

1 Enhancement and depletion of lower/middle tropospheric ozone in Senegal during pre-  
2 monsoon and monsoon periods of summer 2008: Observations and Model results.

3 Gregory S. Jenkins<sup>1</sup>, Seydi Ndiaye<sup>2</sup>, Moussa Gueye<sup>2</sup>, Rachel Fitzhugh<sup>3\*</sup>, Jonathan W.  
4 Smith<sup>1</sup>, Abou Kebe<sup>4</sup>

5

6

7

8

9 1. Howard University Program in Atmospheric Science, Washington DC 20059

10 2. Lab for Atmospheric-Oceanic Physics-Simeon Fongang, Cheikh Anta Diop  
11 University, Dakar Senegal

12 3. Department of Atmospheric Ocean and Space Science, University of Michigan,  
13 Ann Arbor, Michigan. \*Now at ITT Corporation 1447 St. Paul Rochester, NY  
14 14621

15 4. Senegal Meteorological Agency, Dakar Senegal

16

16

17 **Abstract**

18 During the summer (8 June through 3 September) of 2008, nine (9) ozonesondes are  
19 launched from Dakar, Senegal (14.75°N, 17.49°W) to investigate ozone (O<sub>3</sub>) variability  
20 in the lower/middle troposphere during the pre-monsoon through monsoon period.  
21 Results during June (pre-monsoon period) show a reduction in O<sub>3</sub>, especially in the 850-  
22 700 hPa layer with SAL events. However, O<sub>3</sub> concentrations are increased in the 950-  
23 900 hPa layer where the peak of the inversion is found and presumably the highest dust  
24 concentrations. We use the WRF-CHEM model to explore the observations of  
25 elevated/reduced O<sub>3</sub> concentrations during the pre-monsoon/monsoon periods. In the  
26 transition period between 26 June and 2 July lower tropospheric (925-600 hPa) O<sub>3</sub>  
27 concentrations are likely enhanced by enhanced biogenic NO<sub>x</sub> emissions from the  
28 Saharan desert and Sahelian soils following several rain events on 28 June and 1 July.  
29 During July and August (monsoon period), with the exception of one SAL outbreak,  
30 vertical profiles of O<sub>3</sub> are well mixed with concentrations not exceeding 55 ppb between  
31 the surface and 550 hPa.

32

33

34

34

## 35 **1. Introduction**

36           Understanding tropospheric ozone (O<sub>3</sub>) variability in the tropics remains an active  
37 area of research with biomass burning, biogenic, anthropogenic and lightning being  
38 important sources of ozone production in the tropics and deposition and heterogeneous  
39 chemistry being important sink of O<sub>3</sub> in the tropics. Tropospheric O<sub>3</sub> is also a greenhouse  
40 gas with positive radiative forcing and contributes to global warming. A likely source of  
41 Northern Hemisphere summer season tropospheric O<sub>3</sub> variability is associated with  
42 Saharan dust events. Each year, between May and October, the Saharan Air Layer (SAL)  
43 (Carlson and Prospero, 1972; Dunion and Veldon, 2004) is a dominant feature  
44 influencing continental areas of West Africa and the Tropical Atlantic. The SAL is  
45 characterized by dry, (low relative humidity), stable air (an inversion capped above the  
46 marine boundary layer), a mid-level easterly jet and reduced visibilities from enhanced  
47 dust. Potential sources of dust are found in Mauritania, Mali and Algeria during June-  
48 July and August in association with the SAL (Middleton and Goudie, 2001).

49           A number of observational studies have shown reduced O<sub>3</sub> in the presence of  
50 Saharan dust (De Reus et al. 2000; Bonasoni et al. 2004). The desert aerosols can reduce  
51 tropospheric O<sub>3</sub> concentrations at a single location in multiple ways: (a) dust aerosols  
52 may serve as deposition sites for O<sub>3</sub> while also reducing photolysis rates. (b)  
53 Heterogeneous chemistry on aerosol surfaces can reduce important precursors (OH, HO<sub>2</sub>)  
54 associated with O<sub>3</sub> production. (c) The production of Nitric Acid (HNO<sub>3</sub>), leading to  
55 particulate nitrate can act as a sink for NO<sub>x</sub> and limit O<sub>3</sub> production (Zhang et al. 1994;  
56 Jacob, 2000; Bian and Zender 2003; De Reus et al. 2000; Tang et al. 2003). Hence, the

57 SAL acts as a potential sink for O<sub>3</sub>, reducing its greenhouse forcing while also scattering  
58 solar radiation causing daytime cooling at the surface.

59         Recent observations of the SAL have examined its radiative impact (Myhre et al,  
60 2003), chemical composition (Twohy et al. 2008), aerosol and water vapor structure  
61 (Ismail et al. 2010), and potential impact on African Easterly Waves (AEWs) and  
62 Tropical Cyclogenesis (Jenkins et al. 2008; Zipser et al. 2009). Here we report on the  
63 variability of lower/middle tropospheric O<sub>3</sub> (surface through 550 hPa) during the pre-  
64 monsoon and monsoon periods of 2008 at a location in the semi-arid Sahelian zone.

## 65 **2. Observational data and Model Simulations**

66         In preparation for ozonesonde launches Vaisala ECC6AB ozonesondes were  
67 prepared 3-7 days in advance and launched in concert with RS92 radiosonde at 1200  
68 UTC during June, July and August of 2008. Table 1 shows the 9 ozonesonde launches  
69 from Dakar, Senegal (14.75°N, 17.49°W) during the period. Aerosol observations are  
70 derived from the Aerosol Optical Thickness (AOT) measurements from the Aerosol  
71 Robotic Network (AERONET) at Mbour, Senegal (14.39°N, 16.59°W) and from the  
72 Space-borne Ozone Monitoring Instrument (OMI) Aerosol Index (AI) for the summer of  
73 2008. The 1° x 1° Deep Blue product from the Moderate Resolution Imaging  
74 Spectrometer (MODIS) instrument aboard the AQUA satellite is used for AOT over land  
75 areas. Daily averages of AOT are constructed from the hourly AOT data from Mbour,  
76 Senegal. Tropospheric Column Ozone (TCO) estimates for the summer of 2008 at 1° x  
77 1.25° are also used (Ziemke et al. 2006.)

78         To address daily and seasonal tropospheric O<sub>3</sub> variations for comparison with the

79 measurements the Weather and Research Forecasting with Chemistry (WRF-CHEM)  
80 model is used to simulate O<sub>3</sub> concentrations over West Africa during the summer of 2008  
81 (Grell et al. 2005). The WRF-CHEM computes gas phase chemistry with 55 prognostic  
82 species and 134 reactions (Fast et al. 2005). The WRF-CHEM simulations use 30 km grid  
83 spacing, 27 vertical levels and the top of the model is 50 hPa. The biogenic sources of  
84 volatile organic compounds are computed (Guenther et al. 1995) and the model is  
85 initialized from a uniform state with O<sub>3</sub> concentrations that increases with height to the  
86 stratosphere. NCEP final analyses at 6-hour intervals provide meteorological initial and  
87 boundary conditions to the WRF-CHEM. Here we define the pre-monsoon period as the  
88 month of June and the monsoon period from July through September.

89

### 90 **3. Results**

91        Figures 1a-d show the satellite derived TCO values across West Africa and the  
92 Eastern Atlantic for June through September 2008. A north-south gradient of TCO  
93 values with higher TCO values are generally found over the Sahara relative to Sahelian  
94 and Guinean regions of West Africa. The TCO gradient is weakest and strongest during  
95 June and August respectively. During June, TCO values of 37-40 DU are found over the  
96 desert regions with slightly lower values to the south (Figure 1a). However, by August  
97 TCO values are considerably lower over the Sahelian and Guinea regions of West Africa.  
98 Figures 2a-c show the OMI AI along with the AOT from Mbour, Senegal for the period  
99 of ozonesonde launches during 2008. There are a number of high AI/AOT days  
100 associated with SAL outbreaks during June and July and considerably fewer days in  
101 August. The numbers of days where AI was greater than 2.5 during June, July and  
102 August were five (5), three (3) and one (1) respectively. The numbers of days where

103 AOT values are greater than 0.6 during June, July and August are seven (7), four (4) and  
104 two (2), respectively. The OMI AI is positive correlated to the Mbour AOT values with  
105 correlation values of 0.82, 0.86 and 0.79 during June, July and August respectively.  
106 TRMM daily averaged rain amounts for a  $5^{\circ} \times 5^{\circ}$  (11.5-16.5°N, 12.5-17.5°W) box over  
107 Senegal and Gambia shows very little precipitation until the end of June when wetter  
108 conditions begin and continue through July and August (Figure 2d).

109 Figure 3a shows ozonesondes for the summer of 2008 with a range in  $O_3$   
110 concentrations from approximately 20-80 ppb between the surface and 450 hPa. Table 1  
111 shows that highest column ozone in the 925-550 hPa layer is found on 12 June (20.5 DU)  
112 with 14.2 DU on 2 July. The lowest column ozone in the 925-550 hPa layer is found on  
113 27 September (6.4 DU) followed by 8 June (6.6 DU). Figure 3b shows the two vertical  
114 profiles of  $O_3$  between the end pre-monsoon period and the start of the monsoon season;  
115 a significant enhancement is found between the surface and 600 hPa. We discuss  
116 possible causes for elevated  $O_3$  during the transition period in Section 3.2.

### 117 *3.1 Pre-monsoon $O_3$ measurements*

118 There are three (3) ozonesonde launches, 8, 10, 15 June with OMI AI values  $> 2$   
119 which are associated with SAL events in Figure 2. During the pre-monsoon period  
120 (Figure 4a),  $O_3$  concentrations are reduced on 8 and 10 June relative to the other profiles,  
121 especially in the 850-600 hPa layer. These are also the two days with the lowest 925-550  
122 hPa column ozone during the pre-monsoon period (Table 1). In the 950-900 hPa layer,  
123 just below the depleted layer there is also evidence of enhanced  $O_3$  concentrations. Figure  
124 4b shows low relative humidity ( $< 20\%$ ) at approximately 950 hPa for 8, 10, 12 and 15

125 June, which begins to increase at pressure levels less than 700 hPa, except for 12 June  
126 where RH values rapidly increase above 900 hPa. Figure 4c shows a temperature  
127 inversion is present for all pre-monsoon ozone profiles, but the very strongest  
128 temperature inversions are found for 8, 10 and 15 June. It is at the peak of temperature  
129 inversion that increases in O<sub>3</sub> concentrations are found (Figure 4a). This is also the  
130 altitude where high dust concentrations are found (Ismail et al. 2010).

131 Figures 5a-h show OMI- AI with 925 hPa streamlines along with the Deep Blue AOT  
132 overlain by 700 hPa streamlines for comparison during the pre-monsoon period.  
133 Elevated aerosol loadings on 8, 10 and 15 June are associated with either northerly or  
134 northeasterly winds at 925 hPa (Figures 5 a, b, d). In contrast, winds with a southerly  
135 component are found on 12 June and 26 June (Figures 5c, 5e). Moreover an African  
136 Easterly Wave (AEW) (Burpee et al. 1972) at 700 hPa streamlines, exits the continent on  
137 12 June and is associated elevated RH, cloudiness and precipitation in Dakar. Reduced  
138 O<sub>3</sub> concentrations between 850 and 600 hPa on 8, 10 June in concert with higher AI/AOT  
139 from Saharan dust support depletion through heterogeneous chemistry.

