



1

2 **Development of an OMI AI data assimilation scheme for aerosol modeling over bright**
3 **surfaces—a step toward direct radiance assimilation in the UV spectrum**

4

5 Jianglong Zhang¹, Robert J. D. Spurr², Jeffrey S. Reid³, Peng Xian³, Peter R. Colarco⁴, James R.
6 Campbell³, Edward J. Hyer³, and Nancy Baker³

7

8 ¹Department of Atmospheric Sciences, University of North Dakota, Grand Forks, ND

9

²RT SOLUTIONS Inc., Cambridge MA

10

³Marine Meteorology Division, Naval Research Laboratory, Monterey, CA

11

⁴NASA Goddard Space Flight Center, Greenbelt, MD

12

13

14

15

16

Submitted to

17

GMD

18

19

20

June 2020

21

22

23

24

25 Corresponding Author: jianglong.zhang@und.edu



26

Abstract

27 Using the Vector Linearized Discrete Ordinate Radiative Transfer (VLIDORT) code as the main
28 driver for forward model simulations, a first-of-its-kind data assimilation scheme has been
29 developed for assimilating Ozone Monitoring Instrument (OMI) aerosol index (AI) measurements
30 into the Naval Aerosol Analysis and Predictive System (NAAPS). This study suggests both RMSE
31 and absolute errors can be significantly reduced in NAAPS analyses with the use of OMI AI data
32 assimilation, when compared to values from NAAPS natural runs. Improvements in model
33 simulations demonstrate the utility of OMI AI data assimilation for improving the accuracy of
34 aerosol model analysis over cloudy regions and bright surfaces. However, the OMI AI data
35 assimilation alone does not out-perform aerosol data assimilation that uses passive-based aerosol
36 optical depth (AOD) products over cloud free skies and dark surfaces. Further, as AI assimilation
37 requires the deployment of a fully-multiple-scatter-aware radiative transfer model in the forward
38 simulations, computational burden is an issue. Nevertheless, the newly-developed modeling
39 system contains the necessary ingredients for assimilation of radiances in the ultra-violet (UV)
40 spectrum, and our study shows the potential of direct radiance assimilation at both UV and visible
41 spectrums, possibly coupled with AOD assimilation, for aerosol applications in the future.
42 Additional data streams can be added, including data from Tropospheric Monitoring Instrument
43 (TROPOMI), Ozone Mapping and Profiler Suite (OMPS) and eventually with the Plankton,
44 Aerosol, Cloud and ocean Ecosystem (PACE) mission.

45

46



47 **1.0 Introduction**

48 Operational chemical transport modeling (CTM) of atmospheric aerosol particles,
49 including simulation of sources and sinks and long-range transport of aerosol events such as
50 biomass burning aerosols from fires and dust outbreaks, is now commonplace at global
51 meteorology centers for air quality and visibility forecasts (e.g. Sessions et al, 2015; Lynch et al.,
52 2016). Variational and ensemble-based assimilation of satellite derived aerosol products such as
53 aerosol optical depth (AOD), lidar backscatter measurements, and surface aerosol properties, can
54 substantially improve accuracies in CTM analyses and forecasts (Zhang et al., 2008; 2011; 2014;
55 Yumimoto et al., 2008; Uno et al., 2008; Benedetti et al., 2009; Schutgens et al., 2010; Sekiyama
56 et al., 2010; Saide et al. 2013; Schwartz, 2012; Li et al., 2013; Rubin et al., 2017; Lynch et al.,
57 2016).

58 Currently, the main satellite inputs for operational aerosol modeling are AOD products
59 derived from passive-based polar orbiting imagers, such as the Moderate Resolution Imaging
60 Spectroradiometer (MODIS), the Visible Infrared Imaging Radiometer Suite (VIIRS), and the
61 Advance Very High Resolution Radiometer (AVHRR). Experimentation is proceeding with the
62 use of products from the multi-angle imaging spectroradiometer (MISR) (e.g., Lynch et al., 2016;
63 Randles et al. 2017; Buchard et al. 2017) and from geostationary instruments such as Himawari
64 and Geostationary Operational Environmental Satellite (GOES). A major advantage with such
65 passive-based satellite sensors is that the AOD is retrieved with high spatial and temporal
66 resolutions over relatively broad fields-of-view (e.g. Zhang et al., 2014). For example, MODIS
67 and VIIRS provide near-global daily daytime coverage (e.g. Levy et al., 2013; Hsu et al., 2019)
68 and GOES and Himawari are capable of retrieving AOD over North American and East Asia
69 regions at sub-hourly temporal resolution (e.g. Bessho et al., 2016).



70 To date, these traditional passive-based satellite AOD retrievals have been limited to darker
71 surfaces and relatively cloud-free conditions. The widely-used MODIS Dark Target aerosol data,
72 for instance, are available globally over only oceans and dark land surfaces (e.g. Levy et al., 2013).
73 The MISR and MODIS Deep Blue aerosol products are also available over some arid
74 environments, but are not applicable to snow and ice covered regions (e.g. Kahn et al., 2010; Hsu
75 et al., 2013). Also, none of the above-mentioned aerosol products are valid over cloudy regions.

76 In comparison to AOD, the semi-quantitative UV-based aerosol index (AI) has long been
77 used to monitor major aerosol events such as smoke plumes and dust storms, starting with the
78 Total Ozone Mapping Spectrometer (TOMS) from the late 1970s (Herman et al., 1997). AI is
79 derived using the ratio of observed UV radiances to simulated ones assuming only a clear Rayleigh
80 sky (e.g. Torres et al., 2007). AI retrievals are currently computed using observations from sensors
81 with ozone-sensitive channels. For example, the Ozone Monitoring Instrument (OMI), Ozone
82 Mapping and Profiler Suite (OMPS), TROPOspheric Monitoring Instrument (TROPOMI) and the
83 future Plankton, Aerosol, Cloud and ocean Ecosystem (PACE) mission can detect UV-absorbing
84 aerosol particles, such as black carbon laden smoke or iron-bearing dust, over bright surfaces, such
85 as desert, snow and ice covered regions, and aerosol plumes above clouds (e.g. Torres et al., 2012;
86 Yu et al., 2012; Alfaro-Contreras et al., 2014; 2016).

87 To complement existing AOD assimilating systems, we have developed an AI data
88 assimilation (AI-DA) system that is capable of assimilating OMI AI over bright surfaces and
89 cloudy regions for aerosol analyses and forecasts. This study can be considered as one of the first
90 attempts for direct radiance assimilation in the UV spectrum for aerosol applications, as AI can be
91 directly computed from UV radiances and the developed OMI AI-DA system has all necessary
92 components for a typical radiance assimilation package. In time we expect our assimilation model



93 to merge with AOD or solar radiance assimilation to influence aerosol loading, height and
94 absorption (e.g., VIIRS+OMPS product; such as Lee et al. 2015). Details of the developed OMI
95 AI assimilation system are presented in the paper, which is organized as follows: Data sets used
96 in the study are summarized in Section 2; Section 3 discusses the components of the AI-DA
97 system. Section 4 provides an evaluation of the developed system; and Section 5 contains a
98 summary discussion.

99

100 **2.0 Datasets and Models**

101 Three datasets are used in this study. These are: (i) the OMI level 2 UV aerosol product
102 (OMAERUV; Torres et al., 2007), (ii) the Aerosol Robotic Network (AERONET; Holben et al.,
103 1998) AOD product, and (iii) reanalysis data from the Naval Aerosol Analysis and Prediction
104 System (NAAPS; Lynch et al., 2016), which was the first operational global aerosol mass transport
105 model available to the community. The assimilation system is based on spatial and temporal
106 variations of aerosol particles from NAAPS (Zhang et al., 2006; 2008), and the Vector Linearized
107 Discrete Ordinate Radiative Transfer (VLIDORT; Spurr, 2006) code is used to construct a forward
108 model for the AI-DA system.

109

110 **2.1 OMI aerosol product**

111 UV Aerosol Index data from the OMI level 2 version 3 UV aerosol products (OMAERUV)
112 are used in this study. The OMI instrument is on board the Aura satellite (launched in 2004) and
113 it observes the earth's atmosphere over the UV/visible spectrum with a pixel size of 13x24 km at
114 nadir for the global scan mode, and a swath of ~2600 km (Levelt et al., 2018). The daytime
115 equatorial crossing for the Aura platform is ~1:30 p.m. The dataset comprises the UV AI, viewing



116 and solar geometries, spectrally-dependent surface albedos at the 354 and 388 nm spectral
117 channels, terrain pressure, geolocations, x-track and algorithm quality flags, plus other aerosol and
118 ancillary parameters. The UV AI is designed to detect UV-absorbing aerosol particles, and is
119 based on radiance observations at 354 nm (I_{obs354}) and calculated radiance (I_{cal354}) at 354 nm for a
120 Rayleigh (no aerosol) atmosphere (e.g. Torres et al., 2007) as defined as

$$121 \quad AI = -100 \log_{10} \frac{I_{obs354}}{I_{cal354}}. \quad (1)$$

122 Unbiased, noise-reduced, quality-assured AI data are necessary for AI data assimilation.
123 This is especially important for OMI observations, due to this particular sensor suffering from the
124 well-referenced “row anomalies” issues (Torres et al., 2018). To remove pixels with row
125 anomalies, only retrievals with x-track flag values of 0 are retained. Also, abnormal AI values
126 were identified over mountain regions. Thus, retrievals with terrain/surface pressure less than 850
127 hpa are excluded in the study. Finally, only retrievals with OMI AI values larger than -2 are used.
128 Therefore, OMI observations over cloudy skies, which could have negative OMI AI values, are
129 also included.

130 Both cloud-free and above-cloud AI data satisfying these quality checks are aggregated /
131 averaged in $1 \times 1^\circ$ (Latitude/Longitude) bins. As a radiative transfer model run is applied for each
132 observation, the gridded data are used in the assimilation process in order to reduce the
133 computational burden. Averaged parameters for the gridded data include the solar and sensor
134 zenith angles, the relative azimuth angles, the spectrally-dependent surface albedos at 354 and 388
135 nm, the cloud fraction, and the AI values themselves. Additional quality assurance steps are also
136 applied during the spatial-averaging process. Isolated high AI values are removed as follows.
137 First, for a 4×4 pixel box, if the mean AI is less than 0.7 but an individual AI value is larger than



138 0.7, then that one value is removed. Second, if the standard deviation of AI values for a 3x3 pixel
139 box surrounding a pixel is larger than 0.5, that individual AI value is likewise removed.

140

141 **2.2 AERONET data**

142 Version 3 level 2 daytime, cloud-cleared and quality-assured AERONET data are used to
143 evaluate the performance of the OMI AI data assimilation in our study (Holben et al., 1998; Giles
144 et al., 2019). During daytime, AOD from AERONET instruments are derived by measuring the
145 attenuated solar radiance typically at seven wavelengths ranging from 340 to 1020 nm. In this
146 study, AERONET data are collocated with NAAPS analyses with and without OMI AI
147 assimilation. In order to collocate AERONET and NAAPS AOD data, AERONET AOD values
148 within ± 30 minutes of a given NAAPS analysis time are averaged and used as ground-based AOD
149 values for the NAAPS $1 \times 1^\circ$ (Latitude/Longitude) collocated bins. As AERONET data require a
150 cloud-free line of sight to the solar disk, the performance of OMI AI data assimilation over overcast
151 regions is not evaluated.

152

153 **2.3 NAAPS and NAAPS reanalysis data**

154 The NAAPS (<http://www.nrlmry.navy.mil/aerosol/>) model is a multi-species, three-
155 dimensional, Eulerian global transport model using operational Navy Global Environmental
156 Model (NAVGEM) as the meteorological driver (Hogan et al., 2014). NAAPS provides 6-day
157 forecasts at a 3-hour interval with a spatial resolution of $1/3^\circ$ (latitude/Longitude) and 42 vertical
158 levels on a global scale. NAAPS predicts four aerosol particle classes: anthropogenic and biogenic
159 fine particles (ABF, such as primary and secondary organic aerosols and sulfate aerosols); dust,
160 biomass burning smoke; and sea salt (Lynch et al, 2016).



