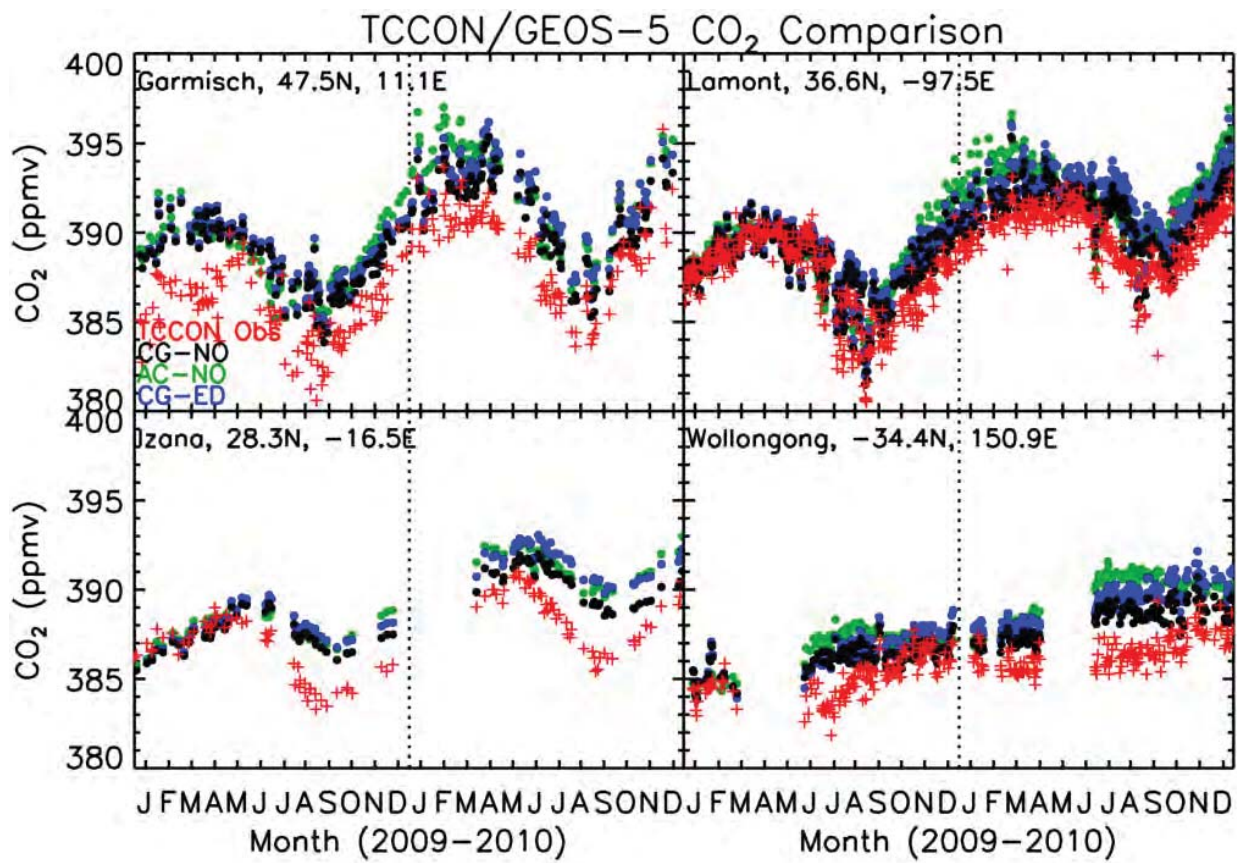
Figure 10. Comparison of observed and simulated CO₂ mixing ratios (top) during HIPPO-3 in March and April, 2010 show HIPPO observations (left), GEOS-5 using flux combination CG-NO (middle) and the simulated minus observed difference (right). Bottom plots show the difference between GEOS-5 and observations using alternate flux scenarios AC-NO (left) and CG-ED (right). Dashed boxes indicate regions where flux differences are significantly larger than the variance of observations.



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1460 **Figure 11.** Simulated and observed vertical gradients in in situ CO₂ derived from HIPPO data and
 1461 GEOS-5 model simulations (right). Gradients are calculated by binning observations and pseudo-data
 1462 into 5 latitude bins and calculating the difference between the mean mixing ratios below 2 km (left) and
 1463 between 6 and 8 km (middle).