### 140 *3.2 Pre-Monsoon/Monsoon Transition*

141 Satellite and aircraft observations along with modeling studies have identified  
142 NO<sub>x</sub> emissions from soil as an important source of tropospheric NO<sub>x</sub> during June, July  
143 and August in Sahelian and Saharan zones (Jeagle et al. 2004; Van der A et al. 2008;  
144 Williams et al 2009; Delon et al. 2008; Delon et al. 2010). Pulses of biogenic NO<sub>x</sub>  
145 emissions are released into the atmosphere from the soil at the end of the dry season with  
146 the first rains. During the 2006 African Monsoon Multidisciplinary Analyses (AMMA)

147 campaign (Redelsperger et al. 2006), aircraft measurements flown over wet/dry soils  
148 during the AMMA field campaign show higher  $\text{NO}_x$  concentrations in the boundary layer  
149 over the recently wet soils (Stewart et al. 2008).  $\text{O}_3$  concentrations are significantly  
150 higher over wet soils when compared to dry surfaces and low correlations with CO imply  
151 that  $\text{NO}_x$  production is not associated with anthropogenic or biomass burning sources  
152 (Stewart et al. 2008). A sharp gradient in  $\text{O}_3$  concentrations are found implying produced  
153 locally  $\text{O}_3$  estimated at  $1 \text{ ppb h}^{-1}$  from soil  $\text{NO}_x$ .

154 Figure 3b shows that  $\text{O}_3$  concentrations between 950 and 600 hPa are considerably  
155 higher at the start of the monsoon period (2 July) when compared to the end of the pre-  
156 monsoon period (26 June).  $\text{O}_3$  concentrations are more than 10-20 ppb larger on 2 July  
157 when compared to 26 June. The enhancement of  $\text{O}_3$  coincides with several rainfall  
158 events in Senegal that the vicinity of Dakar on 28, 30 June (Figures 6 a,b). TRMM daily  
159 rain estimates were highest on 1 July with daily amount approaching 80 mm in some  
160 areas to the east of Dakar (Figure 6d). The rainfall was associated with the passage of an  
161 AEW on 1 July and a plausible cause for  $\text{O}_3$  enhancement between 26 June and 2 July by  
162 biogenic  $\text{NO}_x$  emissions from wet Sahelian soils.

### 163 *3.3 Monsoon $\text{O}_3$ measurements*

164 During the monsoon period, the vertical profiles of  $\text{O}_3$  are nearly uniform on 2 July,  
165 27 August and 3 September, except for 2 August when there is a SAL event affecting  
166 Senegal (Figure 7a). The 2 August measurement shows the enhance/depletion pattern of  
167  $\text{O}_3$  concentrations in the vertical profile as in earlier SAL events with enhancement near  
168 the peak of the SAL temperature inversion and depletion immediately above this level.

169 Figure 7b shows humid conditions over Senegal during July, August and September



170 except for 2 August when a SAL event occurred over Senegal. The 2 August vertical  
171 profile of relative humidity shows values less than 10% in the 950-900 hPa layer. In  
172 contrast a very moist profile can be found on 27 August with RH values between 90 and  
173 100% from 950 through 750 hPa. A large temperature inversion is found on 2 August in  
174 association with the SAL event with smaller temperature inversions found during the  
175 other three monsoon launches (Figure 7c). Table 1 shows that during the monsoon  
176 period the lowest column O<sub>3</sub> is found on 27 August in association with the highest  
177 relative humidity in the 925-550 hPa layer.

178 Figures 8a-h show OMI- AI with 925 hPa streamlines along with the Deep Blue AOT  
179 overlain by 700 hPa streamlines for comparison during the monsoon period. During this  
180 period, only 2 August shows elevated AI and AOT in association with a thermal low  
181 (heat low) over Southern Mauritania (Figure 8b). During the other three days (2 July, 27  
182 August, 3 September), the highest AI or AOT values are found over higher latitudes (>  
183 20 degrees). Also evident at 700 hPa are strong AEWs, which have closed vortices on 2  
184 July, 27 August and 3 September (Figure 8a, c, d).

185 *3.4 WRF-CHEM simulations of elevated O<sub>3</sub> concentrations on 12 June and reduced O<sub>3</sub>*  
186 *concentrations during the monsoon period.*

187 Table 1 shows that during the period of 10 June through 12 June, the 925-550 hPa  
188 column O<sub>3</sub> is increased by a factor of 2.77; between the SAL air mass and the passage of  
189 the AEW. This is followed by a decreased by nearly a factor of 1.8 between 12 and 15  
190 June. The elevated O<sub>3</sub> concentration on 12 June could be due to: (a) a stratospheric  
191 intrusion; (b) a biogenic pulse of NO<sub>x</sub> from Sahelian soils enhancing O<sub>3</sub> or; (c) lightning-

192 NO<sub>x</sub> which enhances O<sub>3</sub> concentrations that is then transported to the lower/middle  
193 troposphere (Grant et al. 2008). To address a possible stratospheric influence, the WRF-  
194 CHEM is initialized at four different times at 0000 UTC, 1 June; 1200 UTC, 4 June; 1200  
195 UTC 5 June; 1200 UTC 6 June to simulate elevated O<sub>3</sub> concentration for the period 10-12  
196 June.. Figures 9 a-d show for the period of 8 June and 13 June that higher concentrations  
197 of O<sub>3</sub> are found in the middle and upper troposphere based on vertical profiles of O<sub>3</sub> at  
198 14.5 °N, 17.5° W near Dakar, Senegal, in a region of weak sinking motion. WRF-  
199 CHEM forecasts that are initialized on 5 and 6 June show O<sub>3</sub> concentrations in the 90 to  
200 120 ppb range at the 200-300 hPa levels with values of 35-50 ppb extending downward  
201 into middle to lower troposphere (Figures 9c, d).

202 The elevated levels are consistent with the observations between 10 and 12 June  
203 where a large increase is found in the lower/middle troposphere (Figures 4a). For  
204 example, O<sub>3</sub> values of 20 ppb are found near 600 hPa on 10 June but increase to nearly  
205 65 ppb on 12 June. Figure 10 provides further support for a stratospheric intrusion when  
206 comparing O<sub>3</sub> concentrations on 10, 12 June for 1000-150 hPa levels. On 12 June O<sub>3</sub>  
207 values greater than 400 ppb are found just below 200 hPa. The enhancement of O<sub>3</sub>  
208 relative to 10 June is found throughout the lower, middle and upper troposphere. Hence,  
209 O<sub>3</sub> elevated concentrations from the lower stratospheric is supported by observations and  
210 WRF-CHEM results during the period of 10-12 June.

211 Next we use WRF-CHEM to explore the lower O<sub>3</sub> concentrations that are  
212 relatively well mixed during August and September. Table 1 shows low 925-550 hPa  
213 column O<sub>3</sub> under moist and low aerosol loading conditions. In particular, 27 August and 3  
214 September measurements show low concentrations of O<sub>3</sub> that are well mixed relative to

215 the O<sub>3</sub> measurements under humid conditions (Figure 7a). There are several factors that  
216 could lead to well mixed, reduced O<sub>3</sub> concentrations in the monsoon season: (a)  
217 increased O<sub>3</sub> and NO<sub>x</sub> deposition; and (b) the northward transport of O<sub>3</sub> poor air from  
218 areas to the south of the Sahel.

219 First, a reduction in O<sub>3</sub> concentrations during the monsoon period is expected  
220 because of increased deposition with the rapid growth of vegetation. After June, the  
221 northward movement of the monsoon and the accompanied southwest moist flow leads to  
222 increased instability and precipitation. The precipitation supplies an important source of  
223 moisture for vegetation growth of West Africa (Guinea region northward to the Sahel).  
224 The expansion of vegetation leads to increased deposition and a north-south gradient in  
225 deposition rates and hence ozone in the lower troposphere. Measurements in Senegal  
226 during 2006 estimate a maximum deposition velocity of 1.5 cm-s<sup>-1</sup> with higher values  
227 noted during precipitation events (Grant et al. 2008). The removal of NO<sub>x</sub> through  
228 dry/wet deposition would lead to reduced O<sub>3</sub> concentrations in lower latitudes relative to  
229 the drier semi-arid and arid regions in the Sahelian zone and Sahara desert (Delon et al.  
230 2010).

231 Second, northward surges in the monsoon flow or the passage of AEWs should lead  
232 to an overall poleward transport of O<sub>3</sub> poor air in the lower troposphere, because of more  
233 vegetation and higher deposition rates in lower latitudes; onshore flow from the Atlantic  
234 ocean would also transport in O<sub>3</sub> poor air. Conversely surges of dry air from desert  
235 regions should lead an equatorward transport of enriched O<sub>3</sub> concentrations to lower  
236 latitudes. An example of this process is the low O<sub>3</sub> concentrations in the lower  
237 troposphere measured at Dakar, Senegal for 27 August, and 3 September. In each case,

238 very moist conditions with 925 hPa flow coming from Atlantic Ocean (Figure 8c) and  
239 lower latitudes (Figure 8d) are found with the approach or passage of an AEW (Figures  
240 8g, h) at 700 hPa. AEWs on 2 July and 27 August are associated with Tropical Cyclones  
241 Bertha and Ike, respectively (Rhome 2008; Berg 2008). The monthly averaged TCO  
242 values (Figure 1) along with recent measurements during the AMMA field campaign  
243 support a north-south gradient in the lower troposphere with lower O<sub>3</sub> concentrations  
244 found in equatorward latitudes of West Africa (Saunois et al. 2008).

245 To examine simulated seasonal O<sub>3</sub> concentrations and its relationship to relative  
246 humidity we show WRF-CHEM forecasted 925 hPa 12-hour instantaneous O<sub>3</sub>  
247 concentrations and relative humidity averaged over the area of 14-16° and 18-16°W for  
248 June, July and August and early September (Figure 11). The highest O<sub>3</sub> concentrations  
249 are found during the month of June with significant reductions simulated during the  
250 months of July and August. Simulated O<sub>3</sub> concentrations of 10-20 ppb are common at  
251 925 hPa during the monsoon period (Figure 11a) relative to June. The lower simulated  
252 O<sub>3</sub> concentrations are found when the simulated relative humidity (RH) at 850 hPa values  
253 begins to increase (Figure 11b). Elevated 850 hPa relative humidity can be used as a  
254 proxy of the monsoon flow during July, August and September. The simulated increases  
255 in relative humidity are in agreement with satellite observed wetter conditions during July  
256 and August (Figure 2d)

257

258

259 **4. Discussion and Summary**

260           During the pre-monsoon period reductions of O<sub>3</sub> concentrations in the lower  
261 troposphere associated with SAL outbreaks. Figures 12 a-d show the vertical profiles of  
262 O<sub>3</sub> concentration, relative humidity and temperature in the 1000-750 hPa layer for the  
263 four SAL events that occurred during the summer of 2008. Similar vertical profiles of  
264 O<sub>3</sub> concentrations (> 50 ppb for 2 August) are found in three of the four ozonesondes:  
265 An increase in O<sub>3</sub> concentrations is found to overlap with a temperature inversion and  
266 very low relative humidity (Figures 12a, b, d). Above the layer where O<sub>3</sub> concentrations  
267 are enhanced, there is a depleted layer of O<sub>3</sub> in the 900-750 hPa layer. The depleted O<sub>3</sub>  
268 layer in Figure 12 is consistent with the earlier results of De Reus et al. (2000) and  
269 Bonasoni et al. (2004) with heterogeneous chemistry being a primary source of O<sub>3</sub>  
270 depletion (Zhang et al. 1994; Tang et al. 2004). Tang et al. (2004) show through  
271 modeling studies and aircraft observations that losses of O<sub>3</sub> and HNO<sub>3</sub> through direct  
272 interaction with dust were large during observed dust storms in the ACE-Asia field  
273 campaign.

274

275           Figure 13a depicts the changes in aerosol concentrations, relative humidity and O<sub>3</sub>  
276 concentrations, during a ideal SAL events with significant dust loading, based on the  
277 findings from this study. During this SAL event, a vertical profile of O<sub>3</sub> would show:  
278 Layer A – from the surface to 950 hPa with slowly increasing O<sub>3</sub> concentrations and low  
279 aerosol concentrations; Layer B- a 50-100 hPa layer with elevated O<sub>3</sub>, high aerosols  
280 concentrations and a strong temperature inversion; Layer C - a 100-200 hPa layer with  
281 reduced O<sub>3</sub> concentrations and smaller sized aerosols. One might expect the largest losses  
282 of O<sub>3</sub> in layer B in association with heterogeneous reactions but the observations show

283 that this is not observed and hence there must be a source of O<sub>3</sub>. The most likely source  
284 would be biogenic sources of NO<sub>x</sub> from Saharan soils that have been lifted that act as a  
285 source of atmospheric O<sub>3</sub> in downstream regions. The exact mechanism for O<sub>3</sub>  
286 enrichment by Saharan dust remains uncertain.