161 The 2003–2018 NAAPS reanalysis version 1 (v1) (Lynch et al., 2016) is a modified version
162 of the operational NAAPS model. In this version, quality-controlled retrievals of AOD from
163 MODIS and MISR (Zhang et al., 2006; Hyer et al., 2011; Shi et al., 2014) are assimilated into
164 NAAPS through the Naval Research Laboratory Atmospheric Variation Data Assimilation
165 System-AOD system (NAVDAS-AOD; e.g., Zhang et al., 2008; Zhang et al., 2011; Zhang et al.,
166 2014). Aerosol source functions, including biomass burning, smoke and dust emissions, are tuned
167 regionally based on the AERONET data. Other aerosol processes, including dry deposition over
168 water, are also tuned based on AOD data assimilation correction fields. NOAA Climate Prediction
169 Center (CPC) MORPHing (CMORPH) precipitation data are used to constraint the wet removal
170 process within the tropics (Joyce et al., 2004). The usage of CMORPH avoids the ubiquitous
171 precipitation bias that exists in all global atmospheric models (e.g. Dai, 2006) and is proven to
172 improve aerosol wet deposition, therefore yielding better AOD (Xian et al., 2009). The reanalysis
173 agrees reasonably well with AERONET data on a global scale (Lynch et al., 2016) and also
174 reproduces AOD trends that are in a good agreement with satellite based analysis (e.g., Zhang and
175 Reid, 2010; Hsu et al., 2012). In this study, we use a free running version of NAAPS reanalysis v1
176 without AOD assimilation to provide aerosol fields every 6 hours at $1^\circ \times 1^\circ$ (Latitude/Longitude)
177 resolution.

178

179 **2.4 VLIDORT radiative transfer code**

180 VLIDORT is a linearized, multiple-scatter radiative transfer model for the simultaneous
181 generation of Stokes 4-vectors and analytically-derived Jacobians (weighting functions) of these
182 4-vectors with respect to any atmospheric or surface property (Spurr, 2006). The model uses
183 discrete-ordinate methods to solve the polarized plane-parallel RT equations in a multi-layer



184 atmosphere, plus the solution of a boundary value problem and subsequent source-function
185 integration to obtain radiation fields at any geometry and any atmospheric level. VLIDORT has a
186 “pseudo-spherical” *ansatz*: the treatment of solar-beam attenuation in a spherical-shell atmosphere
187 before scattering. Single-scattering in VLIDORT is accurate for both line-of-sight and solar-beam
188 spherical geometry. The model has a full thermal emission capability. VLIDORT has two
189 supplements, one dealing with bidirectional (non-Lambertian) reflection at the surface, and the
190 other with the inclusion of surface light sources (SIF or water-leaving radiances). Full details on
191 the VLIDORT model may be found in a recent review paper (Spurr and Christi, 2019, and
192 references to VLIDORT therein).

193 VLIDORT is used to simulate the AI in this study. Simulations at 354 and 388 nm are
194 performed both for Rayleigh atmospheres, and for scenarios with aerosol loadings (four mass-
195 mixing profiles for different aerosol types) taken from the NAAPS model. In addition to the AI,
196 Jacobian calculations are needed with respect to these aerosol profiles. Firstly, radiance Jacobians
197 with respect to these four mass-mixing profiles are computed analytically using VLIDORT’s
198 linearization facility, and secondly the associated Jacobians of AI are further derived through a
199 second VLIDORT linearization with respect to the Lambertian-equivalent reflectivity. The details
200 of this process is given in the next section

201

202 **3.0 OMI AI assimilation system**

203 The OMI assimilation system has three components: a forward model, a 3-D variational
204 assimilation system, and a post-processing system. Based on the background NAAPS 3-D aerosol
205 concentrations for dust, smoke, ABF, and sea salt aerosols, the forward model not only computes
206 the associated AI values, but also their Jacobians of AI with respect to the four aerosol mass-



207 loading profiles. The 3-D variational assimilation system is a modified 3-D AOD system (Zhang
208 et al., 2008; 2011; 2014) that computes increments for dust and smoke aerosol concentrations
209 based on OMI AI data. The post-processing system constructs a new NAAPS analysis based on
210 the background NAAPS aerosol concentrations and increments as derived from the 3-D variational
211 assimilation system. Details of the forward model and the modified NAVDAS-AOD system are
212 described in this section.

213

214 **3.1 Forward model for simulating OMI AI**

215 To construct an AI-DA system, a forward model is needed to simulate AI using aerosol
216 concentrations from NAAPS. In this study, the forward model is built around the VLIDORT
217 model, following a similar method to that suggested in Buchard et al. (2015). Here VLIDORT is
218 configured to compute OMI radiances and Jacobians as functions of the observational conditions
219 at 354 and 388 nm, using geolocation information from OMI data such as satellite zenith, solar
220 zenith and relative azimuth angles, as well as ancillary OMI data (surface albedos at 354 and 388
221 nm).

222 To convert from NAAPS mass-loading concentrations to aerosol extinction and scattering
223 profiles, we require aerosol optical properties for the four species at 354 and 388 nm, which are
224 summarized in Table 1. The optical properties of ABF (assumed to be sulfate in this study), sea
225 salt, dust and smoke aerosols, including mass extinction cross sections and single scattering
226 albedos at 354 and 388 nm are adapted from NASA's Goddard Earth Observing System version 5
227 (GEOS-5) model (e.g. Colarco et al., 2014; Buchard et al., 2015). Note that the study period is
228 July and August of 2007 over Africa, coinciding with the early biomass burning season associated
229 with lower single scattering albedo values (Eck et al., 2013). With that in mind, we choose a quite



low value of 0.85 for the single-scattering albedo value at 354nm (e.g. Eck et al., 2013; Cochrane et al., 2019). A slightly higher single scattering albedo of 0.86 is assumed at 388 nm. The slight increase in single scattering albedo from 354 to 388 nm has also been observed from Solar Spectral Flux Radiometer (SSFR) observations during the recent NASA Observations of Clouds above Aerosols and their interactions (ORACLES) Campaign (Pistone et al., 2019). Scattering matrices for dust, smoke, sea salt and sulfate (to represent ABF) aerosols are based on associated expansion coefficients (e.g. Colarco et al., 2014; Buchard et al., 2015) taken from NASA's GEOS-5 model. Also to reduce computational expenses, scalar radiative transfer calculations are performed.

To simulate OMI AI, the Lambertian Equivalent Reflectivity (LER) at 388 nm (R_{388}) is needed for estimating LER at 354 nm. The R_{388} is calculated from VLIDORT, based on equation 2 below, adapted from Buchard et al. (2015), or

$$R_{388} = \frac{I_{aer388}(\rho_{388}) - I_{ray388}(0)}{T + S_b(I_{aer388}(\rho_{388}) - I_{ray388}(0))} \quad (2)$$

$I_{ray388}(0)$ is the calculated path radiance at 388 nm assuming a Rayleigh atmosphere with surface albedo 0. T and S_b are the calculated transmittance and spherical albedo at 388 nm. $I_{aer388}(\rho_{388})$ is the computed radiance including 3-D aerosol fields from NAAPS and the 388 nm surface albedo from OMI data. In Buchard et al. (2015), an adjusting factor is applied to R_{388} by adding the difference between climatological surface albedos at 354 and 388 nm. The similar approach is also adopted in this study, as shown in their Equation 3.

$$R'_{388} = R_{388} - (\rho_{388} - \rho_{354}) \quad (3)$$

Here, R'_{388} is surface albedo adjusted Lambertian Equivalent Reflectivity at 388 nm. ρ_{388} and ρ_{354} are surface albedo values at 388 and 354 nm channels that are obtained from the OMI OMAERUV data. Finally, the simulated AI (AI_{naaps}) is given by

$$AI_{naaps} = -100 \log_{10} \frac{I_{aer354}(\rho_{354})}{I_{ray354}(R'_{388})} \quad (4)$$



253 Here, $I_{\text{aer}354}(\rho_{354})$ is the calculated radiance at 354 nm using NAAPS aerosol fields as well as the
254 OMI-reported surface albedo at 354 nm (ρ_{354}). $I_{\text{ray}354}(R_{388})$ is the calculated radiance assuming a
255 Rayleigh atmosphere and the derived value of R_{388} as surface albedo (Buchard et al., 2015).

256 The forward model-simulated OMI AI values are inter-compared with OMI AI values as
257 shown in Figure 1 for the study region. A total of one month (01-31 July 2007) of NAAPS
258 reanalysis data and OMI AI data were used. Note that OMI AI data over both cloud-free and
259 cloudy skies were used. Since surface albedos included in the OMI data represent reflectivities
260 under clear-sky situations, the albedo under cloudy sky is then computed

$$261 \quad \rho_{\text{cld}} = \rho_{\text{clr}} * (1 - f_c) + 0.8 * f_c \quad . \quad (5)$$

262 Here, ρ_{clr} and f_c are the clear sky surface albedo (e.g. ρ_{354} or ρ_{388}) and the cloud fraction, both
263 quantities obtained from the OMI dataset. Clouds are assumed to be tropospheric (close to the
264 surface) with an UV albedo of 0.8, such that this equation applies to both the 354 and 388 nm
265 channels.

266 Figure 1a shows the spatial distribution of NAAPS AOD over Central and North Africa,
267 using collocated NAAPS and OMI AI datasets. OMI AI data are grid-averaged in $1^\circ \times 1^\circ$
268 (latitude/longitude) bins. Also, we focus over Africa in this paper as this area includes dust plumes
269 over deserts and smoke plumes overlying stratus cloud decks. The Arctic is not included as
270 additional efforts may be needed to fully understand properties of sea ice reflectivity; we leave this
271 topic for a future paper. Only bins that have valid NAAPS and OMI AI data are used to generate
272 Figure 1. Dust plumes are visible over North Africa and the Persian Gulf, and a smoke plume from
273 Central Africa is also evident. These UV-absorbing aerosol plumes are also captured by OMI AI,
274 as seen in Figure 1c. Shown in Figure 1b are the simulated OMI AI using the NAAPS aerosol
275 fields and viewing geometries and surface albedos from OMI. The simulated OMI AI shows



276 similar patterns to those derived from OMI, especially for the dust plums over North Africa and
 277 smoke plumes over Central Africa. An overall correlation of 0.785 is found between simulated
 278 and satellite-retrieved OMI AI values, as shown in Figure 1, suggesting the forward model is
 279 functioning reasonably as designed.

280

281 3.1 Forward model for Jacobians of AI

282 Jacobians of OMI AI with respect to aerosol mass concentrations are needed for the OMI
 283 AI assimilation system. In this study, AI Jacobians (K) are calculated from radiance Jacobians
 284 with respect to aerosol mass concentrations for four aerosol species (smoke, dust, ABF/sulfate,
 285 sea-salt) at 354 nm ($K_{354,nk} = \frac{\partial I_{aer354}}{\partial M_{nk}}$) and 388 nm ($K_{388,nk} = \frac{\partial I_{aer388}}{\partial M_{nk}}$) wavelengths. Here M_{nk}
 286 is the mass concentration for aerosol type, k , and for vertical layer, n . I_{aer354} and I_{aer388} are radiances
 287 for the 354 and 388 nm channels, respectively. $K_{354,nk}$ and $K_{388,nk}$ are the corresponding radiance
 288 Jacobians at 354 and 388 nm, respectively. AI Jacobians can then be calculated by analytic
 289 differentiation of the basic formula in Equation (1), and, after some algebra, we find the following
 290 result:

$$291 \quad \frac{\partial AI}{\partial M_{nk}} = \mathcal{A}_1 K_{354,nk}(\rho_{354}) + \mathcal{A}_2 K_{388,nk}(\rho_{388}) \quad . \quad (6)$$

292 Here, \mathcal{A}_1 and \mathcal{A}_2 are given respectively by Equations (7) and (8), as

$$293 \quad \mathcal{A}_1 = \left(-\frac{100}{I_{aer354}(\rho_{354}) \times \ln 10} \right) \quad , \text{ and} \quad (7)$$

$$294 \quad \mathcal{A}_2 = \left(-\frac{100}{I_{ray354}(R_{388}) \times \ln 10} \right) \frac{\partial I_{ray354}(R_{388})}{\partial R} \left[\frac{(1 - S_{388} R_{388})^2}{T_{388}} \right] \quad . \quad (8)$$

295 Based on these equations, radiance Jacobians with respect to aerosol particles, $K_{354,nk}$ and $K_{388,nk}$,
 296 are computed at 354 and 388 nm, respectively, using OMI-reported surface albedo values (ρ_{354}
 297 and ρ_{388}), followed by a calculation of the albedo Jacobian $\frac{\partial I_{aer354}(R_{388})}{\partial R}$ at 354 nm.



