1 On the relationship between the mesospheric sodium layer and the meteoric input function

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10 **Abstract**

12 This study examines the relationship between the concentration of atmospheric sodium and its
13 Meteoric Input Function (MIF). We use the measurements from the Colorado State University (CSU)
14 Lidar and the Andes Lidar Observatory (ALO) with a new numerical model that includes sodium
15 chemistry in the mesosphere and lower thermosphere (MLT) region. The model is based on the
16 continuity equation to treat all sodium-bearing species and runs at a high temporal resolution. The
17 model simulation employs data assimilation to compare the MIF inferred from the meteor radiant
18 distribution and the MIF derived from the new sodium chemistry model. The simulation captures the
19 seasonal variability of sodium number density compared with lidar observations over the CSU site.
20 However, there were discrepancies for the ALO site, which is close to the South Atlantic Anomaly (SAA)
21 region, indicating it is challenging for the model to capture the observed sodium over ALO. The CSU site
22 had significantly more lidar observations (27,930 hours) than the ALO sites (1872 hours). The simulation
23 revealed that the uptake of the sodium species on meteoric smoke particles was a critical factor in
24 determining the sodium concentration in MLT, with the sodium removal rate by uptake found to be
25 approximately three times that of the $NaHCO_3$ dimerization. Overall, the study's findings provide
26 valuable information on the correlation between MIF and sodium concentration in the MLT region,
27 contributing to a better understanding of the complex dynamics in this region. This knowledge can
28 inform future research and guide the development of more accurate models to enhance our
29 comprehension of the MLT region's behavior.

30

31 **Keywords:** Sodium layer, sodium chemistry, Meteor radiant distribution, Meteoroid input function

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33 **Key points:**
34 ● A high-time resolution, time-dependent Na chemistry model is developed.
35 ● Ablated global meteoroid material inputs inferred from ALO and CSU observations are about 83
36   t d^{-1} and 53 t d^{-1}, respectively.
37 ● The frequency of meteor occurrences might not provide a precise reflection of the mass of
38   meteoroid material input.

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1. Introduction

Micro-meteoroids enter the Earth's atmosphere day and night, depositing their constituents into the atmosphere via ablation, creating a region that hosts various metal species, for example, Fe, K, Si, Mg, Ca, and Na, in both neutral and ion form (Plane et al., 2015; Plane et al., 2021; and references therein). The region is commonly referred to as the mesosphere and lower thermosphere (MLT), located between 75 and 110 km altitude. The metal layers in the MLT often serve as the tracers that facilitate the investigation of the dynamical and chemical processes within the region (Takahashi et al., 2014; Qiu et al., 2021). Quantitative measurements of metal atoms have been made since the 1950s (Hunten, 1967) through a variety of ground or space-borne technologies (Koch et al., 2021; Koch et al., 2022). The large resonant scattering cross-section (Bowman et al., 1969) and the substantial presence of the sodium atom in the MLT make it one of the most researched metal layers in the atmosphere (Yu et al., 2022).

The sodium layer is usually studied via observations carried out by resonance lidars, satellites, and through Na D-line emission at 589.0nm and 589.6nm (Plane, 2010; Plane et al., 2012; Hedin and Gumbel, 2011; Langowski et al., 2017; Andrioli et al., 2019; Li et al., 2020a). The sodium vertical profiles retrieved by lidars have been commonly used as a tracer to study atmospheric dynamics, e.g., gravity waves, wind shear, etc. The long-term seasonal and short-term diurnal variability of metallic species have been investigated by several studies (Feng et al., 2013; Marsh et al., 2013; Cai et al., 2019a, b; Yu et al., 2022). A typical sodium chemistry scheme consists of neutral chemistry, ion chemistry, and photolysis. The sodium chemistry research in recent years has primarily been based on the sodium chemistry model by Plane (2004), which has been cited in various subsequent works, including Bag et al. (2015) and references therein.