287 A suggested mechanism for enhancing O<sub>3</sub> concentrations in this shallow layer  
288 may come from the activation of a biogenic NO<sub>x</sub> pulse when encountering moist  
289 conditions (Figure 13b). Nitrate formation on dust aerosols is an end-product after  
290 heterogeneous reactions between HNO<sub>3</sub> and dust aerosols. Gravitational settling of  
291 aerosols between 900-600 hPa that may lead to significant amount aerosols with nitrate  
292 on its surfaces through heterogeneous processes which accumulate just above the  
293 inversion. At Dakar, very moist conditions (RH > 80%) are found from the surface to the  
294 950 hPa just below the inversion associated with the SAL. Even though the inversion  
295 inhibits vertical motions, some mixing between the SAL layer and shallow moist layer  
296 below is possible.

297 Vlasenko et al. (2006) show in laboratory studies that nitrate-coated dust aerosols  
298 will increase their hygroscopicity in the presence of high relative humidity. Twohy et al.  
299 (2009) and Ismail et al. (2010) show that some SAL events have areas of enhanced  
300 moisture embedded within the SAL and that dust particles from the desert have  
301 hygroscopic properties and can serve as cloud condensation nuclei (CCN). A critical  
302 threshold of moisture on the surface of the aerosols could lead water stressed microbes to  
303 denitrify and release airborne biogenic NO<sub>x</sub> leading to higher O<sub>3</sub> concentrations.

304 Other findings reported also include:

- 305 • Elevated O<sub>3</sub> concentrations on 12 June are likely caused by a stratospheric  
306 intrusion based on WRF-CHEM results and potentially enhanced NO<sub>x</sub>  
307 emissions from dry Sahelian soils with the passage of an AEW. However,  
308 lightning-NO<sub>x</sub> leading to O<sub>3</sub> enhancement followed by downward vertical  
309 transport by convective downdrafts cannot be ruled out as an additional  
310 source for 12 June (Jenkins et al. 2008b; Grant et al. 2008).
- 311 • During the transition between the pre-monsoon and monsoon periods (26  
312 June and 2 July) a significant enhancement of O<sub>3</sub> concentrations in the  
313 lower/middle troposphere are found after precipitation events in area  
314 surrounding Dakar on 28 June and 1 July 2008. We suggest that pulses of  
315 biogenic NO<sub>x</sub> emissions from dry Sahelian soils are the primary cause of  
316 enhanced O<sub>3</sub> concentrations consistent with early studies (Stewart et al.  
317 2008; Delon et al. 2008; Delon et al. 2010).
- 318 • Low O<sub>3</sub> concentrations are found in the 925-550 hPa layers during the  
319 monsoon period (27 August and 3 September) and likely linked to dry  
320 deposition of O<sub>3</sub> and dry/wet deposition NO<sub>x</sub> and HNO<sub>3</sub> during the  
321 monsoon period (Grant et al. 2008; Delon et al. 2010).

322 Additional chemical, radiative and aerosol measurements along with chemical  
323 modeling on a regional basis in West Africa will provide additional insights into the  
324 processes that control O<sub>3</sub> concentrations in the lower troposphere.

325 **Acknowledgements**

326 This work was supported by ATM-0621529. We thank all of the Met operators for their  
327 help in this project. We thank Didier Tanre for their efforts in establishing and  
328 maintaining the Mbour, Senegal site.

329



329 **References**

330 Berg, R.: Tropical Cyclone Report Hurricane Ike (AL092008), National Hurricane Center  
331 <http://www.nhc.noaa.gov/2008atlan.shtml>, 2008.

332

333 Bian, H. and Zender C. S.: Mineral dust and global tropospheric chemistry: Relative roles  
334 of photolysis and heterogeneous uptake, *J. Geophys. Res.*, 108, 4672,  
335 doi:10.1029/2002JD003143, 2003.

336 Bonasoni, P. et al.: Aerosol-ozone correlations during dust transport episodes, *Atmos.*  
337 *Chem. Phys.* 4, 1201-1215, 2004.

338 Burpee, R. W.: The Origin and Structure of Easterly Waves in the Lower Troposphere of  
339 North Africa, *J. Atmos. Sci.*, 29, 77-90, 1972.

340

341 Camara, M., Jenkins, G. S., Konare, A.: Impact of dust on West African Climate during  
342 2005 and 2006, *Atmos. Chem. Phys.*, 10, 3053-3086, 2010.

343 Carlson T. and Prospero, J. M.: The Large-Scale Movement of Saharan Air Outbreaks  
344 over the Northern Equatorial Atlantic, *J. Appl. Meteorol.*, 11, 283-297, 1972.

345 De Reus, M. et al.: Airborne observations of dust aerosols over the North Atlantic during  
346 ACE2: Indications for heterogeneous ozone destruction, *J. Geophys. Res.*, 105,  
347 15-263-15,275, 2000.

348 Delon, C., Reeves, C. E., Stewart, D. J., Serça, D., Dupont, R., Mari, C., Chaboureau, J.-  
349 P., and Tulet, P.: Biogenic nitrogen oxide emissions from soils – impact on NO<sub>x</sub>  
350 and ozone over West Africa during AMMA (African Monsoon Multidisciplinary  
351 Experiment): modelling study, *Atmos. Chem. Phys.*, 8, 2351-2363, 2008.

352 Delon, C., et al.: Atmospheric nitrogen budget in Sahelian dry savannas, *Atmos. Chem.*  
353 *Phys.*, 10, 2691-2708, 2010.

354 Dunion, J. P., and Velden C. S.: The impact of the Saharan air layer on Atlantic tropical  
355 cyclone activity, *B. Am. Meteorol. Soc.*, 85, 353–365, doi:10.1175/BAMS-85-3-  
356 353, 2004.

357 Fast, J., et al.: Evolution of ozone, particulates, and aerosol direct radiative forcing in the  
358 vicinity of Houston using a fully coupled meteorology-chemistry-aerosol model,  
359 *J. Geophys. Res.*, 111, D21305, doi:10.1029/2005JD006721, 2006.

360 Grant et al.: Ozone Transport by Mesoscale Convective Systems in Western Senegal,  
361 *Atmos. Environ.*, 42, 7104–7114, 2008.

362 Grell, G. A., et al.: Fully coupled “online” chemistry within the WRF model, *Atmos.*  
363 *Environ.*, 39, 6957–6975, 2005.

364 Guenther, A. et al: A global model of natural volatile organic compound emissions, *J. J.*  
365 *Geophys. Res.*, 100, 8873-8892, 1995.

366 Ismail, S. et al.: LASE Measurements of Water Vapor, Aerosols and Cloud Distributions  
367 in Saharan Air Layers and Tropical Disturbances, *J. Atmos. Sci.*, 67, 1026-1047,  
368 2010.

369 Jacob, D. J.: Heterogeneous chemistry and tropospheric ozone, *Atmos. Environ.*, 24,  
370 2131-2159, 2000.

371 Jaeglé, L et al.: Satellite mapping of rain-induced nitric oxide emissions from soils, *J.*  
372 *Geophys. Res.*, 109, D21310, doi:10.1029/2004JD004787, 2004.

373 Jenkins, G.S., Pratt, A. Heymsfield, A. Possible linkages between Saharan dust and  
374 Tropical Cyclone Rain Band Invigoration in Eastern Atlantic during NAMMA-  
375 06, Geophys. Res. Lett., doi:10.1029/2008GL034072, 2008.

376 Jenkins, G. S., Camara, M., Ndiaye, S.: Observational Evidence of enhanced  
377 middle/upper tropospheric ozone via convective processes over the Equatorial  
378 Tropical Atlantic during the summer of 2006, Geophys. Res. Lett., 35,  
379 doi:10.1029/2008GL033954, 2008b.

380 Middleton, N. J. and A. S. Goudie, A. S.: Saharan dust: Sources and trajectories, T. I.,  
381 Brit. Geogr., 26, 165-181, 2001.

382 Myhre, G. A. et al.: Modeling the radiative impact of mineral dust during the Saharan  
383 Dust Experiment (SHADE) campaign, J. Geophys. Res., 108, 8579,  
384 doi:10.1029/2002JD002566, 2003.

385 Redelsperger, J. L., Thorncroft, C. D., Diedhiou, A. Lebel, T., Parker, D. J., Polcher, J.:  
386 African Monsoon Multidisciplinary Analysis: An International Research Project  
387 and Field Campaign, B. Am. Meteorol. Soc., 87, 1739–1746, 2006.

388 Rhome, J. R.: Tropical Cyclone Report Hurricane Bertha (AL022008), National  
389 Hurricane Center <http://www.nhc.noaa.gov/2008atlan.shtml>, 2008.  
390

391 Saunio, M. et al.: Factors controlling the distribution of ozone in the West African lower  
392 troposphere during AMMA (African Monsoon Multidisciplinary Analysis) wet  
393 season campaign, Atmos. Chem. Phys., 9, 6135–6155, 2009.

394 Stewart, D. J., Taylor, C. M., Reeves, C. E., and McQuaid, J. B: Biogenic nitrogen oxide  
395 emissions from soils: impact on NO<sub>x</sub> and ozone over west Africa during AMMA  
396 (African Monsoon Multidisciplinary Analysis): observational study, Atmos.  
397 Chem. Phys., 8, 2285-2297, 2008.

398 Tang et al.: Impacts of dust on regional tropospheric chemistry during the ACE-Asia  
399 experiment: A model study with observations, J. Geophys. Res., 109,  
400 doi:10.1029/2003JD003806, 2004.

401  
402 Twohy, C. H., et al.: Saharan dust particles nucleate droplets in eastern Atlantic clouds,  
403 Geophys. Res. Lett., 36, L01807, doi:10.1029/2008GL035846, 2009.

404 van der A. et al.: Trends, seasonal variability and dominant NO<sub>x</sub> source derived from a  
405 ten year record of NO<sub>2</sub> measured from space, J. Geophys. Res., 113, ,  
406 doi:10.1029/2007JD009021, 2008.

407 Vlasenko, A. et al.: Effect of humidity on nitric acid uptake to mineral dust aerosol  
408 particles, Atmos. Chem. Phys., 6, 2147-2160, 2006.

409 Williams, J. E et al.: The influence of biogenic emissions from Africa on tropical  
410 tropospheric ozone during 2006: a global modeling study, Atmos. Chem. Phys., 9,  
411 5729-5749, 2009.

412 Zhang, Y., Sunwoo, Y., Kotamarthi, V., Carmichael, G. R.: Photochemical Oxidant  
413 Processes in the Presence of Dust: An Evaluation of the Impact of Dust on  
414 Particulate Nitrate and Ozone Formation, J. Appl. Meteorol., 33,813-824, 1994.

415 Ziemke, J. R. et al.: Tropospheric ozone determined from Aura OMI and MLS:  
416 Evaluation of measurements and comparison with the Global Modeling  
417 Initiative's Chemical Transport Model", J. Geophys. Res., 111,  
418 doi:10.1029/2006JD007089, 2006.

419  
420

421 Zipser, E.J. et al.: The Saharan Air Layer and the Fate of African Easterly Waves—  
422 NASA's AMMA Field Study of Tropical Cyclogenesis, B. Am. Meteorol. Soc., 90,  
423 1137–1156, 2009.

424  
425  
426

427 **Figure Captions**

428 **Figure 1.** OMI/MLS Total Column Ozone for (a) June 2008; (b) July 2008; (c) August  
429 2008; (d) September 2008. Units are in DU.

430 **Figure 2.** Daily area averaged (13-16°N, 16-19°W) OMI derived Aerosol Index (AI) and  
431 Mbour, Senegal AOT for:(a) June; (b) July; (c) 1 August-5 September; (d) Area averaged  
432 (11.5-16.5°N, 12.5-17.5°W) TRMM daily averaged precipitation for 1 June-5 September  
433 2006.