298 To check this analytic Jacobian calculation in Eqns. (6)-(8), we compute the aerosol AI
299 Jacobians using a finite difference (FD) method. Here, the derivative of AI as a function of aerosol
300 concentration of a species, k , in layer n , is computed using

$$301 \quad \frac{\partial AI}{\partial M_{nk}} = \frac{(AI - AI')}{(C_{nk} - C'_{nk})} \quad . \quad (9)$$

302 Here C_{nk} and C'_{nk} are the baseline and perturbed aerosol concentrations, respectively, and AI and
303 AI' are computed using C_{nk} and C'_{nk} , respectively.

304 Figure 2b shows the comparison of Jacobians of dust aerosols estimated from the analytic
305 and the FD solutions. Dust, smoke, ABF and sea salt aerosol concentrations as a function of
306 altitude are shown in Figure 2a. To compute FD Jacobians with respect to dust aerosols, a 10%
307 perturbation is introduced in the dust profiles. A very close match is found between analytic and
308 FD Jacobians. This validates the analytical solution used in the study. The analytic solution is of
309 course much faster, as a single call to VLIDORT will deliver all necessary Jacobians at one
310 wavelength, as compared to 97 separate calls to VLIDORT with the FD calculation (baseline; 4
311 species perturbations in the 24-layer atmosphere).

312

313 **3.2 The variational OMI AI assimilation system**

314 The OMI AI assimilation system is based on AI simulations (with Jacobians) from the
315 forward model. Two principles underlay the assimilation procedure. First, we assume that OMI AI
316 is sensitive to UV-absorbing aerosol particles, such as NAAPS smoke and dust, or that only smoke
317 and dust are injected high enough into the troposphere to impact AI. Therefore, innovations are
318 limited to modifications of dust and smoke aerosol properties. For classes that do not strongly
319 project onto AI, such as sea salt and ABF aerosols, aerosol concentrations are not modified during
320 the process. Second, contributions of smoke/dust aerosols to AI ($AI_{\text{smoke}} / AI_{\text{dust}}$) prior to



321 assimilation are estimated by multiplying smoke/dust aerosol concentrations from NAAPS with
 322 Jacobians of AI respective of smoke/dust aerosols. The ratio of AI innovation from smoke aerosols
 323 (ΔAI_{smoke}) to total AI innovation (ΔAI or $OMI\ AI - AI_{\text{naaps}}$) is assumed to be the ratio of AI_{smoke} to
 324 $AI_{\text{smoke}} + AI_{\text{dust}}$. The same assumption holds for dust aerosols.

325 Given these two principles, the overall design concept for the OMI AI assimilation can be
 326 expressed as

327

328 $C^a = C^b +$

329
$$\frac{P_{dust} H_{dust}^T}{H_{dust}^T P_{dust} H_{dust} + R} [y - H(C^b)] \times \frac{H_{dust} C_{dust}^b}{H_{dust} C_{dust}^b + H_{smk} C_{smk}^b} +$$

330
$$\frac{P_{smk} H_{smk}^T}{H_{smk}^T P_{smk} H_{smk} + R} [y - H(C^b)] \times \frac{H_{smk} C_{smk}^b}{H_{dust} C_{dust}^b + H_{smk} C_{smk}^b}, \quad (10)$$

331

332 where C^b and C^a are NAAPS aerosol concentrations for the analysis and background fields,
 333 respectively, C_{dust}^b and C_{smk}^b are background NAAPS particle mass concentrations for dust and
 334 smoke, $H(C)$ is the NAAPS forward model that links NAAPS particle mass concentrations to AI,
 335 and H is defined as $\partial H(C)/\partial C$, which is the Jacobian matrix of AI with respect to aerosol
 336 concentrations. Y is the observed OMI AI, and $Y - H(C^b)$ is the innovation of AI, representing the
 337 difference between observed and modeled AI values.

338 The $\frac{H_{dust} C_{dust}^b}{H_{dust} C_{dust}^b + H_{smk} C_{smk}^b}$ and $\frac{H_{smk} C_{smk}^b}{H_{dust} C_{dust}^b + H_{smk} C_{smk}^b}$ terms are the fractional contribution
 339 of innovation from dust and smoke aerosol, respectively. These terms are estimated using NAAPS
 340 aerosol concentrations for relatively high aerosol loading cases ($AOD > 0.15$). For low aerosol
 341 loading ($AOD < 0.15$) as reported from NAAPS, it is possible that NAAPS could underestimate
 342 aerosol concentrations. Thus, the fractional contribution of innovations is assigned to 1 for the



343 dominant aerosol type based on a NAAPS aerosol climatology (Zhang et al., 2008). Note that the
 344 term $[y-H(C^b)] \times \frac{H_{dust} C_{dust}^b}{H_{dust} C_{dust}^b + H_{smk} C_{smk}^b}$ is in observational space. P_{dust} and P_{smk} are model error
 345 spatial covariance matrices for dust and smoke (model space) aerosols (e.g. Zhang et al., 2008;
 346 2011; 2014). R is the observation-based error covariance in model space. The
 347 $\frac{P_{dust} H_{dust}^T}{H_{dust}^T P_{dust} H_{dust} + R} [y-H(C^b)] \times \frac{H_{dust} C_{dust}^b}{H_{dust} C_{dust}^b + H_{smk} C_{smk}^b}$ and $\frac{P_{smk} H_{smk}^T}{H_{smk}^T P_{smk} H_{smk} + R} [y-H(C^b)] \times$
 348 $\frac{H_{smk} C_{smk}^b}{H_{dust} C_{dust}^b + H_{smk} C_{smk}^b}$ terms represent the estimated increments in model space.

349 The background error covariance matrix is constructed from modeled error variances and
 350 error correlations, following the methodology in previous studies (Zhang et al., 2008; 2011). The
 351 horizontal background error covariance is generated using the second-order regressive function
 352 (SOAR), as shown in Equation 11 (Zhang et al., 2008), or

$$353 \quad C(x, y) = (1 + R_{xy}/L) \exp(-\frac{R_{xy}}{L}) \quad (11)$$

354 Here, x and y are two given locations, and R_{xy} is the great circle distance. L is the averaged error
 355 correlation length and is set to 200 km based on Zhang et al. (2008). Similarly, the vertical error
 356 correlation between two pressure levels p_1 and p_2 is also based on the SOAR function, this time in
 357 pressure space, based on Zhang et al., 2011, is

$$358 \quad C(p_1, p_2) = [1 + \left| \int_{p_1}^{p_2} \frac{d \ln p}{L} \right|] e^{-\left| \int_{p_1}^{p_2} \frac{d \ln p}{L} \right|} \quad (12)$$

359 Here, L is a unit-less number representing vertical correlation length and is set to 0.015.

360 The horizontal error variance is based on the RMS error of aerosol concentrations, which
 361 is arbitrarily set to $100 \mu\text{g}/\text{m}^3$ for near-surface dust aerosols (ground to 700 hPa). The RMS error
 362 of dust aerosol mass is assumed to decrease as altitude increases, and is set to 50%, 25%, and 1%
 363 of the near-surface values for 500-700, 350-500 and 70-350 hPa respectively. Note that different



364 aerosol species have different mass extinction values. Here we assume the modeled error in
365 aerosol extinction is the same for different aerosol species and thus, the RMS error of smoke
366 aerosol concentration is scaled by mass extinction cross section ratio between smoke and dust
367 aerosols. The observational errors are assumed to be non-correlated in this study (e.g. Zhang et
368 al., 2008). OMI AI values over cloud-free and cloudy skies are used in the study and therefore,
369 RMS errors of AI are required for both these situations. Note, as suggested by Yu et al. (2012),
370 for the same above cloud CALIOP AOD, variations in AI are found to be of the order of 1 for
371 cloud optical depth changing from 2 to 20. Thus, we assume the RMS error of OMI AI is 0.5 for
372 cloud-free skies, increasing linearly with cloud fraction up to a value of 1 for the 100% overcast.

373 Lastly, we assume that detectable UV absorbing aerosols have AI values larger than 0.8
374 (e.g. Torres et al., 2013). Therefore, for regions with OMI AI values larger than 0.8, UV absorbing
375 aerosol particles can both be added or removed from air columns based on innovations, which are
376 the differences between OMI reported and simulated AI values. For regions with OMI AI values
377 less than 0.8, innovations are only used to remove UV absorbing aerosol particles from air
378 columns.

379

380 **4.0 System evaluation & discussion**

381 **4.1 Evaluating the performance of the AI assimilation system over Africa**

382 Using two months of OMI data (July-August, 2007), the performance of OMI AI
383 assimilation was evaluated around the Africa region (20°S-40°N; 10°W-60°E). The study region
384 was chosen to examine the performance of OMI AI data assimilation over bright surfaces such as
385 the deserts of North Africa, as well as study aerosol advection over clouds, in this case smoke off
386 the west coast of Southern Africa. In this demonstration, two NAAPS runs were performed for



387 the period of July 1 to August 31, 2007, one with and one without the use of OMI AI assimilation
388 (AI-DA run). Both runs were initialized with the use of NAAPS reanalysis data at 0000 UTC 1
389 July and do not include any other form of aerosol assimilation.

390 Figure 3a shows the true color composite from Aqua MODIS for July 28, 2007 over the
391 study region that is obtained from the NASA world view site
392 (<https://worldview.earthdata.nasa.gov/>; last accessed June 2020). Visible in the image are the dust
393 plumes from North Africa transported to the Atlantic Ocean, and smoke plumes from Central and
394 Southern Africa transported to the west coast of South Africa. As indicated by the aggregated
395 OMI AI data for 1200 UTC 28 July 2007 (Figure 3b), dust plumes from North Africa are
396 transported to the North corner of the west coast of North Africa. Smoke plumes are also visible
397 in the OMI AI plot in Southern Africa and are transported to the west coast and over the Atlantic.
398 Comparing Figure 3a and Figure 3b, smoke plumes, as identified from OMI, are also found over
399 cloudy regions as indicated from the MODIS visible imagery. Note that Figure 3b shows the OMI
400 AI data used in the assimilation process and again, AI retrievals over both cloud free and cloudy
401 conditions are included as suggested by Figure 3b.

402 Figure 3c is the 1200 UTC 28 July 2007 NAAPS AOD product from the natural run. In
403 comparison, Figure 3d shows the same situation, this time with the use of OMI AI data
404 assimilation. Comparing 3b with 3d, dust and smoke aerosol patterns as shown from OMI AI
405 resemble more closely the NAAPS AOD fields after AI assimilation. Over the northeast coast of
406 Africa, heavy aerosol plumes, as hinted at in NAAPS AOD from the natural run, cover larger
407 spatial areas than those inferred from OMI AI data. In comparison, NAAPS AOD patterns from
408 the OMI AI data assimilation cycle closely resemble aerosol patterns as suggested from OMI AI
409 data. Also shown in Figures 3e and 3f are the simulated AI using NAAPS data from the natural



410 and OMI AI DA runs (data from Figures 3c and 3d) respectively. Clearly, with the use of NAAPS
411 data from the natural run, simulated OMI AI are overestimated in comparison with OMI AI data
412 (Figure 3b). Simulated AI patterns with the used of NAAPS data from the OMI AI DA run rather
413 closely resemble AI patterns from the OMI data, again, indicating the OMI AI DA system is
414 functioning reasonably as designed.

415 The performance of AI-DA is also evaluated using OMI AI for the whole study period, as
416 shown in Figure 4. These data are constructed using collocated OMI AI and NAAPS data
417 according to the conditions introduced in Sec. 3. Here, Figures 4a and 4e are spatial distributions
418 of two-monthly averaged (July and August 2007) AODs for NAAPS AI-DA and natural runs,
419 respectively. Figure 4b is the spatial distribution of the simulated AI using NAAPS data from AI-
420 DA runs, and Figure 4c is the spatial distribution of OMI AI for the two-month period. Figures 4f
421 and 4g show similar plots to those in Figures 4c and 4d, but this time for NAAPS natural runs.
422 While simulated AI values from NAAPS natural runs (Figure 4f) are overestimated compared to
423 OMI AI values (Figure 4g) for the study region, the patterns of simulated AI from NAAPS AI-DA
424 runs (Figure 4b) are similar to patterns shown from OMI AI (Figure 4c). This is also seen from
425 Figure 4d, which is the difference between simulated AI from NAAPS AI-DA runs and OMI AI.
426 In contrast with the situation in Figure 4d, Figure 4h, which is the difference between simulated
427 AI from NAAPS natural runs and OMI AI, shows much larger differences in AI values.