As meteoroids are the primary source of metal layers in the atmosphere, including the sodium layer, the Meteoroid Input Function (MIF) plays a crucial role in the modeling of metallic layers in the atmosphere. The MIF is a function designed to comprehend the impact of the temporal and spatial variability of the meteoroid on the atmosphere (Pifko et al., 2013). Sporadic meteors are estimated to make up more than 95% of the total meteoroid population by comparing the number of meteors originating in sporadic sources to those originating in known shower meteor sources (Chau and Galindo, 2008). This highlights the importance of incorporating sporadic meteor data in the MIF to accurately understand sodium concentration in the mesosphere and lower thermosphere (MLT) region and its correlation with meteoroid material input. It is well established that there are six apparent sources of sporadic meteors, namely North and South Apex (NA and SA); North and South Toroidal (NT and ST); and Helion and Anti-Helion sources (H and AH) (Campbell-Brown, 2008; Kero et al., 2012; Li et al., 2022). However, the relative strength of these meteor radiant sources varies among the studies. For example, the NA and SA sources are found to be much stronger than other sources in results obtained with High Power Large Aperture (HPLA) radars (Chau et al., 2007; Kero et al., 2012; Li and Zhou, 2019), while specular meteor radars found the difference to be much smaller (Campbell-Brown and Jones, 2005; Campbell-Brown, 2008). The detection sensitivity varies significantly among different facilities. For instance, the Arecibo Observatory (AO) at 18° N, 66° W detects approximately 20 times more meteors per unit area per unit time than the Jicamarca Radio Observatory (JRO) at 12° S, 77° W, and at least 800 times more meteors than the Resolute Bay Incoherent Scatter North (RISR-N) radar at 75° N, 95° W, despite all being HPLA facilities (Li et al., 2020, 2023a; Hedges et al., 2022) Of note, meteor flux varies with time and latitude, but the variations cannot account for such a large difference.

83 Consequently, the total mass of the meteors that enter the Earth's atmosphere is subject to significant
84 uncertainties. In the existing Whole Atmosphere Community Climate Model-Na (WACCM-Na) global
85 sodium model (Dunker et al., 2015), the meteor input function was modeled by placing a flux curve on
86 each radiant meteor source with a definite ratio (more details can be found in Marsh et al., 2013). The
87 flux curve model is based on observations carried out exclusively by the Arecibo Observatory. Although
88 the model can reproduce some of the flux characteristics of the meteors observed at Arecibo, it is a
89 relatively simple model and therefore has several limitations (Li et al., 2022). One of the limitations is
90 that the model cannot reproduce the velocity distribution of the meteors in observations.

91 This study introduces a new numerical model for sodium chemistry that utilizes the continuity equations
92 for all Na-related reactions without steady-state approximations. The main objective is to investigate the
93 relationship between the apparent sodium concentration and the Meteoroid Input Function (MIF) in the
94 MLT region. We then compare the results of the new model with measurements from two lidar
95 instruments, namely the Colorado State University (CSU) and the Andes Lidar Observatory (ALO).
96 Furthermore, we compared the MIF derived from the new sodium chemistry model and lidar
97 measurements from CSU and ALO, against the results of the high-resolution meteor radiant distribution
98 recently deduced from observations conducted at AO. Finally, we discuss the implications of these
99 comparisons and suggest possible explanations for the observed discrepancy between the MIF derived
100 from radar and those obtained from lidar observations.

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102 **2. The sodium chemistry model (NaChem)**

103 **2.1 Sodium chemistry**

104 Numerical airglow models have been extensively used to investigate atmospheric airglow chemistry and
105 gravity waves (Huang and Hickey, 2008; Huang and Richard, 2014; Huang, 2015). A new numerical
106 sodium chemistry model, hereafter referred to as NaChem, was developed for this study. Table 1 lists
107 the complete reactions and their corresponding rate coefficients used in NaChem, which includes
108 neutral chemistry, ion chemistry, and photochemistry. The dimerization reaction of $NaHCO_3$ (reaction 25
109 in Table 1) is the outlet that removes Na atoms in the chemistry scheme. The Na atoms can also be
110 removed by the uptake of sodium species onto meteoric smoke particles (Hunten et al., 1980;
111 Kalashnikova et al., 2000; Plane, 2004), a process that can be turned on or off in the model. This study
112 estimates the MIF in the numerical model by matching the dimerization reaction to maintain the
113 observed sodium present in the MLT. Throughout the rest of the paper, the MIF estimated from the
114 sodium chemistry numerical model will be referred to as MIF(s). On the other hand, the MIF derived
115 from meteor radiant distribution will be referred to as MIF(m). MIF is a function of time and represents
116 the mass of meteoroid material entering Earth's atmosphere. The MIF(m) is determined through a 3-D
117 meteoroid orbital simulation based on the meteor radiant distribution.