434 **Figure 3.** 1000-450 hPa lower/middle tropospheric vertical profiles of O<sub>3</sub> concentrations  
435 for: (a) all launches; (b) 26 June and 2 July. Units are ppb.

436 **Figure 4.** 1000-450 hPa lower/middle tropospheric vertical profiles of: (a) O<sub>3</sub>  
437 concentrations; (b) relative humidity and (c) temperature for the pre-monsoon period.

438 **Figure 5.** Pre-Monsoon OMI AI/925 hPa streamlines and Deep Blue AOT/700 hPa  
439 streamlines: (a, f) 8 June; (b, g); 10 June; (c, h) 12 June; (d, i) 15 June; (e, j) June 26.

440 **Figure 6.** TRMM daily Precipitation amounts (a) 28 June; (b) 29 June; (c) 30 June; (d) 1  
441 July. Units are mm.

442

443 **Figure 7.** 1000-450 hPa lower/middle tropospheric vertical profiles of: (a) O<sub>3</sub>  
444 concentrations; (b) relative humidity and (c) temperature for the monsoon period.

445 **Figure 8.** Monsoon OMI AI/925 hPa streamlines and Deep Blue AOT/700 hPa  
446 streamlines: (a, e) 2 July; (b, f); 2 August; (c, g) 27 August; (d, h) 3 September.

447 **Figure 9.** Time-height profiles of O<sub>3</sub> concentrations and vertical velocity at 14.5° N,  
448 17.5°W for initial conditions beginning at: (a) 1 June 0000 UTC; (b) 4 June 1200 UTC;  
449 (c) 5 June 1200 UTC; (d) 6 June 1200. Units in ppb and cm-s<sup>-1</sup>.

450 **Figure 10.** 1000-150 hPa vertical profile of O<sub>3</sub> at Dakar for 10,12 June, 1200 UTC.

451 Units in ppb.

452 **Figure 11.** WRF-Chem simulation of: (a) 925 hPa O<sub>3</sub> concentrations for June, July and  
453 August; (b) 850 hPa relative humidity for June, July and August.

454 **Figure 12.** 1000-750 hPa vertical profiles of O<sub>3</sub> (blue), RH (green) and Temperature  
455 (red) for identified SAL events. (a) 8 June; (b) 10 June; (c) 15 June; (d) 2 August. Units  
456 are ppb for O<sub>3</sub>, % for RH and Degree C for temperature.

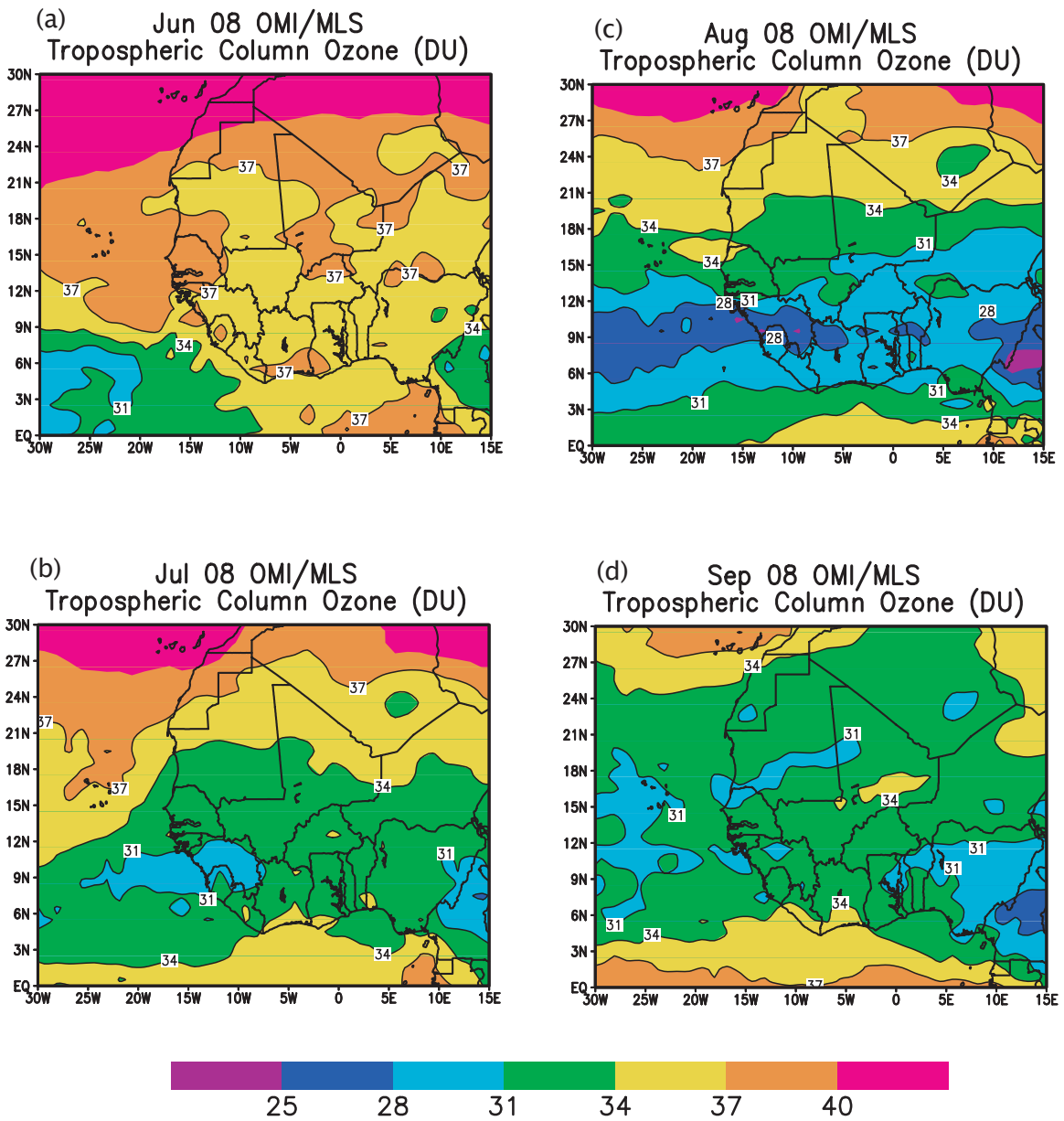
457 **Figure 13.** (a) A depiction of relative humidity, aerosol and O<sub>3</sub> concentrations in various  
 458 layer of the lower/middle troposphere associated with a SAL intrusion. The dash line  
 459 represents the vertical profile of temperature. (b) Proposed mechanism for increasing O<sub>3</sub>  
 460 in the 950-900 hPa layer.

461  
 462  
 463  
 464  
 465  
 466  
 467  
 468  
 469

Date (2008)	Time (UTC)	O <sub>3</sub> 925-550 hPa (DU)	RH 925-550 hPa (%)	OMI AI (13-16°N, 16-19°W)
8 June	1200	6.6	32.2	2.11
10 June	1200	7.4	37.3	1.29
12 June	1200	20.5	65.1	
15 June	1200	11.3	28.7	2.02
26 June	1200	10.5	37.2	1.98
2 July	1200	14.2	67.3	
2 August	1200	10.3	31.3	3.80
27 August	1200	6.4	88.5	0.58
3 Sept	1200	11.0	68.8	0.61

470 **Table 1:** Surface -550 hPa Column O<sub>3</sub>, Relative humidity and OMI AI index.

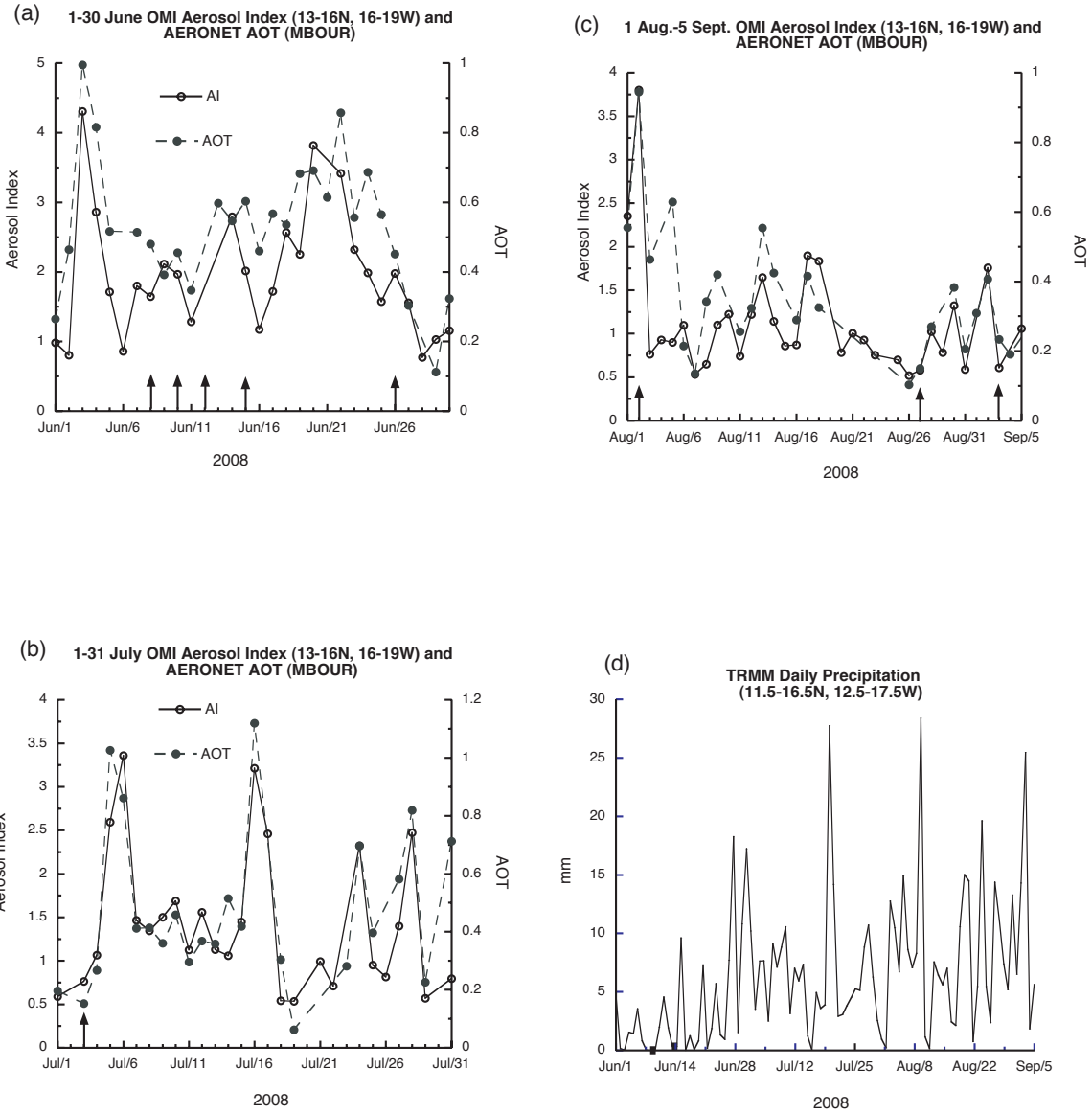
471  
 472  
 473  
 474  
 475



477  
478  
479  
480

**Figure 1.** OMI/MLS Total Column Ozone for (a) June 2008; (b) July 2008; (c) August 2008; (d) September 2008. Units are in DU.