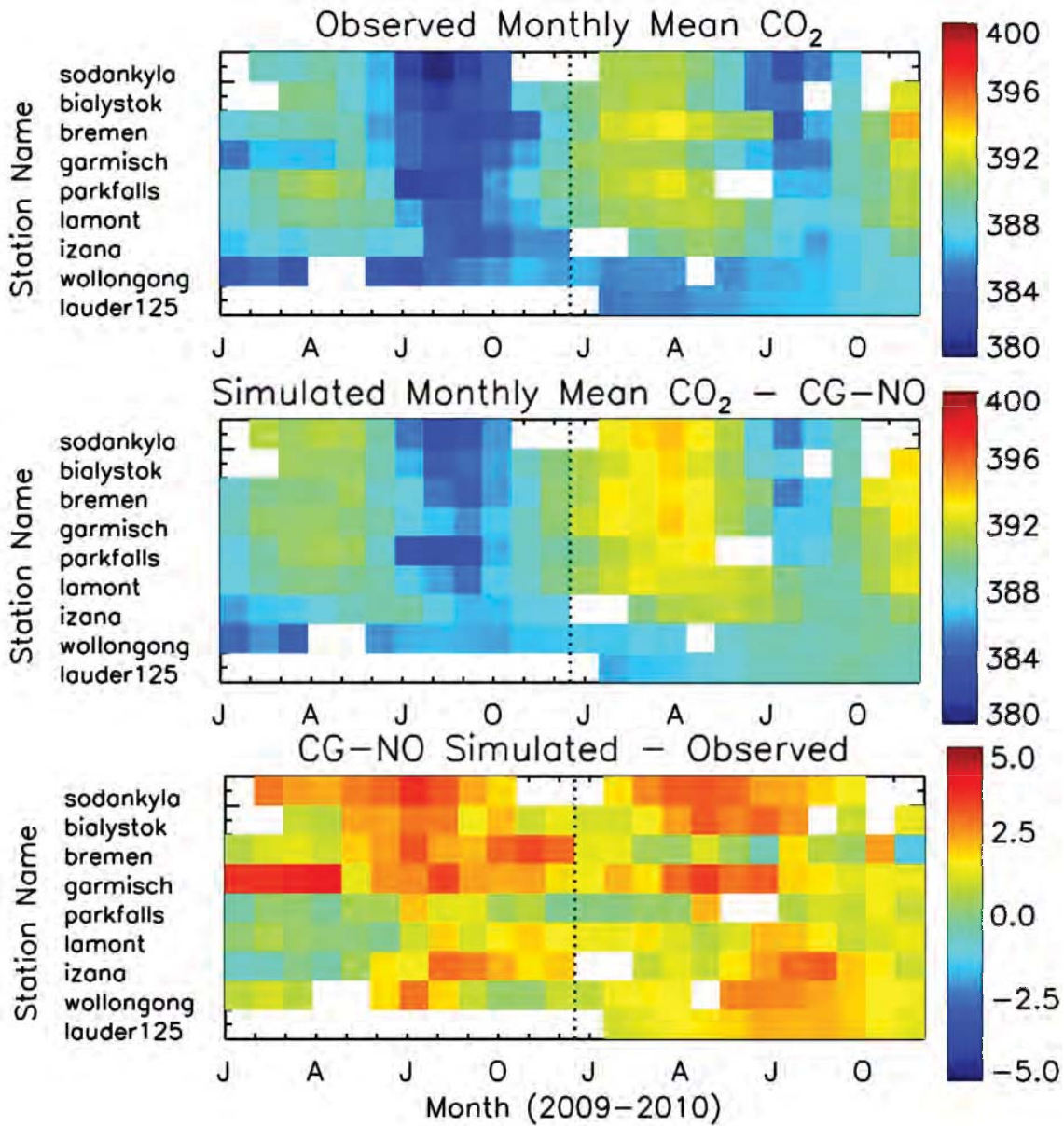
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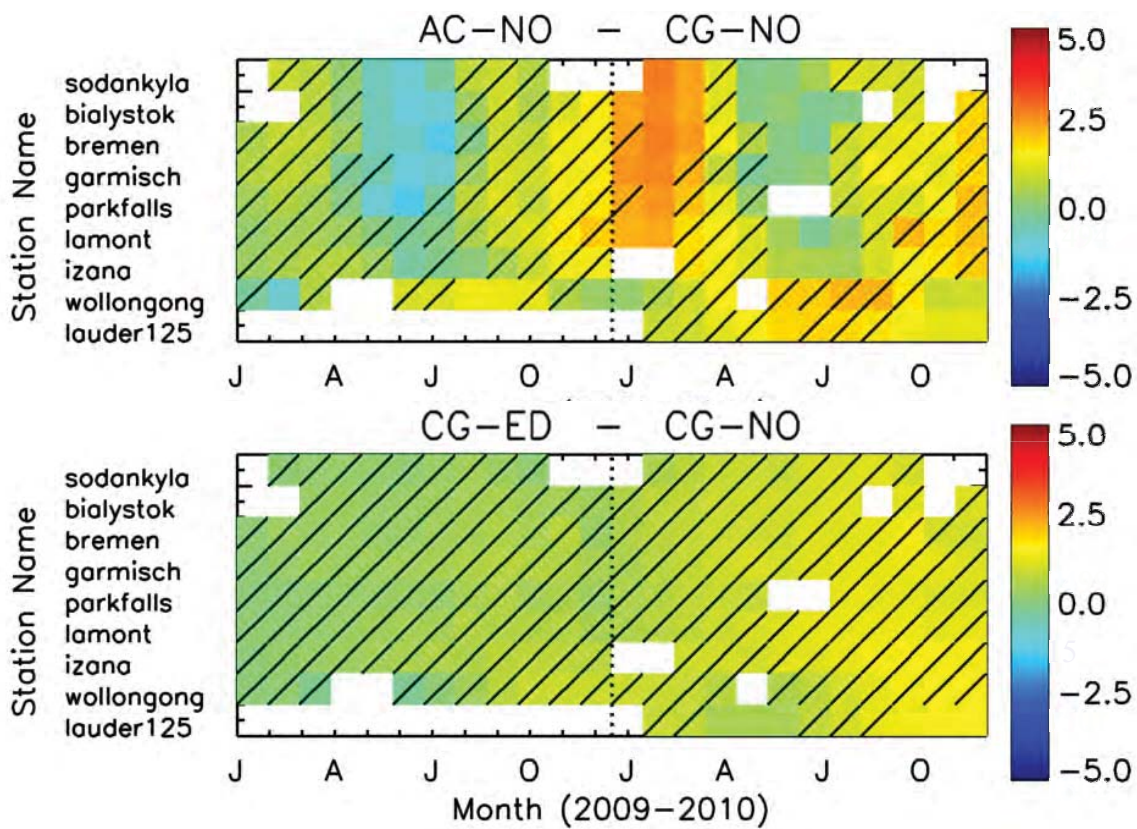
1466 **Figure 12.** Comparison of daily mean simulated and observed column CO₂ at Garmisch, Lamont,
1467 Izana, and Wollongong TCCON stations for 2009-2010.

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1487 **Figure 13.** Monthly mean CO₂ (ppmv) observed at TCCON stations during 2009-2010 (top) compared
 1488 with simulated CO₂ assuming flux combination CG-NO (middle). Bottom plot shows the model-
 1489 observation difference. Stations are oriented from north (top) to south (bottom) on all plots.