428 While it is not too difficult to make the model mimic the AI product, proof of real skill lies
429 in any improvements to AOD calculations. To this end, the performance of OMI AI assimilation
430 was evaluated with the use of AERONET data. Figure 5a shows the inter-comparison of NAAPS
431 AOD versus AERONET AOD at 0.55 μm . A total of 1450 collocated pairs of NAAPS and
432 AERONET data were compiled for the study region over the two months test period. Comparing



433 with AERONET data, NAAPS AOD from the natural run had a correlation of 0.64, a mean
434 absolute error in AOD of 0.17, and an RMSE of 0.25. In comparison, with AI assimilation,
435 NAAPS AOD correlations to AERONET increased to 0.72 (Figure 5b), the absolute error reduced
436 to 0.12, and RMSE reduced to 0.18, both roughly a 30% reduction. Note that AERONET AOD
437 values are only available for lines-of-sight that are free of cloud presence for the sun photometer
438 instruments. Also, the slope of AERONET versus NAAPS AOD is 0.93 for the NAAPS natural
439 runs, and a similar slope of 0.92 is found for the NAAPS AI-DA runs.

440

441 **4.2 Inter-comparison with AOD data assimilation**

442 Typically, NAAPS reanalyses are constructed through assimilation of MISR and MODIS
443 aerosol products (NAAPS AOD assimilation). Thus, the performances of NAAPS AOD and AI-
444 DA assimilations are compared against AERONET data. Figure 5c shows the comparison of
445 AERONET AOD and NAAPS AOD after AOD assimilation, while Figure 5b shows a similar plot
446 but using NAAPS data from AI-DA. A better correlation between AERONET and NAAPS data
447 of 0.82 and a slope of 1.01 are found using AOD data assimilation. In comparison, the correlation
448 is 0.72 and the slope is 0.92 for the AI-DA runs. Slightly better RMSE (0.15 versus 0.18) and
449 absolute error (0.11 versus 0.12) values are also found for the AOD data assimilation runs. This
450 result is not surprising as OMI AI provides only a proxy for aerosol properties while passive-based
451 AOD retrievals are often considered as a more reliable parameter for representing column-
452 integrated aerosol properties. But still, the evaluation efforts are over cloud-free line-of-sight as
453 detected from AERONET, AI DA may further assist traditional AOD data assimilation by proving
454 AI assimilation over cloudy regions.

455

456 **4.3 Sensitivity test**



457 As mentioned in Section 3, aerosol properties for non-smoke aerosol types were obtained
458 from the NASA GEOS-5 model (e.g. Colarco et al., 2014; Buchard et al., 2015). Yet, different
459 smoke aerosol SSA values are used in this study, as values for central Africa have a strong seasonal
460 dependency (e.g. Eck et al., 2013). While SSA values of 0.85 and 0.86 are used for the 354 and
461 388 nm channels, respectively, in our study, we have also examined the sensitivity of simulated
462 OMI AI with respect to differing SSA values (Figure 6). Figures 6a-c show the simulated AI at
463 1200 UTC 28 July 2007 using NAAPS reanalysis data (Lynch et al, 2016) for three scenarios: SSA
464 values at 354 and 388 nm of 0.84 and 0.84 (Figure 6a), 0.85 and 0.85 (Figure 6b) and 0.86 and
465 0.86 (Figure 6c). Over the central Africa area, where smoke plumes are expected, simulated OMI
466 AI patterns are similar for Figures 6a and 6b, but reduced values in AI are found when using higher
467 SSA values of 0.86 at both 354 and 388 nm. This is further confirmed by the averaged AI for the
468 smoke region over central Africa (-0.5° to -15.5° latitude and 10.5° to 30.5° E longitude; indicated
469 using the black box in Figure 6f) of 0.96, 0.94 and 0.78 for Figures 6a, 6b and 6c respectively.

470 Figures 6d-f show the sensitivity for adjustments of the SSA values at 388nm while
471 maintaining a fixed SSA value of 0.85 at 354 nm. Here the SSA values at 388 nm are set to 0.85,
472 0.855 and 0.86 for Figures 6d, 6e and 6f respectively. Interestingly, the spectral dependence of
473 SSA seems to affect the simulated AI significantly, and this phenomenon has also been reported
474 by previous studies (e.g. Hammer et al., 2017). The averaged AI values over central Africa (again,
475 indicated by the black box in Figure 6f) are 0.94, 1.11 and 1.32 for 388 nm SSAs of 0.85, 0.855
476 and 0.86, respectively. This exercise suggests that simulated AI is a strong function of SSA, so
477 that both the spectral dependence of SSA values at 354 and 388 nm and reliable SSA values are
478 needed on a regional basis for future applications.



479 Interestingly, although simulated AI values are significantly affected by perturbing SSA
480 values as shown in Figure 6, less significant impacts are observed for NAAPS AOD. This is found
481 by running the OMI AI DA for 1200UTC, July 28, 2015 for SSA values used in generating Figure
482 6. For example, for the black box highlighted region in Figure 6f, the averaged values for the
483 simulated OMI AI are 0.96, 0.94 and 0.78 for using SSA values at 354 / 388 nm channels of 0.84
484 / 0.84, 0.85 / 0.85 and 0.86 / 0.86, respectively. The corresponding NAAPS AODs are found to
485 be 0.559, 0.560 and 0.585 after OMI AI DA, which is a change of less than 5%. Similar, by fixing
486 the SSA value of the 354 nm channel as 0.85 and perturbing SSA values at 388 nm from 0.85 to
487 0.86, a ~30% change is found in simulated OMI AI (from 0.94 to 1.32), yet a ~10% change is
488 found for the NAAPS AOD (from 0.560 to 0.504) after OMI AI DA.

489 It is also of interest to investigate the changes in aerosol vertical distributions due to the
490 OMI AI DA. For this exercise, we selected the 1200 UTC 28 July 2007 case and compared vertical
491 distributions of smoke and dust aerosols near the peak AI value of the smoke plume (9.5°S and
492 20.5°E) for the NAAPS natural and AI DA runs (Figure 7a). As shown in Figure 7a, the
493 corrections to dust and smoke aerosol concentrations from the AI DA system seem to be systematic
494 changes across the majority of vertical layers, instead of moving dust or smoke aerosol plumes
495 vertically. As dust aerosol concentrations are reduced at all layers and a systematic correction to
496 smoke aerosol concentrations, although non-linear, is also observed. AI assimilation helps reduce
497 the amount of upper troposphere dust (likely to be artifact) but does change the layer centroid
498 slightly upwards. We have also evaluated NAAPS vertical distributions near a peak dust plume
499 region (25.5°N and 12.5°W) for the 12Z 28 July 2007 case as shown in Figure 7b. Similar to
500 Figure 7a, a non-linear correction to dust aerosol concentrations is also observed across the vertical
501 domain.



502

503 **4.4 Issues and discussions**

504 The OMI AI data assimilation system is a proxy for all-sky, all-band modeling system
505 radiance assimilation. It contains all the necessary components for such radiance assimilation,
506 including a forward model for simulating radiances and AI values and their Jacobians, based on a
507 full vector linearized radiative transfer model called for every observation. Therefore, the
508 computational burden is a direct issue associated with the deployment of calls to a radiative transfer
509 model for each observation. For the study area in this work, after binning OMI AI data into a
510 $1^\circ \times 1^\circ$ (Latitude/Longitude) product, it still takes about ~ 1 CPU day for NAAPS to run for one
511 month of model time. Clearly, there will be an unavoidable computational burden of some sort
512 for OMI AI assimilation and by extension, for future radiance assimilation in the UV/visible
513 spectrum for aerosol analyses. Performance enhancement methods, such as parallel processing
514 (the VLIDORT software is thread-safe and can be used in parallel environments such as OpenMP),
515 or fast look-up-table extraction based on neural-networks and trained data sets of forward
516 simulation, must be explored in order to enable such assimilation applications in near real time on
517 a global scale.

518 In contrast with the assimilation of retrieved aerosol properties, both aerosol absorption
519 and scattering need to be accounted for when assimilating radiance or OMI AI in the UV spectrum.
520 This requires the inclusion of more dynamic aerosol optical properties into the data assimilation
521 process, and properties that vary with region and season. As noted already, even for biomass
522 burning aerosols over South Africa, lower single scattering albedo values were found at earlier
523 stages of burning seasons (e.g. Eck et al., 2013). A look-up-table of aerosol optical properties as



524 functions of region and season will be needed for global implications of OMI AI as well as future
525 radiance assimilation for aerosol modeling.

526 OMI AI is sensitive to above-cloud UV-absorbing aerosols (e.g. Yu et al., 2012; Alfaro-
527 Contreras et al., 2014), and therefore, OMI AI values over cloudy scenes were also used in this
528 study. However, OMI AI cannot be used to infer aerosol properties for aerosol plumes beneath a
529 cloud deck. For regions with high clouds, the use of OMI AI data assimilation will likely result in
530 an underestimation of AOD as below-cloud aerosol plumes are not accounted for. Therefore, only
531 OMI AI data over low cloud scenes are to be used for aerosol assimilation efforts. In addition,
532 although some quality assurance steps were applied in this study for the OMI AI data, lower AI
533 values were observed over glint regions near the west coast of Africa. Abnormally high OMI AI
534 values are also seen near the Arctic region - this may be related to the presence of floating ice
535 sheets. Thus, innovative and detailed data screening and quality assurance steps are needed to
536 exclude potentially noisy OMI AI retrievals and for further application of OMI AI data
537 assimilation on a global scale.

538 Even with these known issues, OMI AI assimilation as presented in the study illustrates a
539 new method for assimilating non-conventional aerosol products. Bearing in mind that OMI AI
540 assimilation is essentially radiance assimilation in the UV spectrum, this study demonstrates the
541 potential of directly assimilating satellite radiance in the UV/visible spectrum for aerosol modeling
542 and analyses.

543

544 **5.0 Conclusions**

545 The OMI aerosol index (AI), which measures the differences between simulated radiances
546 over Rayleigh sky and observed radiances at 354 nm, has been used to detect the presence of
547 absorbing aerosols over both dark and bright surfaces. We have constructed a new assimilation



548 system, based on the VLIDORT radiative transfer code as the major component of the forward
549 model, for the direct assimilation of OMI AI. The aim is to improve accuracies of aerosol analyses
550 over bright surfaces such as cloudy regions and deserts.

551 The performance of the OMI AI data assimilation system was evaluated over South-Central
552 and Northern Africa regions for the period of 01 July -31 August 2007. This evaluation was done
553 through inter-comparing NAAPS analyses with and without the inclusion of OMI AI data
554 assimilation. Besides cloud-free AI retrievals over dark surfaces, OMI AI retrievals over desert
555 regions and over areas were also considered. When compared against AERONET data, a total of
556 ~28% reduction in Root-Mean-Square-Error (RMSE) with a ~32% reduction in absolute error
557 were found for NAAPS analyses with the use of OMI AI assimilation. Also, NAAPS analyses
558 with the inclusion of OMI AI data assimilation show similar aerosol patterns to those in the OMI
559 AI data sets, showing that our OMI AI data assimilation system works as expected.

560 This study also suggests that NAAPS analyses with OMI AI data assimilation cannot out-
561 perform NAAPS reanalyses data that were incorporated with MODIS and MISR AOD
562 assimilation, and validated against AERONET data. This is not surprising, as OMI AI is only a
563 proxy for the AOD and is sensitive to other factors such as surface albedo and aerosol vertical
564 distribution. Also, AERONET data are only available over cloud-free field of views, so the
565 performance of our OMI AI data assimilation system over cloudy regions has not been evaluated.

566 There are a number of issues arising from our study. For example, aerosol optical
567 properties are needed for the OMI AI-DA system - these have strong regional and temporal
568 signatures that need to be carefully quantified before applying them to the AI-DA on a global scale.
569 Also, OMI AI retrievals are rather noisy and contain known and unknown biases. Abnormally
570 high OMI AI values are found over mountain regions as well the polar regions. Sporadic high AI



571 values are also known to occur, for reasons that are still not properly understood. Even though
572 quality assurance steps were proposed in this study, detailed analysis of OMI AI data are needed
573 for future implementation of OMI AI data assimilation for aerosol studies.

574 Lastly, AI values are derived from radiances and thus, the AI-DA system presented in the
575 study can be thought of as a radiance assimilation system for the UV spectrum. This is because
576 the AI-DA system contains all necessary components for radiance assimilation, based on a forward
577 model for calculating not only simulated satellite radiances, but also the aerosol-profile Jacobians
578 of these radiance, both quantities as functions of observation conditions. This study is among the
579 first attempts at radiance assimilation at the UV spectrum and indicates the future potential for
580 direct radiance assimilation at the UV and visible spectra for aerosol analyses and forecasts.