118 The numerical model utilizes the continuity equation to track the time evolution of all 14 Na-related
119 species. Table 2 presents a comprehensive list of these species, along with their corresponding
120 production and loss rates. The background major gas species, including O3, O2, O, H, H2, H20, etc., and
121 the temperature are provided by WACCM. Here we use the dynamic version of WACCM nudged with
122 NASA's Modern Era Retrospective Analysis for Research and Application MERRA2 reanalysis data set
123 (Hunziker & Wendt, 1974; Molod et al., 2015; Gettelman et al., 2019). The WACCM reference profiles

124 are linearly interpolated to a resolution of one minute and updated every minute during the simulation.
125 It is worth noting that the Na-related reactions, which are illustrated in Table 2, do not significantly
126 impact the background gas species, as the effect is orders of magnitude smaller than the variation of the
127 major gas species themselves. Therefore, the major gas species are simulated independently of Na-
128 related reactions.

129 2.2 Numerical scheme

130 As discussed earlier, it is worth noting that the reactions of sodium chemistry in NaChem share
131 similarities with those in previous models (e.g., Plane et al., 2015 and references therein); however, the
132 implementation of the numerical chemistry scheme differs. NaChem uses continuity equations to treat
133 all chemicals involved, including short-lived intermediate species. Treating all species with the continuity
134 equation is a more straightforward yet accurate approach than using steady-state approximations.
135 Moreover, by treating all species in a uniform procedure, the numerical model is more compact and
136 easier to interpret. The computational capability of a personal computer nowadays has advanced
137 enough to process an ultra-fine time step (microseconds) that is necessary for numerical simulations of
138 short-lived species in a reasonable duration. Still, the differential equations for production and loss of
139 short-lived species can be numerically unstable unless microsecond or even sub-microsecond time step
140 is used (Higham, 2002). The concern of the differential equation instability can be largely mitigated by a
141 first-order exponential integrator (Hochbruck and Ostermann, 2010), i.e.,

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$$c = x_0 - \frac{a_0}{b_0}$$

(1)

$$x_1 = \frac{a_0}{b_0} + ce^{-b_0 \Delta t}$$

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144 Where x_0 is the value of the current step, a_0 is the production of the species, b_0 is the loss of the
145 species, Δt is the step size in time, and x_1 is the value of the next step.

146 The exponential integrator, as expressed in Equation (1), is the solution to the continuity equation.
147 Notably, reaction 25 listed in Table 1 is an exception, which was carried out using explicit Euler
148 integrator in the simulation. This reaction's continuity equation is structured differently from the others
149 because it represents the only mechanism for removing Na atoms from the chemistry simulation, apart
150 from the uptakes of sodium species. Our testing indicates that both the exponential integrator and
151 explicit Euler integrator yield nearly identical results. However, for numerical stability, the explicit Euler
152 integrator requires a step size of ~1μs, which is orders of magnitude smaller than the exponential
153 integrator. The default time step of NaChem is 0.1 seconds with the exponential integrator.

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160   Table 1. Reactions in NaChem. f_a and f_x are branching ratios.