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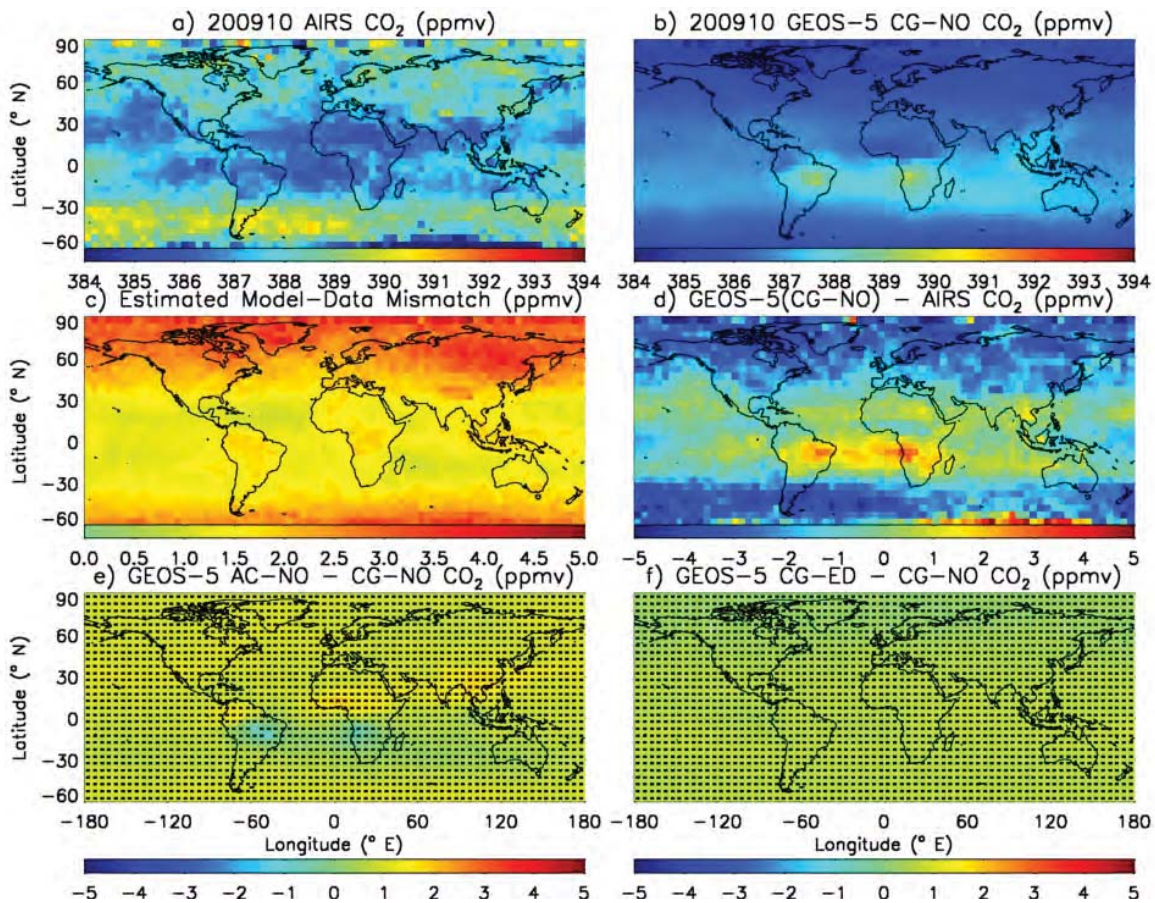
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1504 **Figure 14.** Differences in monthly mean CO₂ mixing ratio at TCCON stations due to land flux
 1505 estimates (top) and ocean flux estimates (bottom). Diagonal bars indicate instances in which flux
 1506 differences would not be statistically significant from observation variability.