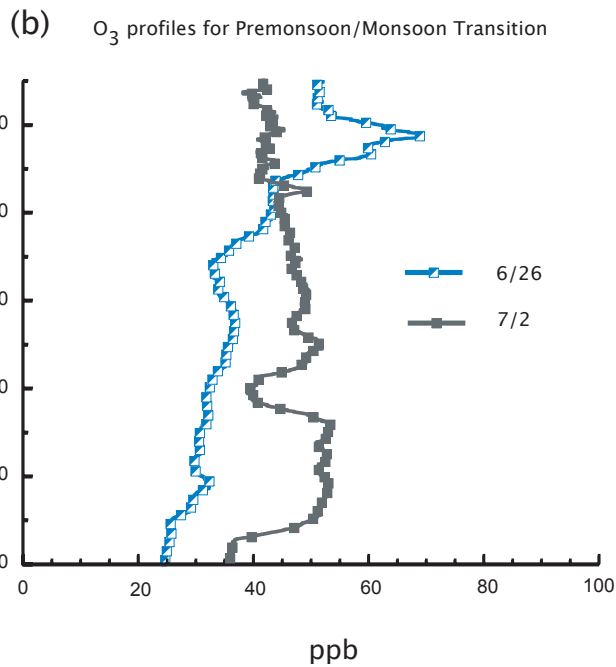
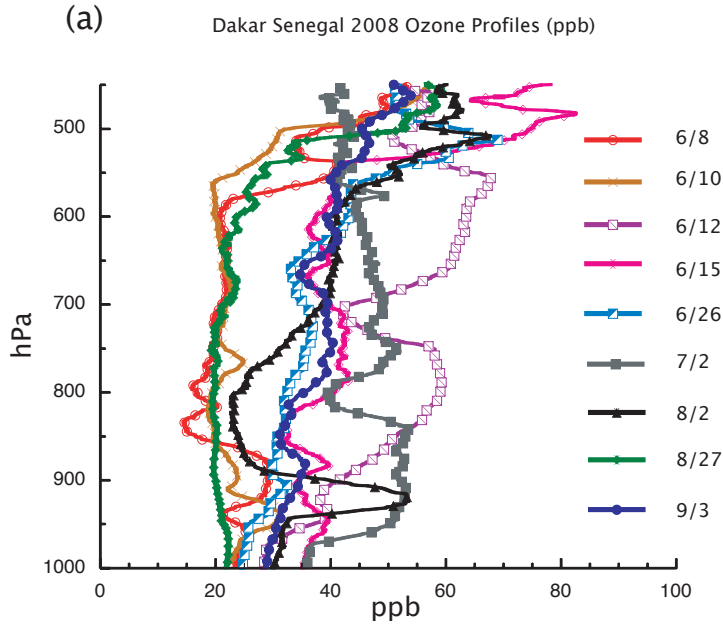




481  
 482 **Figure 2.** Daily area averaged (13-16°N, 16-19°W) OMI derived Aerosol Index (AI) and  
 483 Mbour, Senegal AOT for:(a) June; (b) July; (c) 1 August-5 September; (d) Area averaged  
 484 (11.5-16.5°N, 12.5-17.5°W) TRMM daily averaged precipitation for 1 June-5 September  
 485 2006.

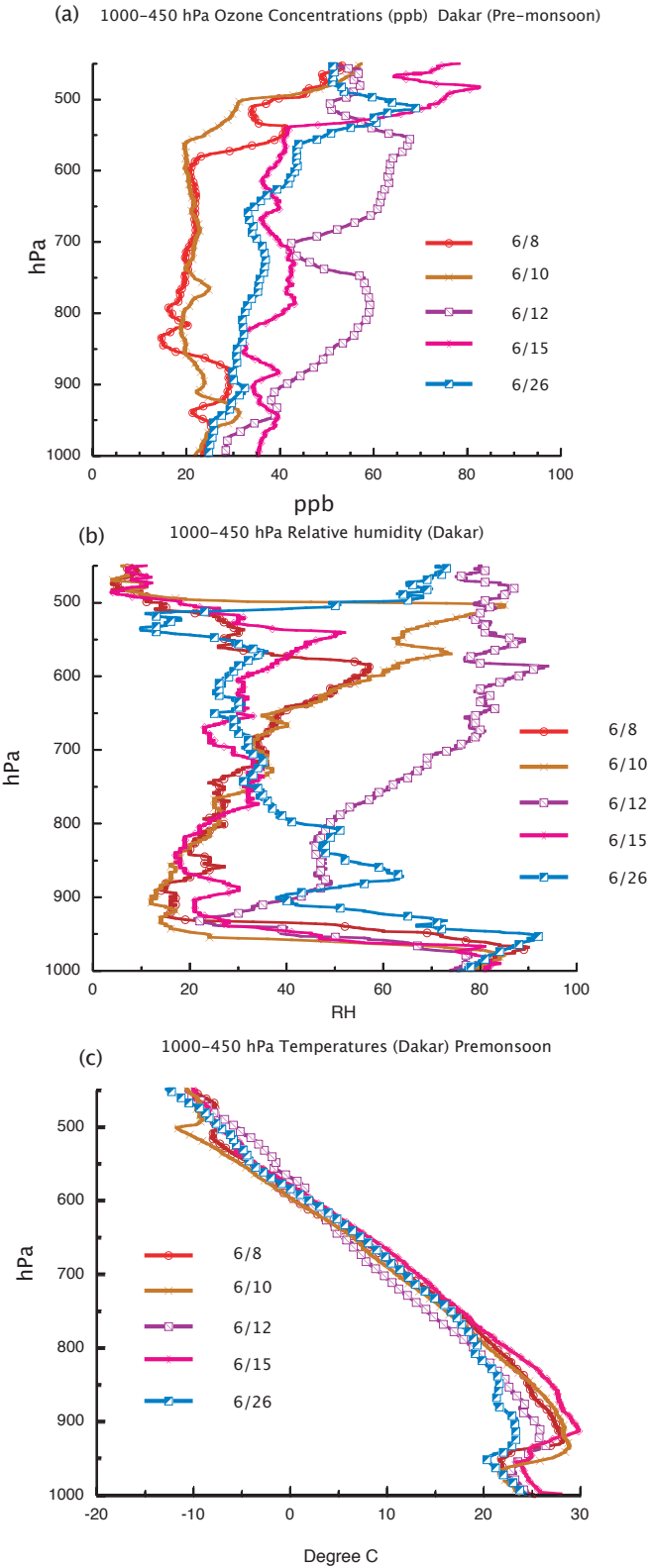
486

487



488

489 **Figure 3.** 1000-450 hPa lower/middle tropospheric vertical profiles of O<sub>3</sub> for: (a) all  
 490 launches; (b) pre-monsoon period and; (c) monsoon periods; (d) 26 June and 2 July.  
 491 Units are ppb.

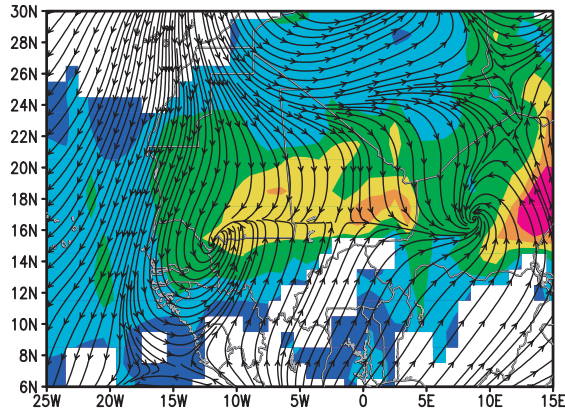


492

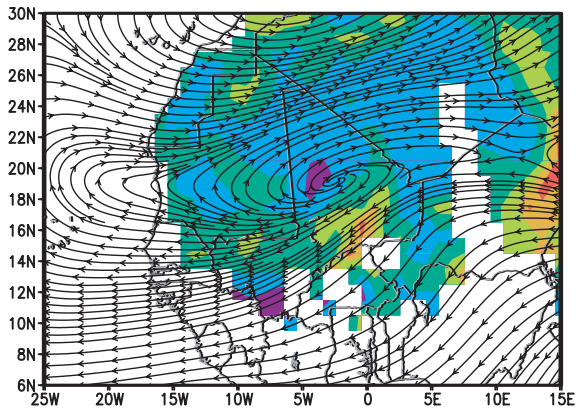
493 **Figure 4.** 1000–450 hPa lower/middle tropospheric vertical profiles of: (a) O<sub>3</sub>

494 concentrations; (b) relative humidity and (c) temperature for the pre-monsoon period.

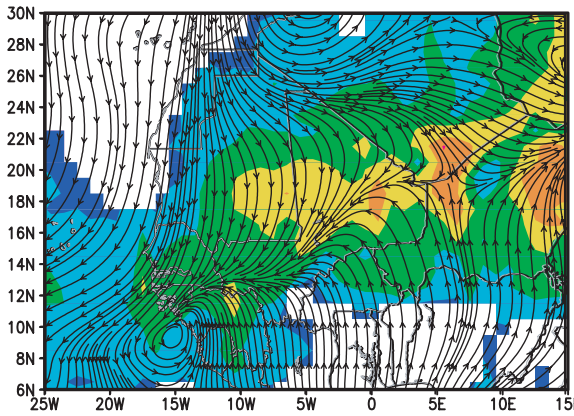
(a) June 8 OMI AI /925 hPa Streamlines



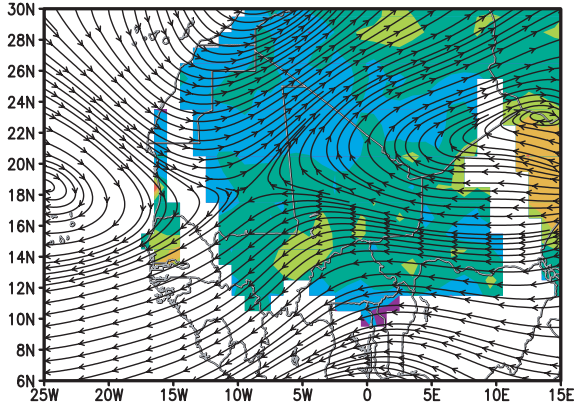
(f) June 8 Deep Blue (AOD)/700 hPa Streamlines



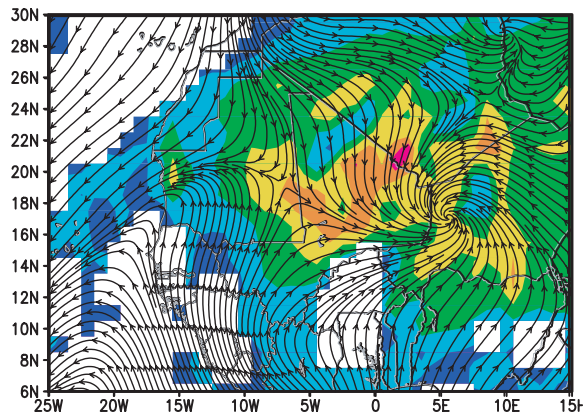
(b) June 10 OMI AI /925 hPa Streamlines



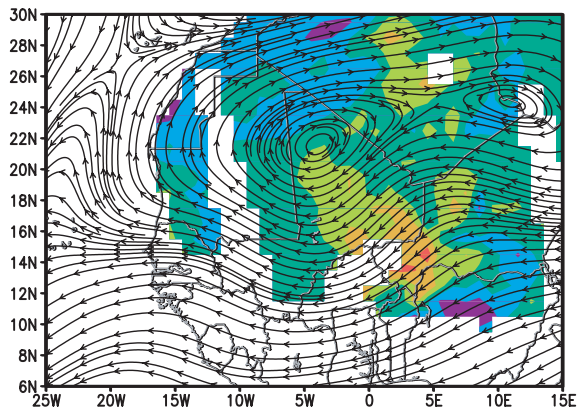
(g) June 10 Deep Blue (AOD)/700 hPa Streamlines