581

582 **Author contributions.** All authors contributed to the overall design of the study. Authors JZ and
583 RS coded the system. Author JSR provided valuable suggestions though the study. Author PX
584 assist with the evaluation of the system.

585

586 **Code and data availability:** The VLIDORT radiative transfer model is available to the public
587 through contacting RT solutions Inc. (http://www.rtslidort.com/mainprod_vlidort.html). The
588 NAAPS model belongs to the Naval Research Laboratory and is not publically available. The
589 NAAPS reanalysis data are available from the USGODAE web site
590 (https://nrlgodae1.nrlmry.navy.mil/cgi-bin/datalist.pl?dset=nrl_naaps_reanalysis&summary=Go).

591 The OMI OMAERUV data are available from the NASA's Goddard Earth Sciences Data and
592 Information Services Center (GES DISC;



593 https://disc.gsfc.nasa.gov/datasets/OMAERUV_003/summary). AERONET data are obtained
594 from the NASA AERONET webpage (<https://aeronet.gsfc.nasa.gov/>).

595

596 **Competing interests.** The authors claim no competing interests.

597

598 **Acknowledgements.** We thank the NASA AERONET group for the sun-photometer data used in
599 the study. “We acknowledge the use of imagery from the NASA Worldview application
600 (<https://worldview.earthdata.nasa.gov>), part of the NASA Earth Observing System Data and
601 Information System (EOSDIS).”

602

603 **Financial support.** This study is supported by the NASA grant NNX17AG52G. Co-Author JSR
604 was supported by the Office of Naval Research Code 322.

605

606

607



608 **References:**

- 609 Alfaro-Contreras, R., Zhang, J., Campbell, J. R., and Reid, J. S.: Investigating the frequency and
610 trends in global above-cloud aerosol characteristics with CALIOP and OMI, *Atmos. Chem.*
611 *Phys.*, 16, 47-69, doi:10.5194/acp-16-47-2016, 2016.
- 612 Alfaro-Contreras R., Zhang, J., Campbell, J. R., Holz, R. E., and Reid, J. S.: Evaluating the Impact
613 of Aerosol Particles above Cloud on Cloud Optical Depth Retrievals from MODIS, *J.*
614 *Geophys. Res. Atmos.*, 119, 5410–5423, doi:10.1002/2013JD021270, 2014.
- 615 Benedetti, A., et al., Aerosol analysis and forecast in the European Centre for Medium-Range
616 Weather Forecasts Integrated Forecast System: 2. Data assimilation, *J. Geophys. Res.*, 114,
617 D13205, doi:10.1029/2008JD011115, 2009.
- 618 Bessho, K., Date, K., Hayashi, M., Ikeda, A., Imai, T., Inoue, H., Kumagai, Y., Miyakawa, T.,
619 Murata, H., Ohno, T., Okuyama, A., Oyama, R., Sasaki, Y., Shimazu, Y., Shimoji, K.,
620 Sumida, Y., Suzuki, M., Taniguchi, H., Tsuchiyama, H., Uesawa, D., Yokota, H., &
621 Yoshida, R.: An Introduction to Himawari-8/9- Japan's New-Generation Geostationary
622 Meteorological Satellites. *Journal of the Meteorological Society of Japan. Ser. II*, 94(2),
623 151–183. <https://doi.org/10.2151/jmsj.2016-009>, 2016.
- 624 Buchard, V., Randles, C., Silva, A., Darmenov, A., Colarco, P., Govindaraju, R., Ferrare, R., Hair,
625 J., Beyersdorf, A., Ziemba, L., Yu, H.: The MERRA-2 Aerosol Reanalysis, 1980 Onward.
626 Part II: Evaluation and Case Studies *Journal of Climate* [https://dx.doi.org/10.1175/jcli-d-](https://dx.doi.org/10.1175/jcli-d-16-0613.1)
627 [16-0613.1](https://dx.doi.org/10.1175/jcli-d-16-0613.1), 2017.
- 628 Buchard, V., da Silva, A. M., Colarco, P. R., Darmenov, A., Randles, C. A., Govindaraju, R.,
629 Torres, O., Campbell, J., and Spurr, R.: Using the OMI aerosol index and absorption



630 aerosol optical depth to evaluate the NASA MERRA Aerosol Reanalysis, *Atmos. Chem.*
631 *Phys.*, 15, 5743–5760, <https://doi.org/10.5194/acp-15-5743-2015>, 2015.

632 Cochrane, S. P., Schmidt, K. S., Chen, H., Pilewskie, P., Kittelman, S., Redemann, J., LeBlanc,
633 S., Pistone, K., Kacenelenbogen, M., Segal Rozenhaimer, M., Shinozuka, Y., Flynn, C.,
634 Platnick, S., Meyer, K., Ferrare, R., Burton, S., Hostetler, C., Howell, S., Freitag, S.,
635 Dobracki, A., and Doherty, S.: Above-cloud aerosol radiative effects based on ORACLES
636 2016 and ORACLES 2017 aircraft experiments, *Atmos. Meas. Tech.*, 12, 6505–6528,
637 <https://doi.org/10.5194/amt-12-6505-2019>, 2019.

638 Colarco, P. R., Nowottnick, E. P., Randles, C. A., Yi, B., Yang, P., Kim, K.-M., Smith, J. A. and
639 Bardeen, C. G.: Impact of radiatively interactive dust aerosols in the NASA GEOS-5
640 climate model: sensitivity to dust particle shape and refractive index. *Journal of*
641 *Geophysical Research: Atmospheres*, 119(2), 753– 786.
642 <https://doi.org/10.1002/2013JD020046>, 2014.

643 Dai, A.: Precipitation characteristics in eighteen coupled climate models. *Journal of Climate*, 19,
644 4605– 4630, 2006.

645 Eck, T. F., Holben, B. N., Reid, J. S., Mukelabai, M. M., Piketh, S. J., Torres, O., Jethva, H. T.,
646 Hyer, E. J., Ward, D. E., Dubovik, O., and Sinyuk, A.: A seasonal trend of single scattering
647 albedo in southern African biomass-burning particles: Implications for satellite products
648 and estimates of emissions for the world’s largest biomass-burning source, *J. Geophys.*
649 *Res.-Atmos.*, 118, 6414–6432, 2013.

650 Giles, D. M., Sinyuk, A., Sorokin, M. G., Schafer, J. S., Smirnov, A., Slutsker, I., Eck, T. F.,
651 Holben, B. N., Lewis, J. R., Campbell, J. R., Welton, E. J., Korkin, S. V., and Lyapustin,
652 A. I.: Advancements in the Aerosol Robotic Network (AERONET) Version 3 database –



653 automated near-real-time quality control algorithm with improved cloud screening for Sun
654 photometer aerosol optical depth (AOD) measurements, *Atmos. Meas. Tech.*, 12, 169–209,
655 <https://doi.org/10.5194/amt-12-169-2019>, 2019.

656 Hammer, M. S., Martin, R. V., van Donkelaar, A., Buchard, V., Torres, O., Ridley, D. A., and
657 Spurr, R. J. D.: Interpreting the ultraviolet aerosol index observed with the OMI satellite
658 instrument to understand absorption by organic aerosols: implications for atmospheric
659 oxidation and direct radiative effects, *Atmos. Chem. Phys.*, 16, 2507–2523,
660 <https://doi.org/10.5194/acp-16-2507-2016>, 2016.

661 Herman, J. R., Bhartia, P. K., Torres, O., Hsu, C., Seftor, C., and Celarier, E.: Global distribution
662 of UV-absorbing aerosols from Nimbus 7/TOMS data, *J. Geophys. Res.*, 102(D14),
663 16911–16922, doi:[10.1029/96JD03680](https://doi.org/10.1029/96JD03680), 1997.

664 Holben, B. N., and coauthors: AERONET—A Federated Instrument Network and Data Archive
665 for Aerosol Characterization. *Remote Sensing of Environment*, 66(1), 1–16.
666 [https://doi.org/10.1016/S0034-4257\(98\)00031-5](https://doi.org/10.1016/S0034-4257(98)00031-5), 1998.

667 Hogan, T.F., Liu, M., Ridout, J. A., Peng, M. S., Whitcomb, T. R., Ruston, B. C., Reynolds, C. A.,
668 Eckermann, S. D., Moskaitis, J. R., Baker, N. L., McCormack, J. P., Viner, K. C., McLay,
669 J. G., Flatau, M. K., Xu, L., Chen, C., and Chang, S. W.: The Navy Global Environmental
670 Model, *Oceanography*, 27(3):116–125, <https://doi.org/10.5670/oceanog.2014.73>, 2014.

671 Hsu, N. C., Lee, J., Sayer, A. M., Kim, W., Bettenhausen, C., and Tsay, S.-C.: VIIRS Deep Blue
672 aerosol products over land: Extending the EOS long-term aerosol data records, *J. Geophys.*
673 *Res.-Atmos.*, 124, 4026–4053, <https://doi.org/10.1029/2018JD029688>, 2019.



- 674 Hsu, N. C., Jeong, M.-J., Bettenhausen, C., Sayer, A. M., Hansell, R., Seftor, C. S., Huang, J., and
675 Tsay S.-C.: Enhanced Deep Blue aerosol retrieval algorithm: The second generation, *J.*
676 *Geophys. Res. Atmos.*, 118, doi:10.1002/jgrd.50712, 2013.
- 677 Hsu, N. C., Gautam, R., Sayer, A. M., Bettenhausen, C., Li, C., Jeong, M. J., Tsay, S.-C., and
678 Holben, B. N.: Global and regional trends of aerosol optical depth over land and ocean
679 using SeaWiFS measurements from 1997 to 2010, *Atmos. Chem. Phys.*, 12, 8037–8053,
680 <https://doi.org/10.5194/acp-12-8037-2012>, 2012.
- 681 Hyer, E. J., Reid, J. S., and Zhang, J.: An over-land aerosol optical depth data set for data
682 assimilation by filtering, correction, and aggregation of MODIS Collection 5 optical depth
683 retrievals, *Atmos. Meas. Tech.*, 4, 379–408, doi:10.5194/amt-4-379-2011, 2011.
- 684 Joyce, J. R., Janowiak, E. J., Arkin, P. A., and Xie, P.: CMORPH: A method that produces global
685 precipitation estimates from passive microwave and infrared data at high spatial and
686 temporal resolution, *J. Hydrometeor.*, 5, 487–503, 2004.
- 687 Kahn, R. A., Gaitley, B. J., Garay, M. J., Diner, D. J., Eck, T. F., Smirnov, A. and Holben B. N.:
688 Multiangle Imaging SpectroRadiometer global aerosol product assessment by comparison
689 with the Aerosol Robotic Network, *J. Geophys. Res.*, 115, *D23209*,
690 doi:[10.1029/2010JD014601](https://doi.org/10.1029/2010JD014601), 2010.
- 691 Lee, J., Hsu, N.C., Bettenhausen, C., Sayer, A. M., Seftor, C. J., and Jeong, M.-J.: Retrieving the
692 height of smoke and dust aerosols by synergistic use of VIIRS, OMPS, and CALIOP
693 observations, *J. Geophys. Res. Atmos.*, 120, 8372–8388, doi:10.1002/2015JD023567, 2015.
- 694 Levelt, P. F., Joiner, J., Tamminen, J., Veefkind, J. P., Bhartia, P. K., Stein Zweers, D. C., Duncan,
695 B. N., Streets, D. G., Eskes, H., van der A, R., McLinden, C., Fioletov, V., Carn, S., de
696 Laat, J., DeLand, M., Marchenko, S., McPeters, R., Ziemke, J., Fu, D., Liu, X., Pickering,



- 697 K., Apituley, A., González Abad, G., Arola, A., Boersma, F., Chan Miller, C., Chance, K.,
698 de Graaf, M., Hakkarainen, J., Hassinen, S., Ialongo, I., Kleipool, Q., Krotkov, N., Li, C.,
699 Lamsal, L., Newman, P., Nowlan, C., Suleiman, R., Tilstra, L. G., Torres, O., Wang, H.,
700 and Wargan, K.: The Ozone Monitoring Instrument: overview of 14 years in space, *Atmos.*
701 *Chem. Phys.*, 18, 5699–5745, <https://doi.org/10.5194/acp-18-5699-2018>, 2018.
- 702 Levy, R. C., Mattoo, S., Munchak, L. A., Remer, L. A., Sayer, A. M., Patadia, F., and Hsu N. C.:
703 The Collection 6 MODIS aerosol products over land and ocean, *Atmos. Meas. Tech.*, 6,
704 2989–3034, doi:10.5194/amt-6-2989-2013, 2013.
- 705 Li, Z., Zang, Z., Li, Q. B., Chao, Y., Chen, D., Ye, Z., Liu, Y., and Liou, K. N.: A three-dimensional
706 variational data assimilation system for multiple aerosol species with WRF/Chem and an
707 application to PM_{2.5} prediction, *Atmos. Chem. Phys.*, 13, 4265–4278,
708 <https://doi.org/10.5194/acp-13-4265-2013>, 2013.
- 709 Lynch, P., Reid, J. S., Westphal, D. L., Zhang, J., Hogan, T. F., Hyer, E. J., Curtis, C. A., Hegg,
710 D. A., Shi, Y., Campbell, J. R., Rubin, J. I., Sessions, W. R., Turk, F. J., and Walker, A.
711 L.: An 11-year global gridded aerosol optical thickness reanalysis (v1.0) for atmospheric
712 and climate sciences, *Geosci. Model Dev.*, 9, 1489–1522, [https://doi.org/10.5194/gmd-9-](https://doi.org/10.5194/gmd-9-1489-2016)
713 1489-2016, 2016.
- 714 Pistone, K., Redemann, J., Doherty, S., Zuidema, P., Burton, S., Cairns, B., Cochrane, S., Ferrare,
715 R., Flynn, C., Freitag, S., Howell, S. G., Kacenelenbogen, M., LeBlanc, S., Liu, X.,
716 Schmidt, K. S., Sedlacek III, A. J., Segal-Rozenhaimer, M., Shinozuka, Y., Stamnes, S.,
717 van Diedenhoven, B., Van Harten, G., and Xu, F.: Intercomparison of biomass burning
718 aerosol optical properties from in situ and remote-sensing instruments in ORACLES-2016,
719 *Atmos. Chem. Phys.*, 19, 9181–9208, <https://doi.org/10.5194/acp-19-9181-2019>, 2019.