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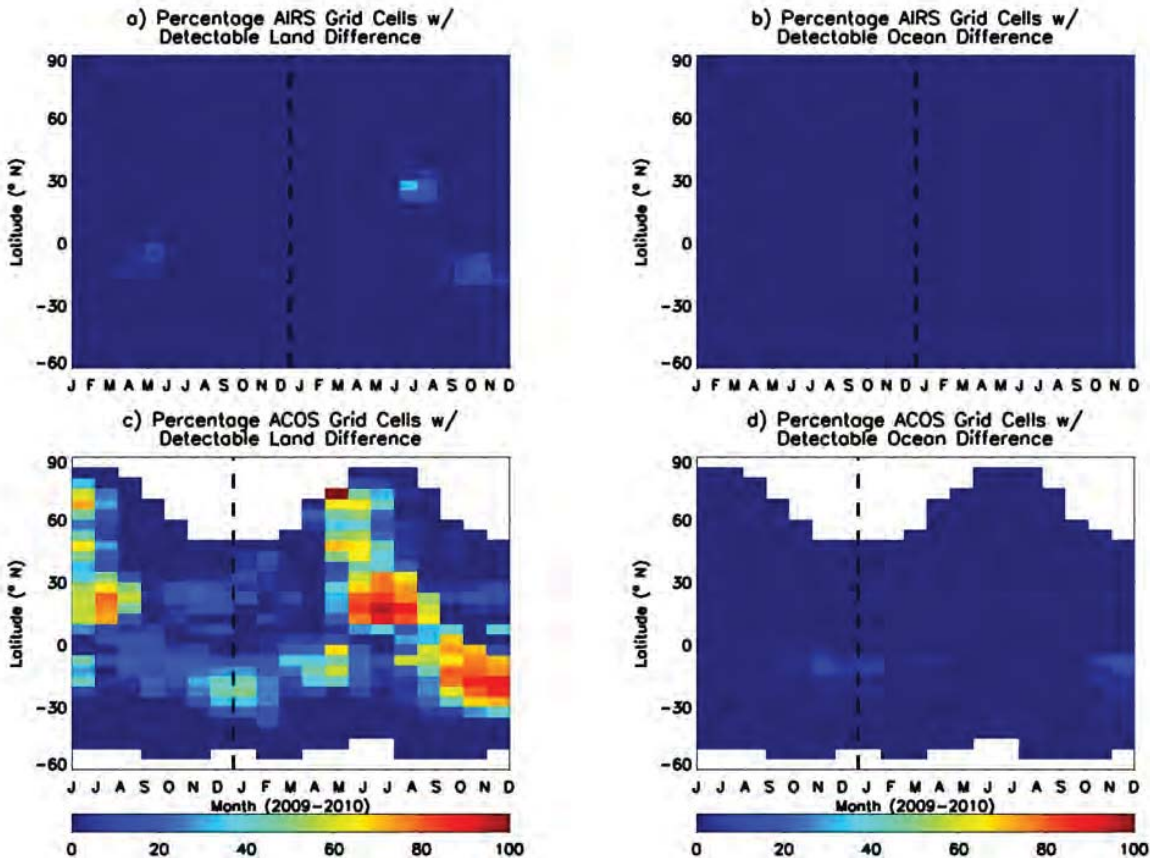
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1511 **Figure 15.** Comparison of AIRS observed mid-troposphere CO₂ mixing ratio with GEOS-5
 1512 simulations for October, 2009 for 5 degree latitude by 5 degree longitude grid boxes. The top panels
 1513 show AIRS monthly mean mixing ratio (left) and monthly mean GEOS-5 mixing ratios assuming flux
 1514 combination CG-NO sampled using AIRS pressure weighting functions (right). Middle panels show
 1515 the estimated model data mismatch of AIRS observations for each grid box (left) and GEOS-5 CO₂
 1516 minus AIRS (right). Bottom panels show the difference between GEOS-5 simulated CO₂ for flux
 1517 combinations AC-NO and CG-NO (left) and combinations CG-ED and CG-NO (right); grid cells
 1518 where these differences are not statistically significant at the 95% confidence level are indicated by
 1519 black lines.