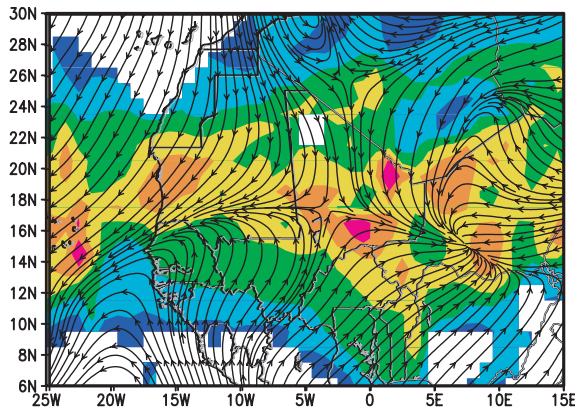
(c) June 12 OMI AI /925 hPa Streamlines



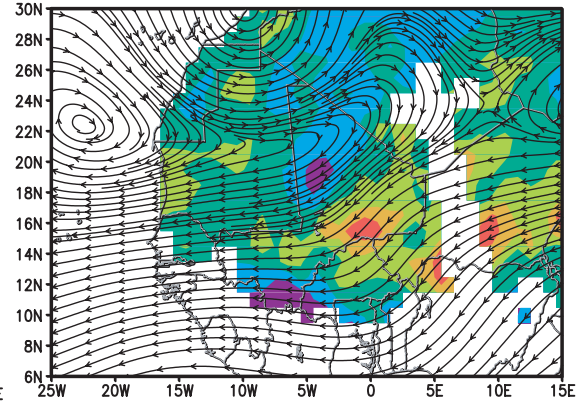
(h) June 12 Deep Blue (AOD)/700 hPa Streamlines



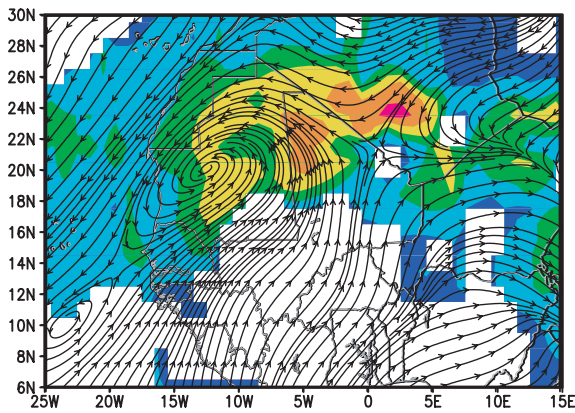
(d) June 15 OMI AI /925 hPa Streamlines



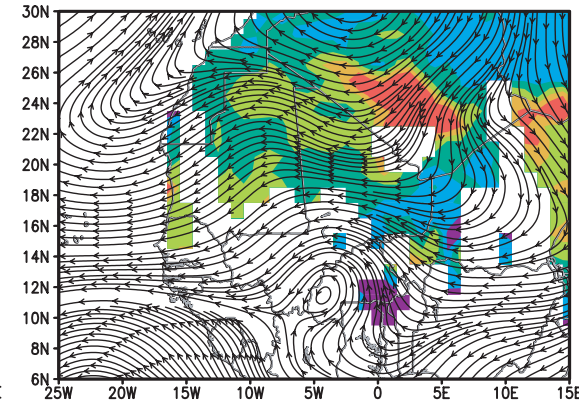
(i) June 15 Deep Blue (AOD)/700 hPa Streamlines



(e) June 26 OMI AI /925 hPa Streamlines



(j) June 26 Deep Blue (AOD)/700 hPa Streamlines



496

497

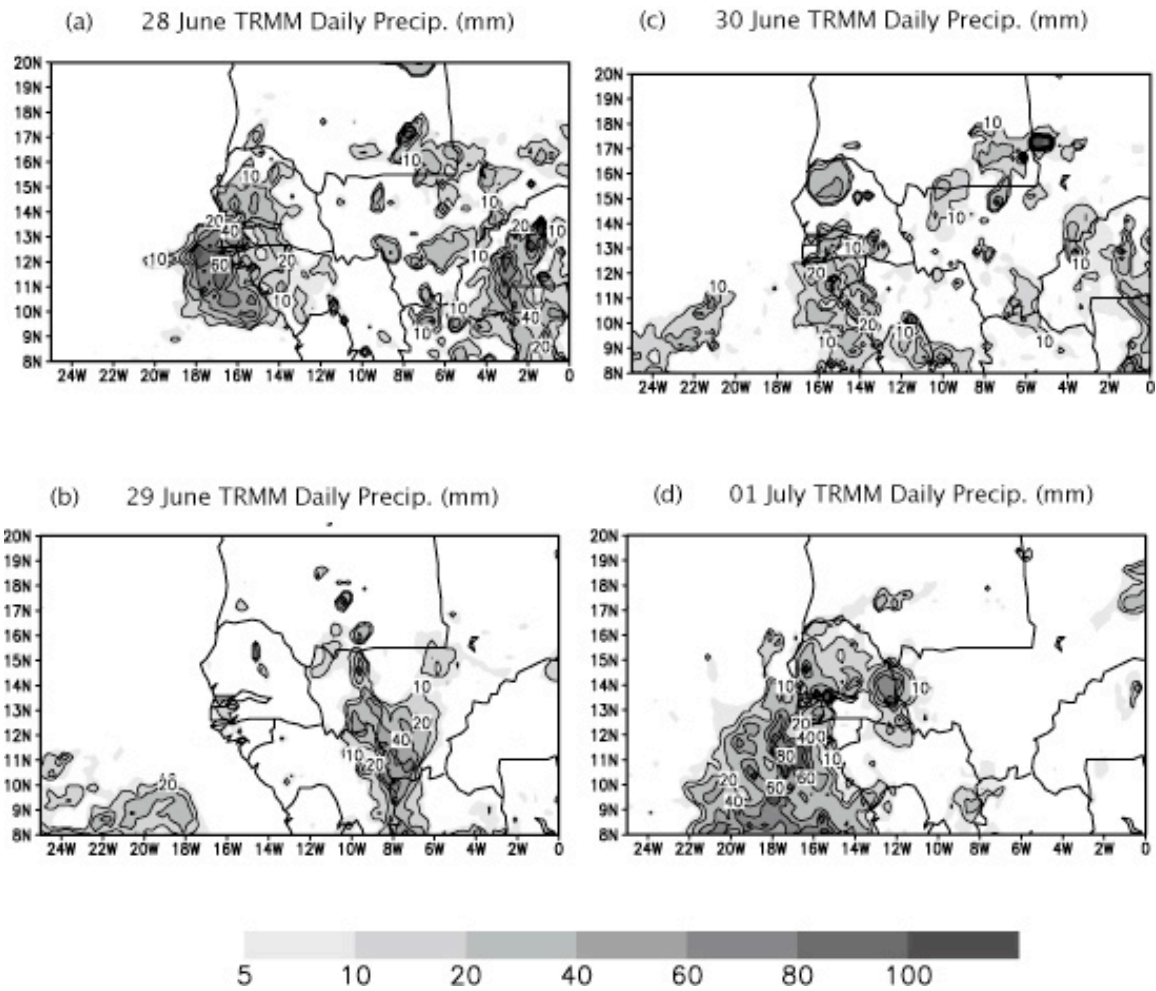
**Figure 5.** Pre-Monsoon OMI AI/925 hPa streamlines and Deep Blue AOT/700 hPa

498

streamlines: (a, f) 8 June; (b, g); 10 June; (c, h) 12 June; (d, i) 15 June; (e, j) June 26.

499



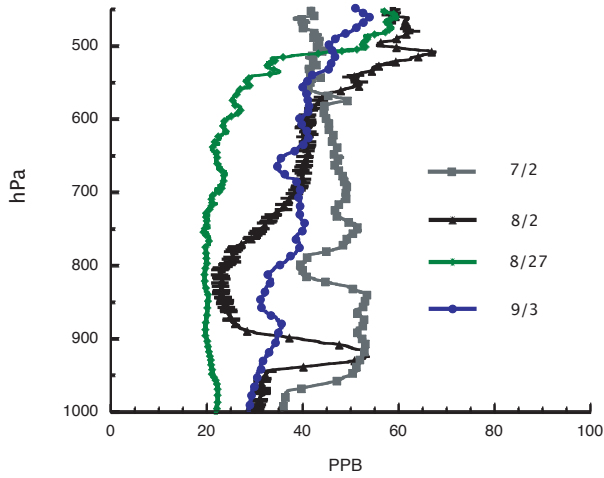


500

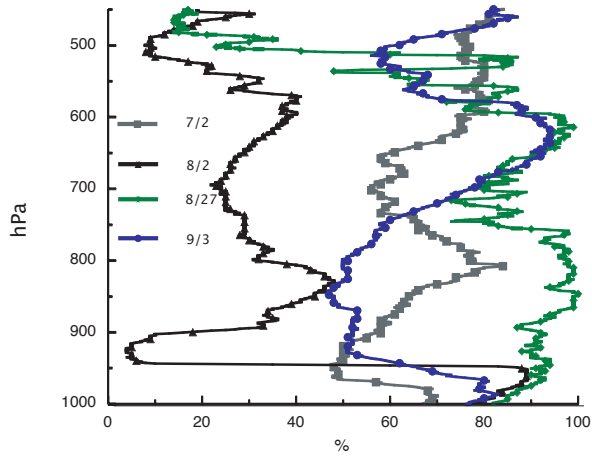
501

502 **Figure 6.** TRMM daily Precipitation amounts (a) 28 June; (b) 29 June; (c) 30 June; (d) 1  
 503 July. Units are mm.

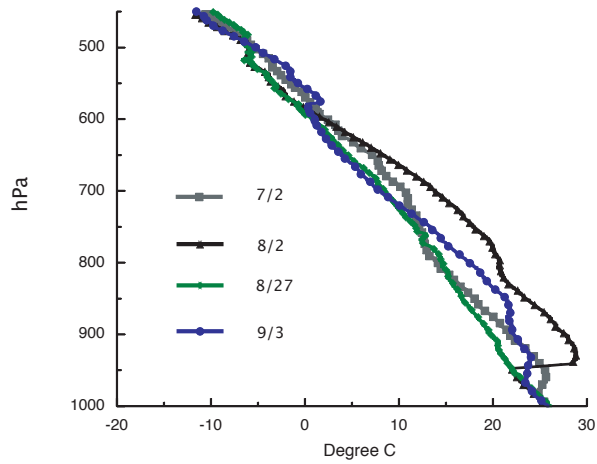
(a) 1000–450 hPa O<sub>3</sub> concentrations (ppb) Dakar (Monsoon)



(b) 1000–450 hPa Relative Humidity (Dakar) – Monsoon



(c) 1000–450 hPa Temperatures (Dakar) Monsoon

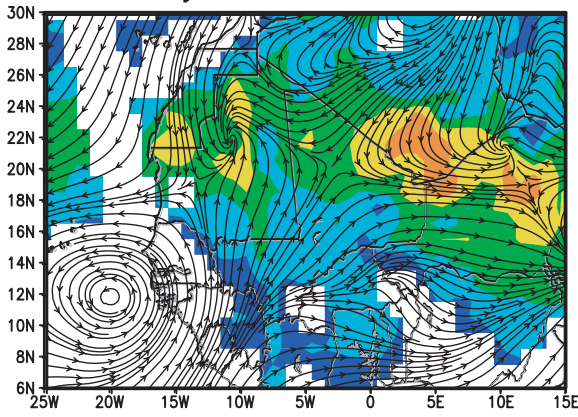


504

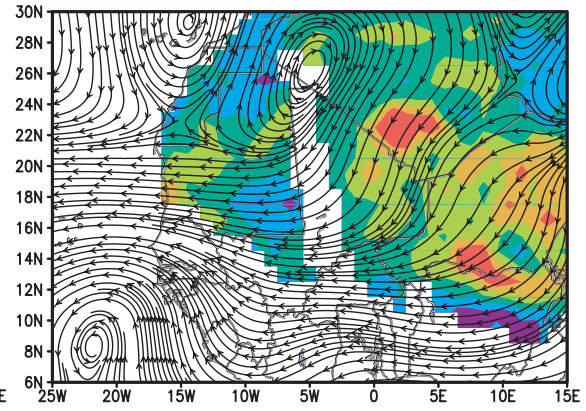
505 **Figure 7.** 1000-450 hPa lower/middle tropospheric vertical profiles of: (a) O<sub>3</sub>

506 concentrations; (b) relative humidity and (c) temperature for the monsoon period