- 720 Randles, C., Silva, A., Buchard, V., Colarco, P., Darmenov, A., Govindaraju, R., Smirnov, A.,
721 Holben, B., Ferrare, R., Hair, J., Shinozuka, Y., Flynn, C.: The MERRA-2 Aerosol
722 Reanalysis, 1980 Onward. Part I: System Description and Data Assimilation Evaluation,
723 Journal of Climate <https://dx.doi.org/10.1175/jcli-d-16-0609.1>, 2017.
- 724 Reid, J. S., Hyer, E. J., Prins, E. M., Westphal, D. L., Zhang, J., Wang, J., Christopher, S. A.,
725 Curtis, C. A., Schmidt, C. C., Eleuterio, D. P., Richardson, K. A., and Hoffman, J. P.:
726 Global monitoring and forecasting of biomass-burning smoke: Description and lessons
727 from the Fire Locating and Modeling of Burning Emissions (FLAMBE) program, IEEE J.
728 Sel. Top. Appl., 2, 144–162, 2009.
- 729 Rubin J. I., Reid, J. S., Hansen, J. A., Anderson, J. L., Holben, B. N., Lynch, P., Westphal, D. L.,
730 and Zhang, J.: Assimilation of AERONET and MODIS AOT observations using
731 Variational and Ensemble Data Assimilation Methods and Its Impact on Aerosol
732 Forecasting Skill, J. Geophys. Res., DOI: 10.1002/2016JD026067, 2017.
- 733 Saide, P. E., Carmichael, G. R., Liu, Z., Schwartz, C. S., Lin, H. C., da Silva, A. M., and Hyer, E.:
734 Aerosol optical depth assimilation for a size-resolved sectional model: impacts of
735 observationally constrained, multi-wavelength and fine mode retrievals on regional scale
736 analyses and forecasts, Atmos. Chem. Phys., 13, 10425–10444,
737 <https://doi.org/10.5194/acp-13-10425-2013>, 2013.
- 738 Schutgens, N. A. J., Miyoshi, T., Takemura, T., and Nakajima, T.: Applying an ensemble Kalman
739 filter to the assimilation of AERONET observations in a global aerosol transport model,
740 Atmos. Chem. Phys., 10, 2561–2576, <https://doi.org/10.5194/acp-10-2561-2010>, 2010.



- 741 Schwartz, C. S., Liu, Z., Lin, H.-C., and McKeen, S. A.: Simultaneous three-dimensional
742 variational assimilation of surface fine particulate matter and MODIS aerosol optical depth,
743 J. Geophys. Res., 117, D13202, doi:10.1029/2011JD017383, 2012.
- 744 Sekiyama, T. T., Tanaka, T. Y., Shimizu, A., and Miyoshi, T.: Data assimilation of CALIPSO
745 aerosol observations, Atmos. Chem. Phys., 10, 39–49, [https://doi.org/10.5194/acp-10-39-](https://doi.org/10.5194/acp-10-39-2010)
746 2010, 2010.
- 747 Sessions, W. R., Reid, J. S., Benedetti, A., Colarco, P. R., da Silva, A., Lu, S., Sekiyama, T.,
748 Tanaka, T. Y., Baldasano, J. M., Basart, S., Brooks, M. E., Eck, T. F., Iredell, M., Hansen,
749 J. A., Jorba, O. C., Juang, H.-M. H., Lynch, P., Morcrette, J.-J., Moorthi, S., Mulcahy, J.,
750 Pradhan, Y., Razinger, M., Sampson, C. B., Wang, J., and Westphal, D. L.: Development
751 towards a global operational aerosol consensus: basic climatological characteristics of the
752 International Cooperative for Aerosol Prediction Multi-Model Ensemble (ICAP-MME),
753 Atmos. Chem. Phys., 15, 335–362, <https://doi.org/10.5194/acp-15-335-2015>, 2015.
- 754 Shi Y., Zhang J., J.S. Reid, B. Liu, E.J. Hyer, Critical evaluation of cloud contamination in the
755 MISR aerosol products using MODIS cloud masking products, Atmos. Meas. Tech., 7,
756 1791-1801, doi:10.5194/amt-7-1791-2014, 2014.
- 757 Spurr, R. J. D., and Christi. M.: The LIDORT and VLIDORT Linearized Scalar and Vector
758 Discrete Ordinate Radiative Transfer Models: An Update for the last 10 Years, Light
759 Scattering Reviews, Volume 12, ed. A. Kokhanovsky, Springer, 2019.
- 760 Spurr, R. J. D.: VLIDORT: A linearized pseudo-spherical vector discrete ordinate radiative
761 transfer code for forward model and retrieval studies in multilayer multiple scattering
762 media, J. Quant. Spectrosc. Radiat. Transfer, 102(2), 316-342,
763 doi:10.1016/j.jqsrt.2006.05.005, 2006.



- 764 Torres, O., Bhartia, P. K., Jethva, H., and Ahn, C.: Impact of the ozone monitoring instrument row
765 anomaly on the long-term record of aerosol products, *Atmos. Meas. Tech.*, 11, 2701–2715,
766 <https://doi.org/10.5194/amt-11-2701-2018>, 2018.
- 767 Torres, O., Ahn, C., and Chen, Z.: Improvements to the OMI near-UV aerosol algorithm using A-
768 train CALIOP and AIRS observations, *Atmos. Meas. Tech.*, 6, 3257–3270,
769 <https://doi.org/10.5194/amt-6-3257-2013>, 2013.
- 770 Torres, O., Jethva, H., and Bhartia, P. K.: Retrieval of Aerosol Optical Depth above Clouds from
771 OMI Observations: Sensitivity Analysis and Case Studies. *J. Atmos. Sci.*, 69, 1037–1053,
772 <https://doi.org/10.1175/JAS-D-11-0130.1>, 2012.
- 773 Torres, O., Tanskanen, A., Veihelmann, B., Ahn, C., Braak, R., Bhartia, P. K., Veefkind, P., Levelt,
774 P.: Aerosols and surface UV products from Ozone Monitoring Instrument observations:
775 An overview. *Journal of Geophysical Research*, 112, D24S47.
776 <https://doi.org/10.1029/2007JD008809>, 2007.
- 777 Uno, I., Yumimoto, K., Shimizu, A., Hara, Y., Sugimoto, N., Wang, Z., Liu, Z., and Winker D.
778 M.: 3D structure of Asian dust transport revealed by CALIPSO lidar and a 4DVAR dust
779 model, *Geophys. Res. Lett.*, 35, L06803, doi:10.1029/2007GL032329, 2008.
- 780 Westphal, D. L., Toon, O. B., and Carlson, T. N.: A case study of mobilization and transport of
781 Saharan dust. *J. Atmos. Sci.*, 45, 2145–2175, 1988.
- 782 Witek M. L., Flatau, P. J., Quinn, P. K., Westphal, D. L., Global sea-salt modeling: results and
783 validation against multicampaign shipboard measurements, *J. Geophys. Res.*, 112, p.
784 D08215, [10.1029/2006JD007779](https://doi.org/10.1029/2006JD007779), 2007.
- 785 Xian, P., Reid, J.S., Turk, J.F., Hyer, E.J. and Westphal, D.L.: Impact of modeled versus satellite
786 measured tropical precipitation on regional smoke optical thickness in an aerosol transport



787 model, Geophysical Research Letters, 36, L16805,
788 <https://doi.org/10.1029/2009GL038823>, 2009.

789 Yu, H., Zhang, Y., Chin M., Liu Z., Omar A., Remer L. A. Yang Y., Yuan T. and Zhang J.: An
790 Integrated Analysis of Aerosol above Clouds from A-Train Multi-sensor Measurements,
791 Remote Sens. Environ., 121, 125–131, <https://doi.org/10.1016/j.rse.2012.01.011>, 2012.

792 Yumimoto, K., Uno, I., Sugimoto, N., Shimizu, A., Liu, Z., and Winker, D. M.: Adjoint inversion
793 modeling of Asian dust emission using lidar observations, Atmos. Chem. Phys., 8, 2869–
794 2884, <https://doi.org/10.5194/acp-8-2869-2008>, 2008.

795 Zhang J., Reid, J. S., Campbell, J. R., Hyer, E. J., and Westphal, D. L.: Evaluating the Impact of
796 Multi-Sensor Data Assimilation on A Global Aerosol Particle Transport Model. J.
797 Geophys. Res. Atmos., 119, 4674–4689, doi:[10.1002/2013JD020975](https://doi.org/10.1002/2013JD020975), 2014.

798 Zhang, J., Campbell, J. R., Reid, J. S., Westphal, D. L., Baker, N. L., Campbell, W. F., and Hyer,
799 E. J.: Evaluating the impact of assimilating CALIOP-derived aerosol extinction profiles on
800 a global mass transport model, Geophys. Res. Lett., Vo. 38, No. 14, L14801, doi:
801 [10.1029/2011GL04773](https://doi.org/10.1029/2011GL04773), 2011.

802 Zhang, J. and Reid, J. S.: A decadal regional and global trend analysis of the aerosol optical depth
803 using a data-assimilation grade over-water MODIS and Level 2 MISR aerosol products,
804 Atmos. Chem. Phys. Discuss., 10, 18879-18917, doi:[10.5194/acpd-10-18879-2010](https://doi.org/10.5194/acpd-10-18879-2010), 2010.

805 Zhang, J., Reid, J. S., Westphal, D. L., Baker, N. L., and Hyer, E. J.: A system for operational
806 aerosol optical depth data assimilation over global oceans, J. Geophys. Res., 113, No. D10,
807 D10208, doi: [10.1029/2007JD009065](https://doi.org/10.1029/2007JD009065), 2008.



808 Zhang, J. and Reid., J.S.: MODIS Aerosol Product Analysis for Data Assimilation: Assessment
809 of Level 2 Aerosol Optical Thickness Retrievals, J. Geophysical Research-Atmospheres,
810 VOL. 111, D22207, doi:10.1029/2005JD006898, 2006.

811

812



813 Table 1. Mass extinction cross-sections (σ , m^2/g) and single scattering albedos (ω_0) used in
814 this study.

	ABF	Dust	Smoke	Sea Salt
σ (354 nm)	7.81	0.56	6.91	0.52
ω_0 (354 nm)	1.0	0.88	0.85	1.0
σ (388 nm)	6.96	0.58	6.07	0.52
ω_0 (388 nm)	1.0	0.91	0.86	1.0

815

816



817 **Figure Captions**

818

819 **Figure 1.** (a) Spatial distribution of NAAPS AODs, using NAAPS reanalysis data from the
820 collocated OMI and NAAPS dataset for July 2007. (b). Simulated AI using NAAPS reanalysis
821 data as shown in (a). (c). Spatial distribution of OMI AI using gridded OMI data from the
822 collocated OMI and NAAPS dataset for July 2007. Grey color highlights those $1 \times 1^\circ$
823 (Latitude/Longitude) bins that have less than two collocated NAAPS and OMI AI data for the
824 study period.