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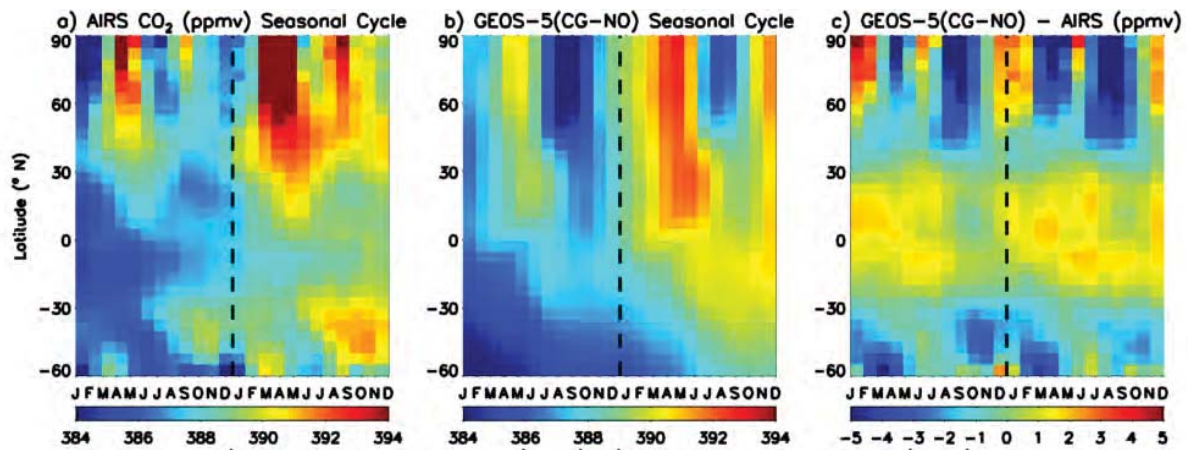


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1522 **Figure 16.** Percentage of 5 degree latitude by 5 degree longitude grid cells per month and per latitude
 1523 bin in which land (ocean) flux differences manifest as significantly different satellite CO₂ mixing ratios
 1524 re shown in right (left) panels. Top panels show the detectability of flux differences by AIRS while
 1525 bottom plots show results for ACOS GOSAT data.

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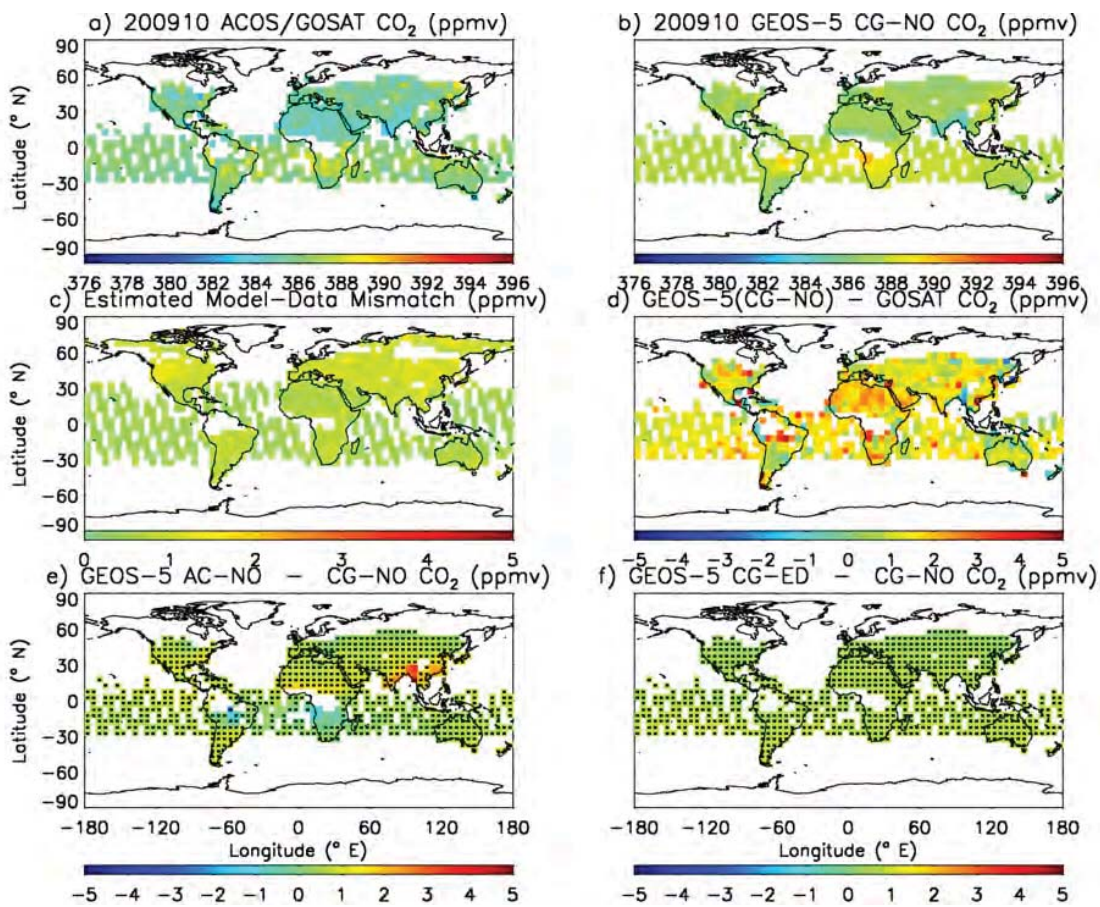
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1529 **Figure 17.** Seasonal cycle of (a) zonal mean AIRS observed mixing ratios, (b) zonal mean GEOS-5
 1530 (CG-NO) mixing ratios calculated with AIRS weighting functions, and (c) the difference between zonal
 1531 mean model results and observations. All units are ppmv.