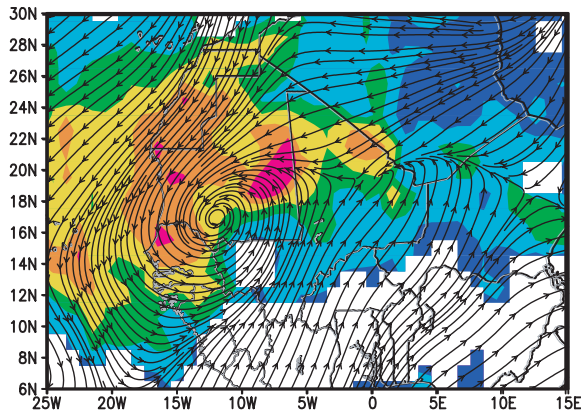
(a) 02 July OMI AI /925 hPa Streamlines



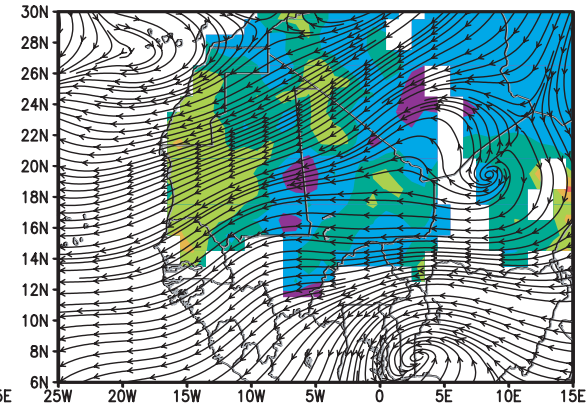
(e) 02 July Deep Blue (AOD)/700 hPa Streamlines



(b) 02 Aug. OMI AI /925 hPa Streamlines



(f) 02 Aug. Deep Blue (AOD)/700 hPa Streamlines



507

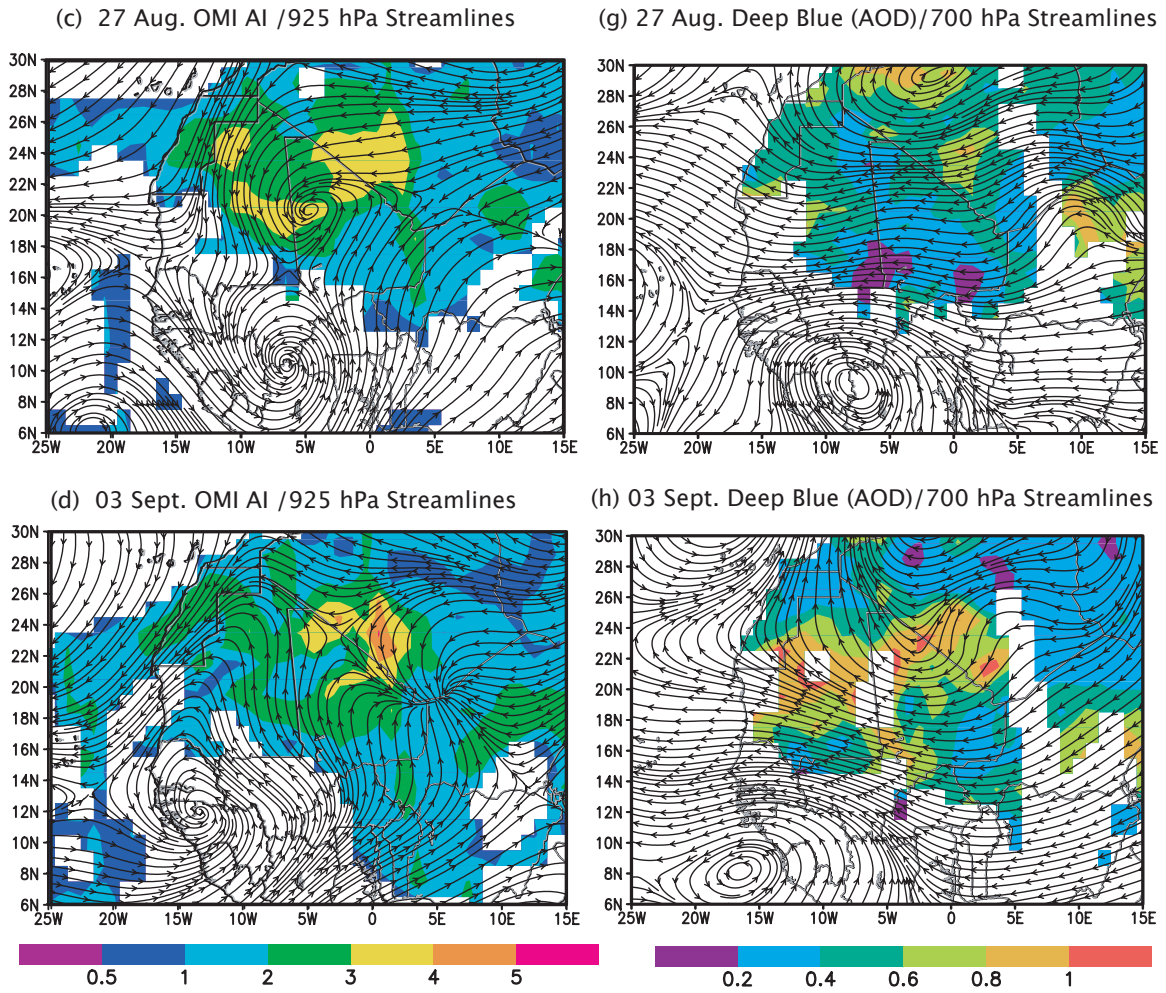
508

509

510

511





512

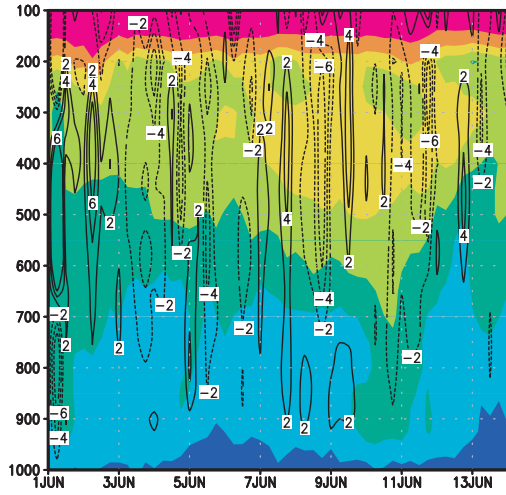
513

514 **Figure 8.** Monsoon OMI AI/925 hPa streamlines and Deep Blue AOT/700 hPa

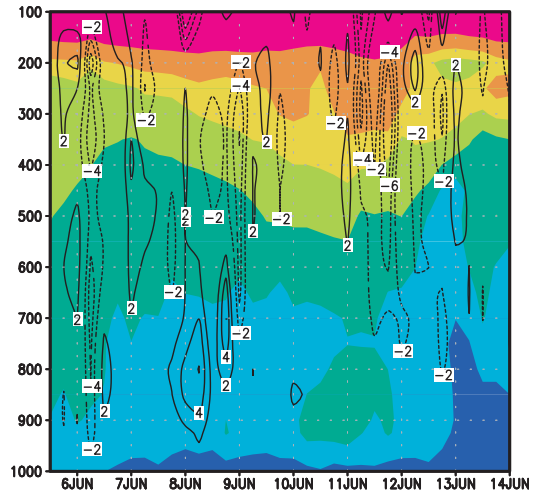
515 streamlines: (a, e) 2 July; (b, f); 2 August; (c, g) 27 August; (d, h) 3 September.

516

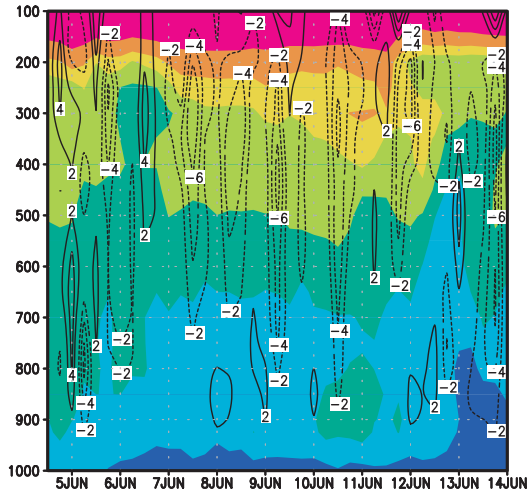
(a) Ozone(ppb) & Vertical Velocity (cm/s)  
Initial 1 June 0000 UTC



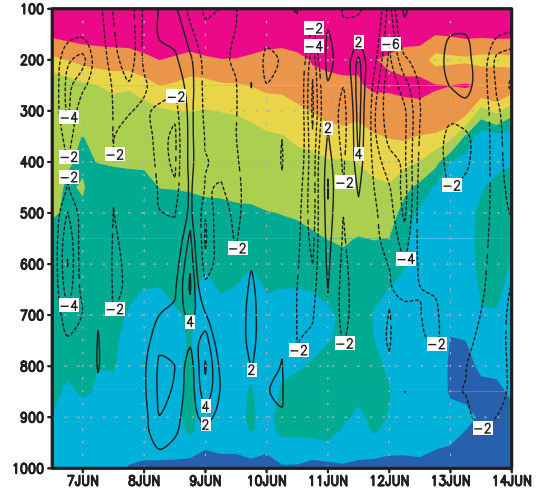
(c) Ozone(ppb) & Vertical Velocity (cm/s)  
Initial 5 June 1200 UTC



(b) Ozone(ppb) & Vertical Velocity (cm/s)  
Initial 4 June 1200 UTC



(d) Ozone(ppb) & Vertical Velocity (cm/s)  
Initial 6 June 1200 UTC



517

518

519

520 **Figure 9.** Time-height profiles of O<sub>3</sub> concentrations and vertical velocity ( $w$ ) at 14.5° N,

521 17.5°W for initial conditions beginning at: (a) 1 June 0000 UTC; (b) 4 June 1200 UTC;

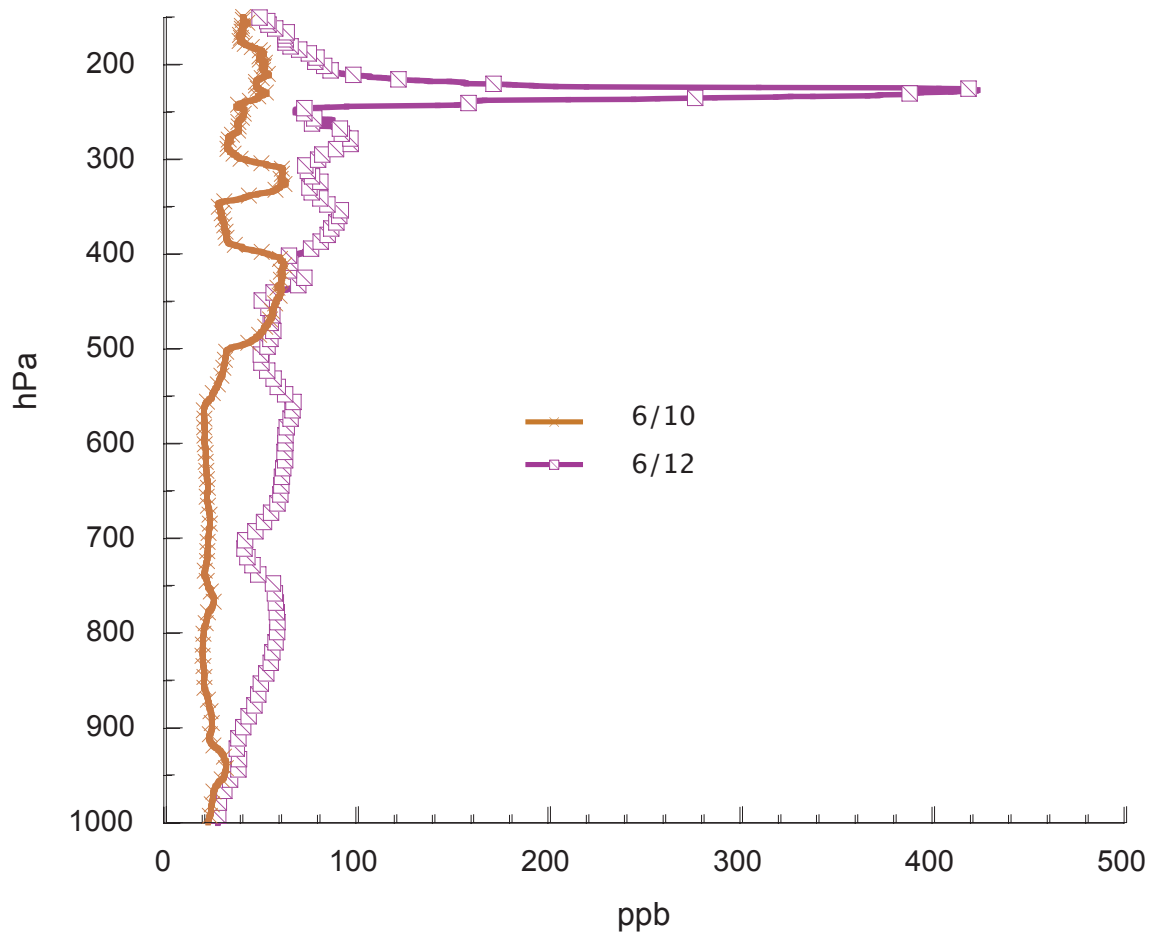
522 (c) 5 June 1200 UTC; (d) 6 June 1200. Units in ppb and cm-s<sup>-1</sup>.