	Reaction	Rate Coefficient	reference
neutral chemistry			
1	$Na + O_3 \rightarrow NaO(A) + O_2$	$K_1 = 1.1 \times 10^{-9} \exp(-116/T)$	1
2	$NaO(A) + O \rightarrow Na(^2P) + O_2$	$K_2 = 2.2 \times 10^{-10}(T/200)^{0.5}$, $f_A = 0.14 \pm 0.4$	1,3
3	$NaO(A) + O \rightarrow Na(^2S) + O_2$	$K_3 = 2.2 \times 10^{-10}(T/200)^{0.5}$, $(1-f_A)$	1,3
4	$NaO(A) + O_2 \rightarrow NaO(X) + O_2$	$K_4 = 1 \times 10^{-11}$	1
5	$Na + O_2 + M \rightarrow NaO_2 + M$	$K_5 = 5.0 \times 10^{-30}(200/T)^{1.22}$	1
6	$NaO_2 + O \rightarrow NaO(X) + O_2$	$K_6 = 5 \times 10^{-10}\exp(-940/T)$	1
7	$NaO(X) + O \rightarrow Na(^2P) + O_2$	$K_7 = 2.2 \times 10^{-10}(T/200)^{0.5}$, $f_x = 0.167$	1,2
8	$NaO(X) + O \rightarrow Na(^2S) + O_2$	$k_8 = 2.2 \times 10^{-10}(T/200)^{0.5}$, $(1-f_x)$	1,2
9	$NaO(X) + O_3 \rightarrow NaO_2 + O_2$	$k_9 = 1.1 \times 10^{-9}\exp(-568/T)$	1
10	$NaO(X) + O_3 \rightarrow Na + 2O_2$	$k_{10} = 3.2 \times 10^{-10}\exp(-550/T)$	1
11	$NaO(X) + O_2 + M \rightarrow NaO_3 + M$	$k_{11} = 5.3 \times 10^{-30}(200/T)$	1
12	$NaO(X) + H \rightarrow Na + OH$	$k_{12} = 4.4 \times 10^{-10}\exp(-668/T)$	1
13	$NaO(X) + H_2 \rightarrow NaOH + H$	$k_{13} = 1.1 \times 10^{-9}\exp(-1100/T)$	1
14	$NaO(X) + H_2 \rightarrow Na + H_2O$	$k_{14} = 1.1 \times 10^{-9}\exp(-1400/T)$	1
15	$NaO(X) + H_2O \rightarrow NaOH + OH$	$k_{15} = 4.4 \times 10^{-10}\exp(-507/T)$	1
16	$NaO(X) + CO_2 + M \rightarrow NaCO_3 + M$	$K_{16} = 1.3 \times 10^{-27}(200/T)$	1
17	$NaO_2 + H \rightarrow Na + HO_2$	$K_{17} = 1.0 \times 10^{-9}\exp(-1000/T)$	1
18	$NaO_3 + O \rightarrow Na + 2O_2$	$k_{18} = 2.5 \times 10^{-10}(T/200)^{0.5}$	1
19	$NaCO_3 + O \rightarrow NaO_2 + CO_2$	$k_{19} = 5.0 \times 10^{-10}\exp(-1200/T)$	1
20	$NaCO_3 + H \rightarrow NaOH + CO_2$	$k_{20} = 1.0 \times 10^{-9}\exp(-1400/T)$	1
21	$NaOH + H \rightarrow Na + H_2O$	$k_{21} = 4.0 \times 10^{-11}\exp(-550/T)$	1
22	$NaOH + CO_2 + M \rightarrow NaHCO_3 + M$	$k_{22} = 1.9 \times 10^{-28}(200/T)^1$	1
23	$NaHCO_3 + H \rightarrow Na + H_2O + CO_2$	$k_{23} = 1.1 \times 10^{-11}\exp(-910/T)$	1
24	$NaHCO_3 + H \rightarrow Na + H_2CO_3$	$k_{24} = 1.84 \times 10^{-13}T^{0.777}\exp(-1014/T)$	1
25	$2NaHCO_3 + M \rightarrow (NaHCO_3)_2 + M$	$k_{25} = 8.8 \times 10^{-10}\exp(T/200)^{-0.23}$	1
26	$Na(^2P) \rightarrow Na(^2S) + h\nu(589.0\text{-}589.6 \text{ nm})$	$K_{26} = 6.26 \times 10^7$	1
ion-molecule chemistry			
27	$Na + O_2^+ \rightarrow Na^+ + O_2$	$K_{27} = 2.7 \times 10^{-9}$	1
28	$Na + NO^+ \rightarrow Na^+ + NO$	$K_{28} = 8.0 \times 10^{-10}$	1
29	$Na^+ + N_2 + M \rightarrow NaN_2^+ + M$	$k_{29} = 4.8 \times 10^{-30}(T/200)^{-2.2}$	1
30	$Na^+ + CO_2 + M \rightarrow NaCO_2^+ + M$	$k_{30} = 3.7 \times 10^{-29}(T/200)^{-2.9}$	1
31	$NaN_2^+ + O \rightarrow NaO^+ + N_2$	$k_{31} = 4.0 \times 10^{-10}$	1
32	$NaO^+ + N_2 \rightarrow NaN_2^+ + O$	$k_{32} = 1.0 \times 10^{-12}$	1
33	$NaO^+ + O \rightarrow Na^+ + O_2$	$k_{33} = 1.0 \times 10^{-11}$	1
34	$NaO^+ + O_2 \rightarrow Na^+ + O_3$	$k_{34} = 5.0 \times 10^{-12}$	1
35	$NaN_2^+ + X \rightarrow NaX^+ + N_2$ (X=CO_2, H_2O)	$k_{35} = 6.0 \times 10^{-10}$	1
36	$NaY^+ + e \rightarrow Na + Y$ (Y=N_2, CO_2, H_2O, O)	$k_{36} = 1.0 \times 10^{-6}(T/200)^{-0.5}$	1
photochemical reactions			
37	$NaO(A)/NaO(X) + h\nu \rightarrow Na + O$	$K_{37} = 5.5 \times 10^{-2}$	1