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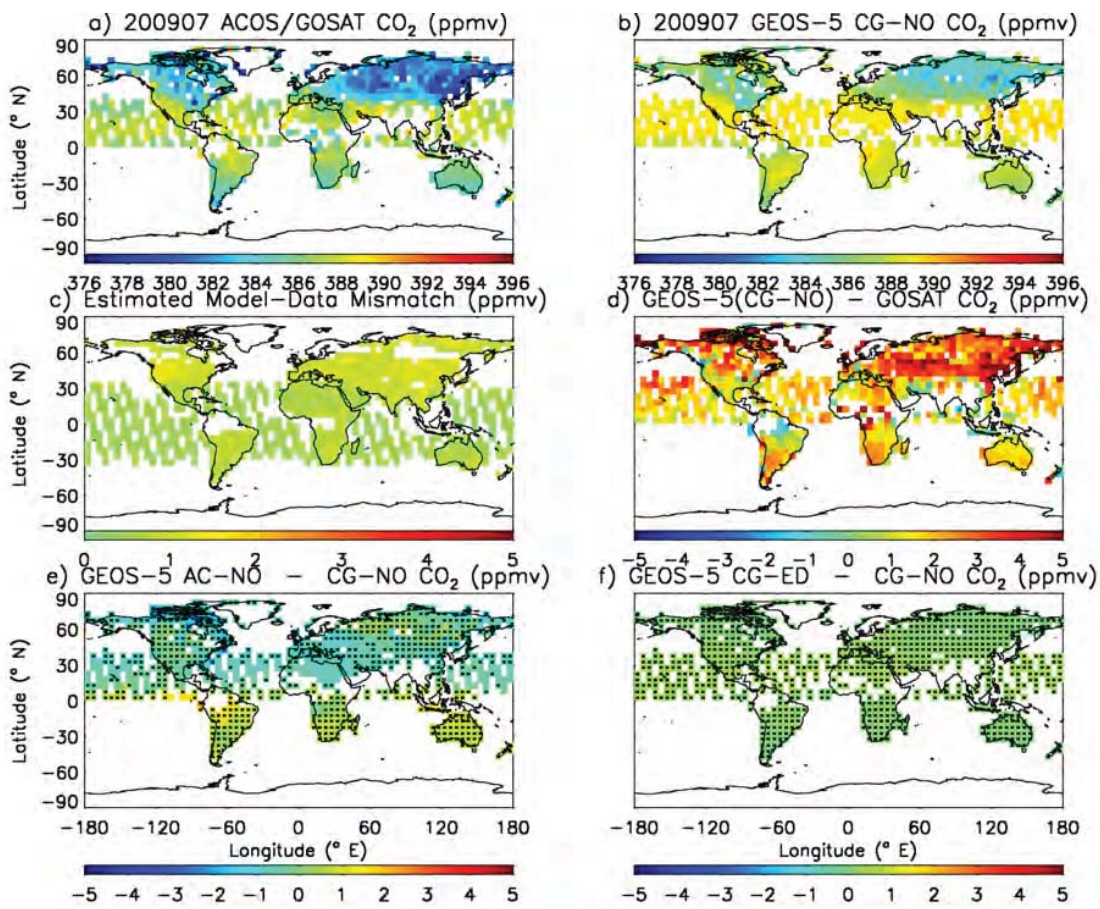


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1534 **Figure 18.** Same as Figure 14, but using ACOS retrievals of GOSAT observations and sampling
 1535 instead of AIRS.