523

524

525

### 1000-150 hPa Ozone Concentrations (ppb) 10, 12 June



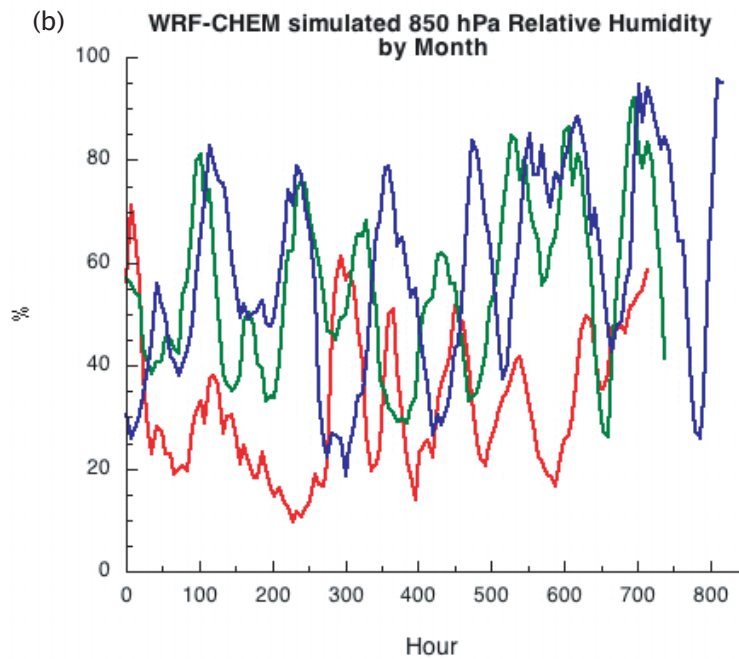
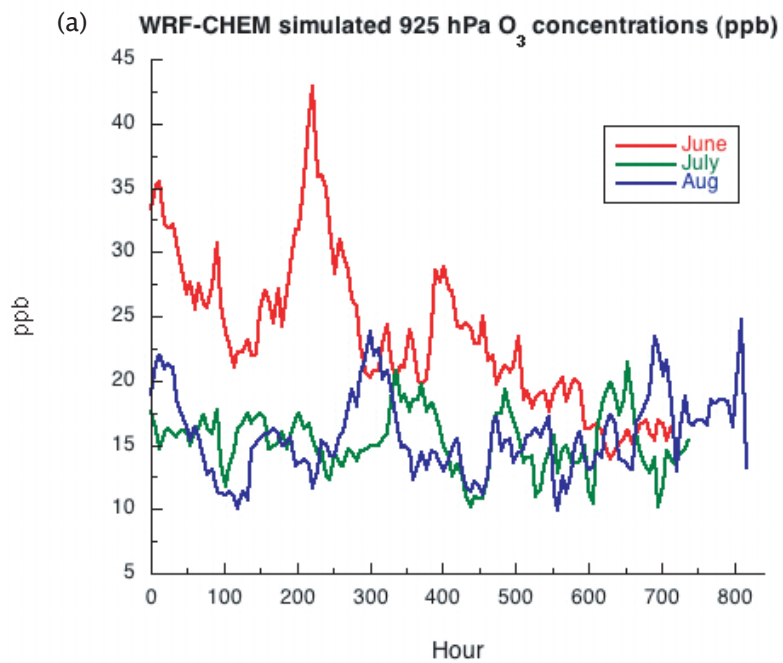
526

527 **Figure 10.** 1000-150 hPa vertical profile of O<sub>3</sub> at Dakar for 10,12 June, 1200 UTC.

528 Units in ppb.

529

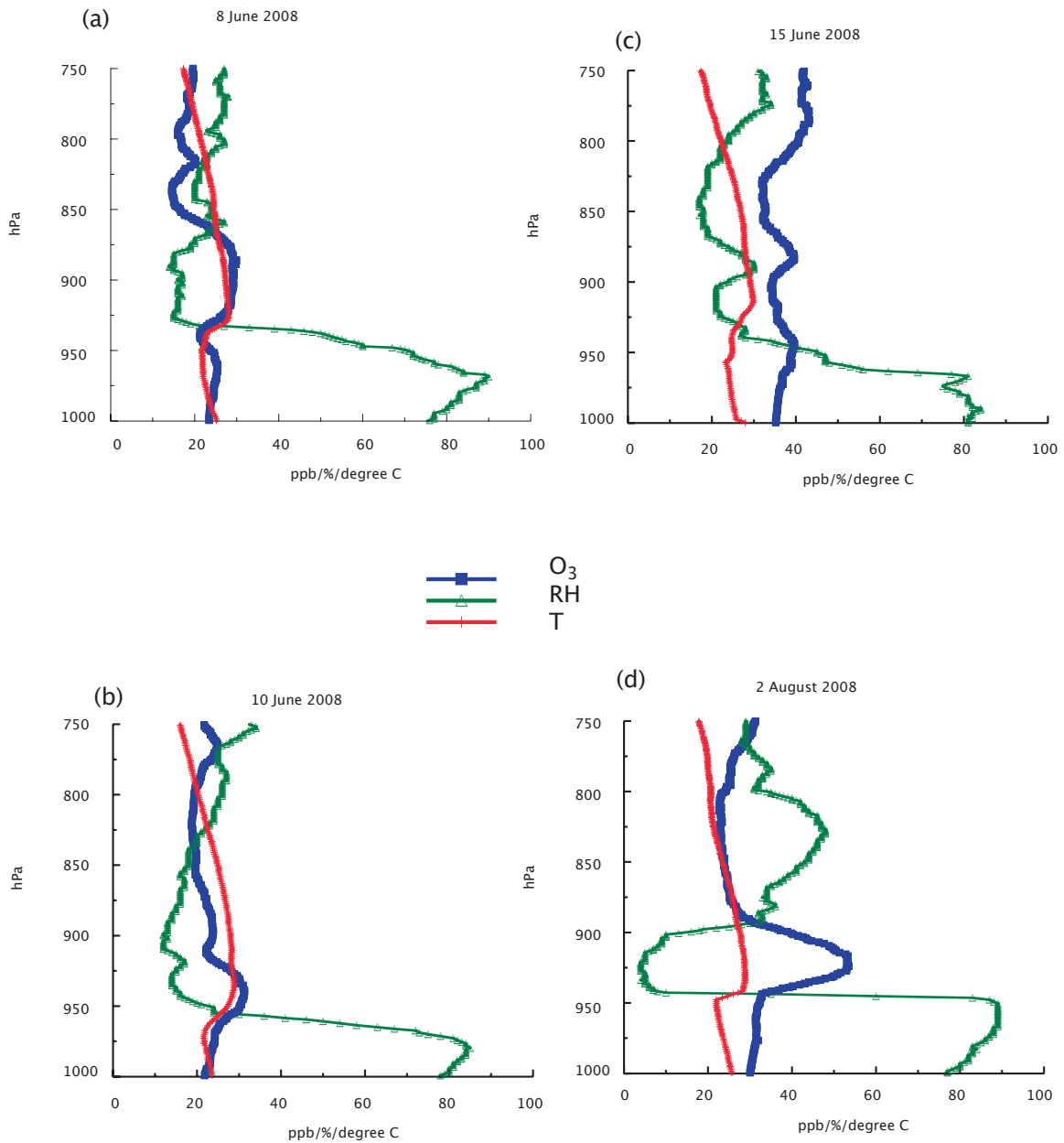
530



531

532 **Figure 11.** WRF-Chem simulation of: (a) 925 hPa O<sub>3</sub> concentrations for June, July and

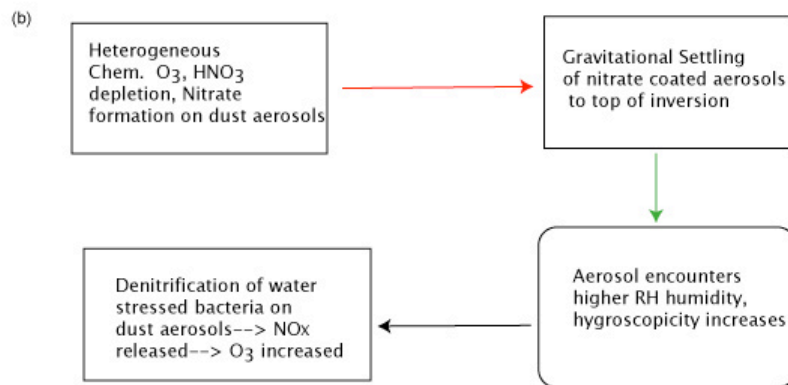
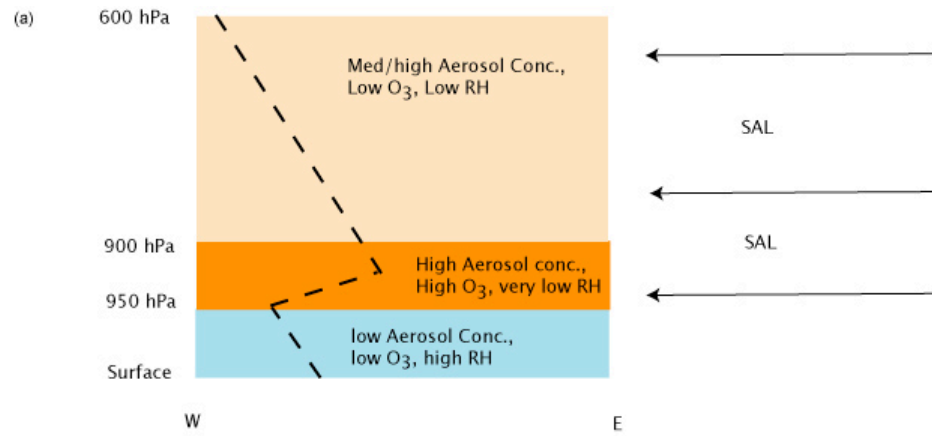
533 August; (b) 850 hPa relative humidity for June, July and August.



535

536

537 **Figure 12.** 1000-750 hPa vertical profiles of O<sub>3</sub> (blue), RH (green) and Temperature  
 538 (red) for identified SAL events. (a) 8 June; (b) 10 June; (c) 15 June; (d) 2 August. Units  
 539 are ppb for O<sub>3</sub>, % for RH and Degree C for temperature.



540

541

542 **Figure 13.** (a) A depiction of relative humidity, aerosol and O<sub>3</sub> concentrations in various  
 543 layer of the lower/middle troposphere associated with a SAL intrusion. The dash line  
 544 represents the vertical profile of temperature. (b) Proposed mechanism for increasing O<sub>3</sub>  
 545 in the 950-900 hPa layer