825 **Figure 2.** (a). Vertical distributions of smoke, dust, anthropogenic and sea salt aerosols for the test
826 case as shown in (b). (b) Scatter plot of Jacobians of AI as a function of dust concentration: analytic
827 versus finite difference solutions.

828 **Figure 3.** (a). Aqua MODIS true-color image over Central and North Africa for July 28, 2007.
829 This composite was obtained from the NASA worldview site
830 (<https://worldview.earthdata.nasa.gov/>). (b). Spatial distribution of Gridded OMI AI for 12 UTC,
831 July 28, 2007. (c). Spatial distribution of NAAPS AOD from the NAAPS natural run for 12 UTC,
832 July 28, 2007. (d). Similar to (c) but using NAAPS AOD from the AI-DA run. (e). Simulated AI
833 using data from (c). (f). Simulated AI using data from (d).

834 **Figure 4.** (a). Spatial distribution of NAAPS AOD using NAAPS data from the AI-DA runs for
835 July and August 2007. Only NAAPS data that have collocated OMI AI data are used. (b). Spatial
836 distribution of simulated AI for July and August 2007 using NAAPS data from the AI-DA runs.
837 (c). Spatial distribution of gridded OMI AI for July and August 2007. (d). Differences between
838 Figures 4(b) and 4(c). (e-h) Similar to Figures 4(a)-4(d) but using NAAPS natural runs. Grey
839 color highlights those $1 \times 1^\circ$ (Latitude/Longitude) bins that have less than two collocated NAAPS
840 and OMI AI data for the study period.



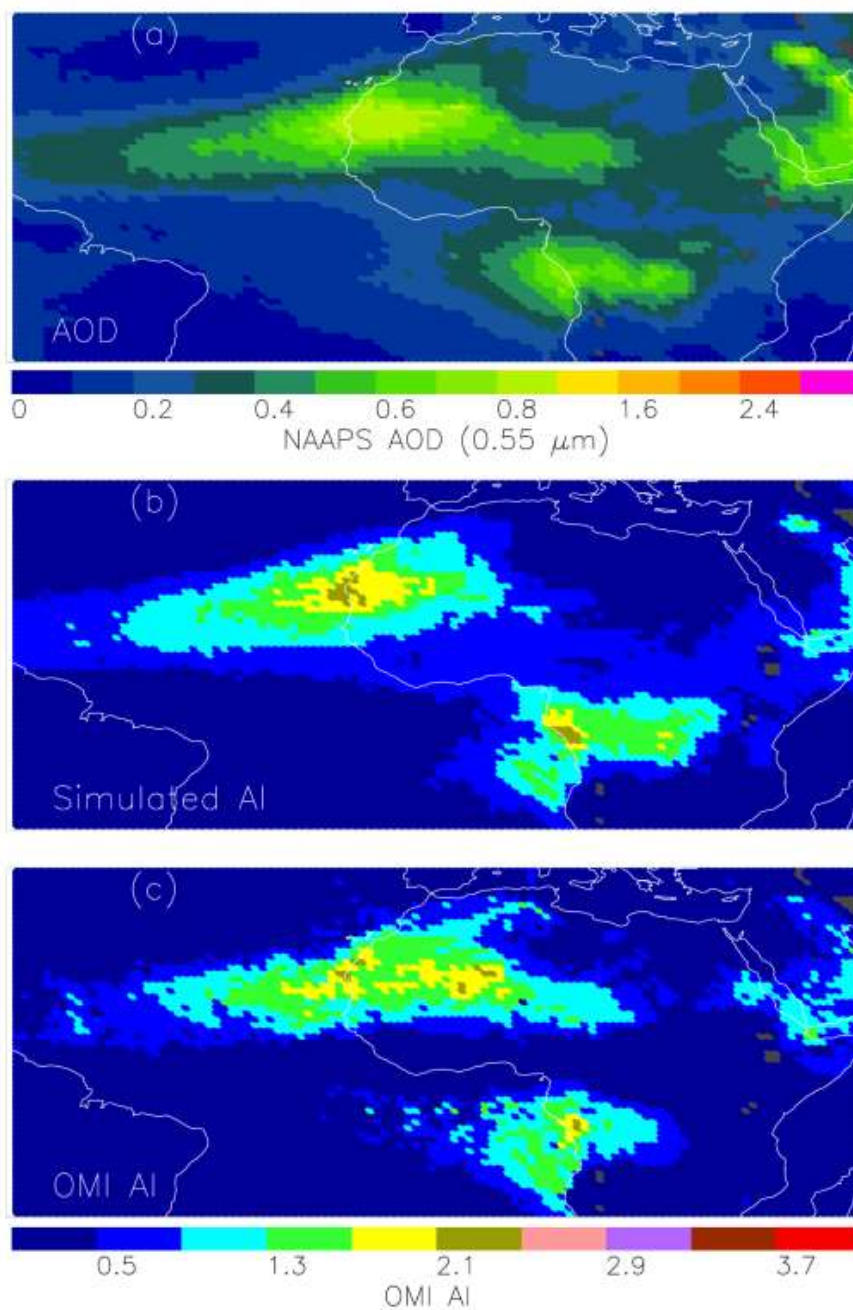
841 **Figure 5.** (a). Scatter plot of AERONET and NAAPS AOD ($0.55 \mu\text{m}$) using NAAPS data from
842 the natural runs for July-August 2007 over the study region. (b). Similar to Figure 5(a) but using
843 NAAPS data from the AI-DA runs. (c). Similar to Figure 5(a) but with AODs taken from the
844 NAAPS reanalysis.

845 **Figure 6.** Spatial distributions of simulated AI at 12 Z on July 28, 2007 using NAAPS reanalysis
846 data, with single scattering albedos of smoke aerosol at 354 and 388 nm taken to be: (a) 0.84 and
847 0.84; (b) 0.85 and 0.85; (c) 0.86 and 0.86; (d) 0.85 and 0.85; (e) 0.85, 0.855; (f) 0.85 and 0.86.

848 **Figure 7.** (a). Vertical distributions of smoke and dust aerosol concentrations over 9.5°S and
849 10.5°E at 12 Z on July 28, 2007 for both natural and AI DA runs. (b). Similar as (a) but over
850 25.5°N and 12.5°W .

851

852



853

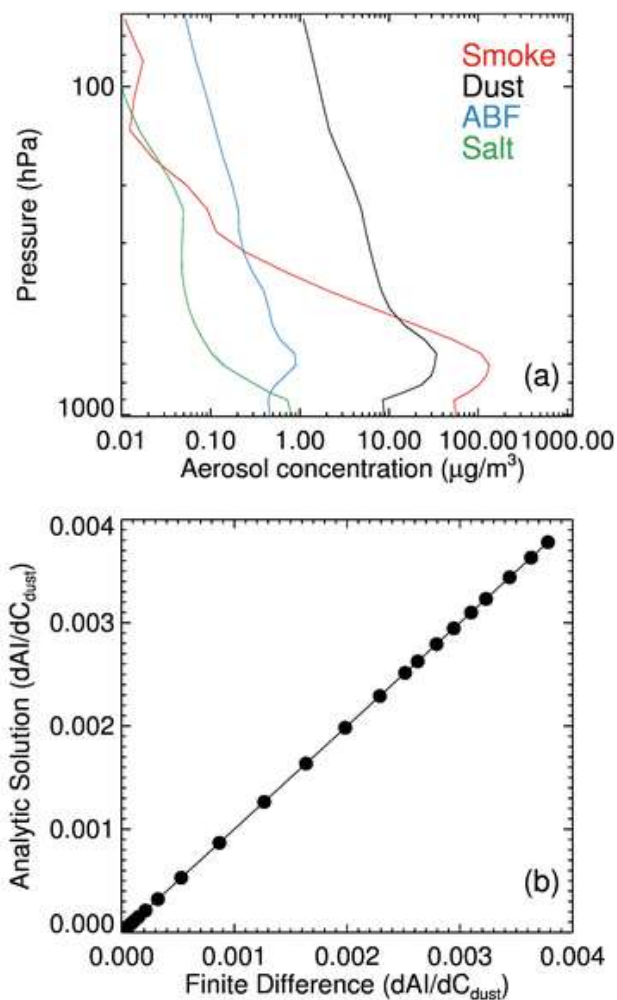
854

855

Figure 1. (a) Spatial distribution of NAAPS AODs, using NAAPS reanalysis data from the collocated OMI and NAAPS dataset for July 2007. (b). Simulated AI using NAAPS reanalysis data as shown in (a). (c). Spatial distribution of OMI AI using gridded OMI data from the collocated OMI and NAAPS dataset for July 2007. Grey color highlights those 1x1° (Latitude/Longitude) bins that have less than two collocated NAAPS and OMI AI data for the study period.



856



857

858 **Figure 2.** (a). Vertical distributions of smoke, dust, anthropogenic and sea salt aerosols for the test
859 case as shown in (b). (b) Scatter plot of Jacobians of AI as a function of dust concentration: analytic
860 versus finite difference solutions.