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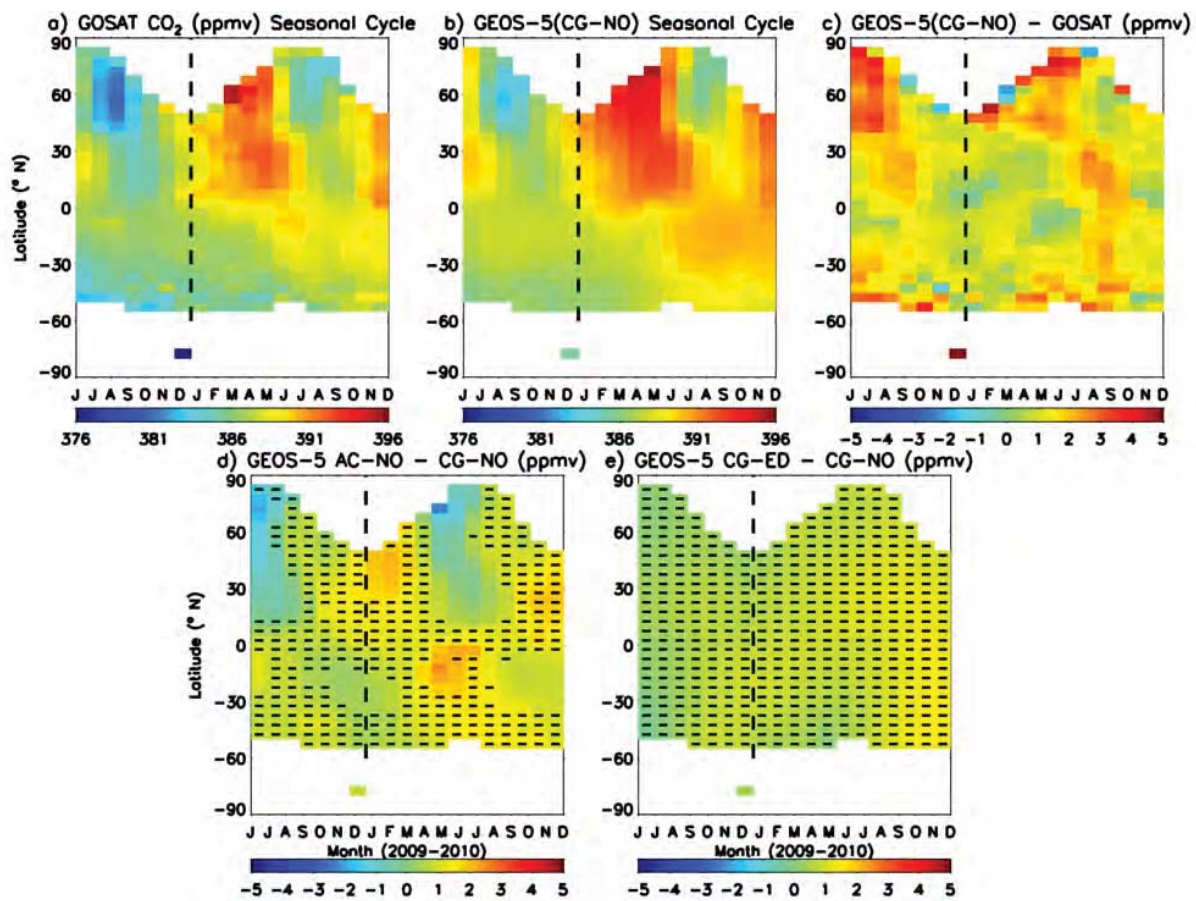


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1539 **Figure 19.** Same as Figure 17, but for July, 2009.

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1543 **Figure 20.** Seasonal cycle of (a) zonal mean ACOS observed mixing ratios, (b) zonal mean GEOS-5
 1544 (CG-NO) mixing ratios calculated with ACOS weighting functions, and (c) the difference between
 1545 zonal mean model results and observations. Bottom plots show the difference in zonal mean caused by
 1546 land (d) and ocean flux differences (e). Dashed lines indicate months where flux differences are not
 1547 statistically significant from natural variability. All units are ppmv.

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