861

862

863

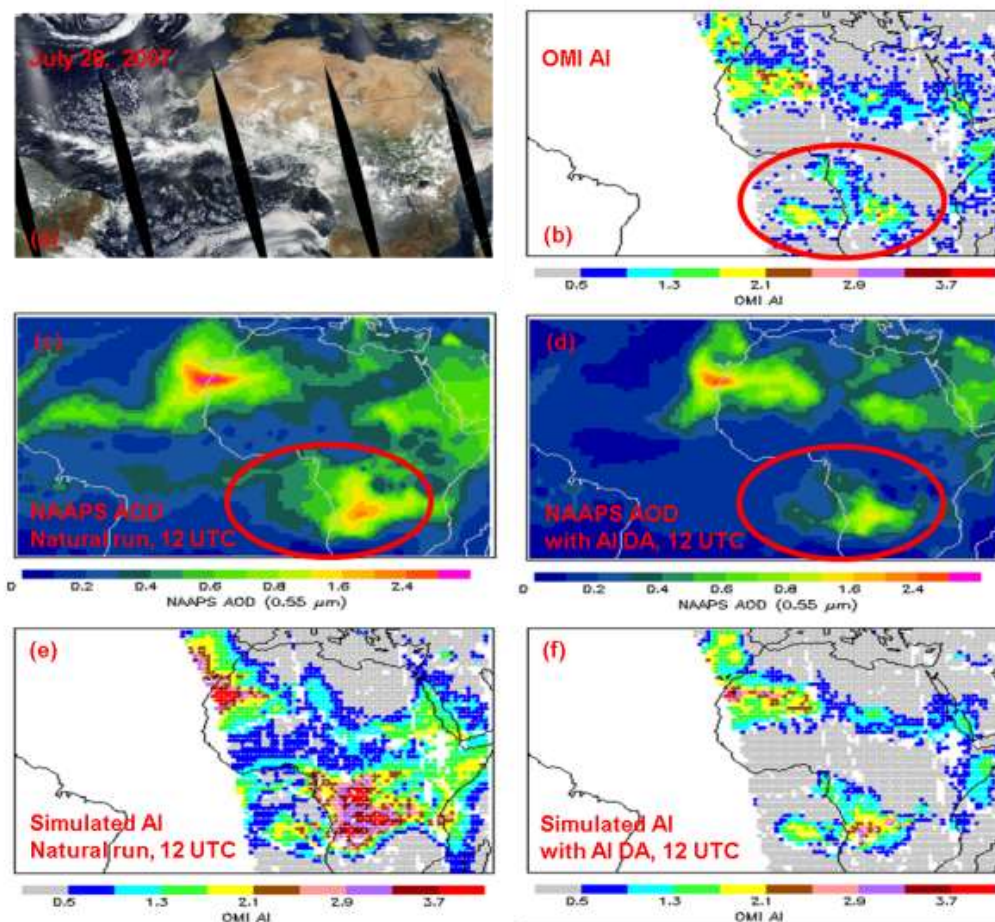
864

865



866

867



868

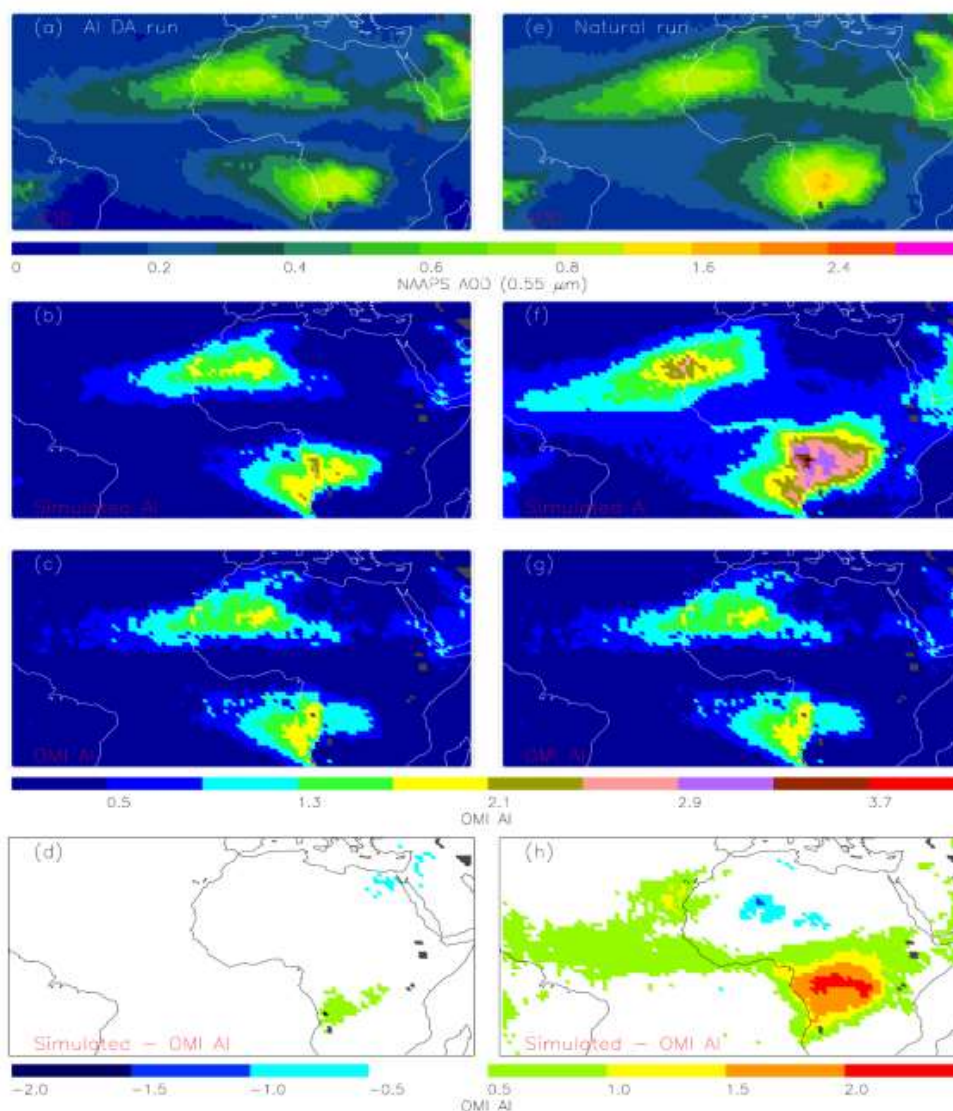
869 **Figure 3.** (a). Aqua MODIS true-color image over Central and North Africa for July 28, 2007.
870 This composite was obtained from the NASA worldview site
871 (<https://worldview.earthdata.nasa.gov/>). (b). Spatial distribution of Gridded OMI AI for 12 UTC,
872 July 28, 2007. (c). Spatial distribution of NAAPS AOD from the NAAPS natural run for 12 UTC,
873 July 28, 2007. (d). Similar to (c) but using NAAPS AOD from the AI-DA run. (e). Simulated AI
874 using data from (c). (f). Simulated AI using data from (d).

875

876

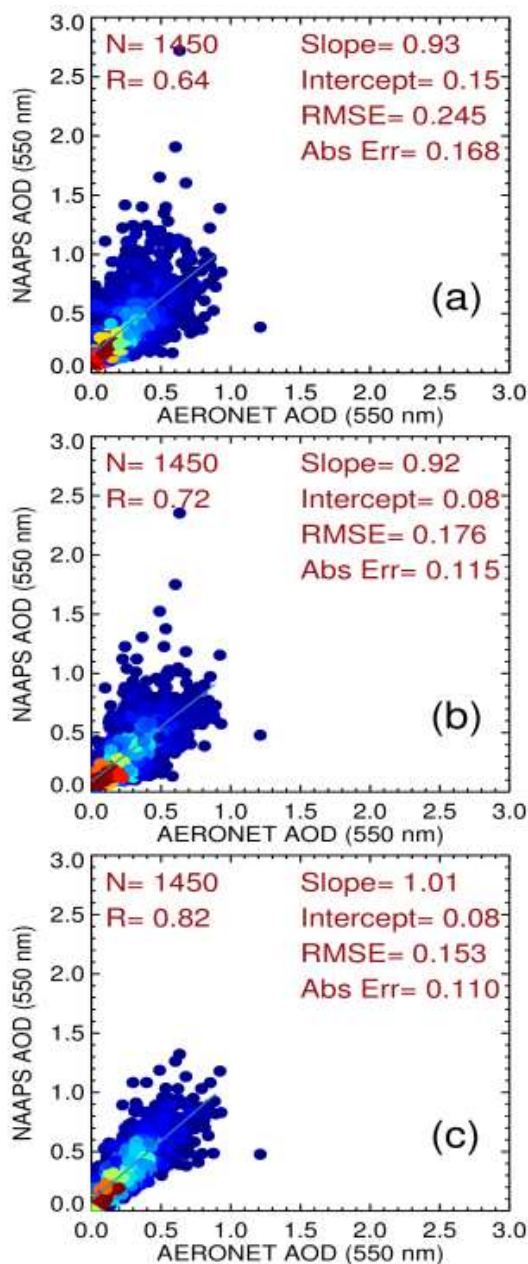
877

878



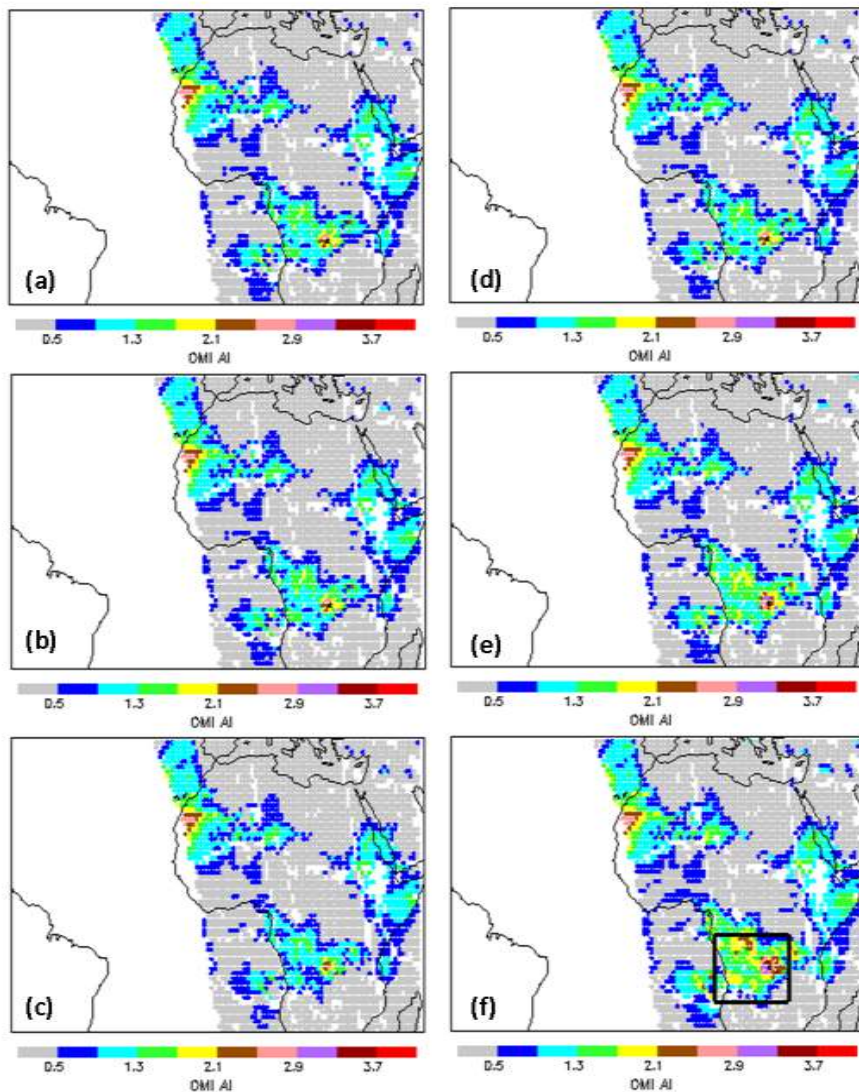
879

880 **Figure 4.** (a). Spatial distribution of NAAPS AOD using NAAPS data from the AI-DA runs for
881 July and August 2007. Only NAAPS data that have collocated OMI AI data are used. (b). Spatial
882 distribution of simulated AI for July and August 2007 using NAAPS data from the AI-DA runs.
883 (c). Spatial distribution of gridded OMI AI for July and August 2007. (d). Differences between
884 Figures 4(b) and 4(c). (e-h) Similar to Figures 4(a)-4(d) but using NAAPS natural runs. Grey
885 color highlights those $1 \times 1^\circ$ (Latitude/Longitude) bins that have less than two collocated NAAPS
886 and OMI AI data for the study period.
887



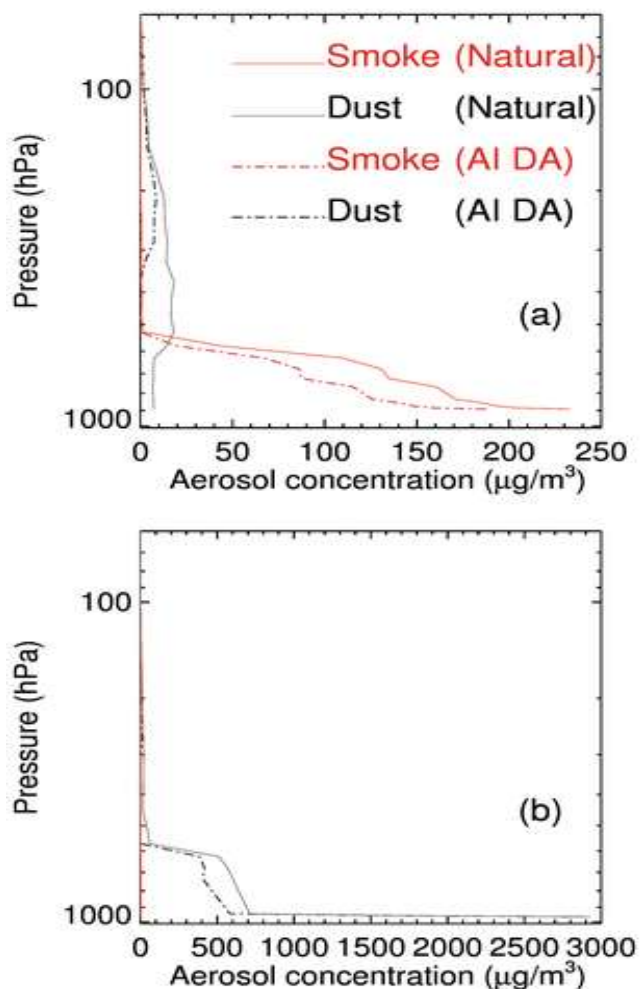
888

889 **Figure 5.** (a). Scatter plot of AERONET and NAAPS AOD ($0.55 \mu\text{m}$) using NAAPS data from
890 the natural runs for July-August 2007 over the study region. (b). Similar to Figure 5(a) but using
891 NAAPS data from the AI-DA runs. (c). Similar to Figure 5(a) but with AODs taken from the
892 NAAPS reanalysis.
893



894

895 **Figure 6.** Spatial distributions of simulated AI at 12 Z on July 28, 2007 using NAAPS reanalysis
896 data, with single scattering albedos of smoke aerosol at 354 and 388 nm taken to be: (a) 0.84 and
897 0.84; (b) 0.85 and 0.85; (c) 0.86 and 0.86; (d) 0.85 and 0.85; (e) 0.85, 0.855; (f) 0.85 and 0.86.
898



899

900 **Figure 7.** (a). Vertical distributions of smoke and dust aerosol concentrations over 9.5°S and
901 10.5°E at 12 Z on July 28, 2007 for both natural and AI DA runs. (b). Similar as (a) but over
902 25.5°N and 12.5°W.

903

904