38	NaO2 + hv -> Na + O2	$K_{38} = 1.9 \times 10^{-2}$	1
39	NaOH + hv -> Na + OH	$K_{39} = 1.8 \times 10^{-2}$	1
40	NaHCO3 + hv -> Na + HCO3	$K_{40} = 1.3 \times 10^{-4}$	1
41	Na + hv -> Na+ + e-	$K_{41} = 2 \times 10^{-5}$	1

*1:Plane (2004), 2: Plane (2012), 3: Griffin et al. (2001).

Table 2. The production and loss terms of the sodium-related species.

	Species	Prod	Loss
a1	Na(2P)	$k_2[a_3][O] + k_7[a_5][O]$;	1*
a2	Na	$k_3[a_3][O] + k_8[a_5][O] + k_{10}[a_5][O_3] + k_{12}[a_5][H] + k_{14}[a_5][H_2] + k_{17}[a_4][H] + k_{18}[a_6][O] + k_{21}[a_7][H] + k_{23}[a_9][H] + k_{24}[a_9][H] + k_{36}[a_{11}][e] + k_{36}[a_{13}][e] + k_{36}[a_{12}][e] + k_{36}[a_{14}][e] + [a_1] + k_{37}[a_3][hv] + k_{37}[a_5][hv] + k_{38}[hv][a_4] + k_{39}[hv][a_7] + k_{40}[hv][a_9]$;	$k_1[O_3] + k_5[O_3] + k_5[O_2][M] + k_{27}[O_2^+] + k_{28}[NO^+] + k_{41}[hv]$;
a3	NaO(A)	$k_1[a_2][O_3]$	$k_2[O] + k_3[O] + k_4[O_2] + k_{37}[hv]$
a4	NaO$_2$	$k_5[a_2][O_2][M] + k_9[a_5][O_3] + k_{19}[a_8][O]$	$k_6[O] + k_{17}[H] + k_{38}[hv]$
a5	NaO(X)	$k_5[a][O_3] + k_4[a_3][O_2] + k_6[a_4][O]$	$k_7[O] + k_8[O] + k_9[O_3] + k_{10}[O_3] + k_{11}[O_2][M] + k_{12}[H] + k_{13}[H_2] + k_{14}[H_2] + k_{15}[H_2O] + k_{16}[CO_2][M] + k_{37}[hv]$
a6	NaO$_3$	$k_{11}[a_5][O_2][M]$	$k_{18}[O]$
a7	NaOH	$k_{13}[a_5][H_2] + k_{15}[a_5][H_2O] + k_{20}[a_8][H]$	$k_{21}[H] + k_{22}[CO_2][M] + k_{39}[hv]$
a8	NaCO$_3$	$k_{16}[a_5][CO][M]$	$k_{19}[O] + k_{20}[H]$
a9	NaHCO$_3$	$k_{22}[a_7][CO_2][M]$	$k_{23}[H] + k_{24}[H] + 2k_{25}[a_9][M] + k_{40}[hv]$
a10	Na$^+$	$k_{27}[a_2][O_2^+] + k_{28}[a_2][NO^+] + k_{33}[a_{13}][O] + k_{34}[a_{13}][O_2] + k_{41}[hv][a_2]$	$k_{29}[N_2][M] + k_{30}[CO_2][M]$
a11	NaN2$^+$	$k_{29}[a_{10}][N_2][M] + k_{32}[a_{13}][N_2]$	$k_{31}[O] + k_{35}[CO_2] + k_{35}[H_2O] + k_{36}[e]$
a12	NaCO2$^+$	$k_{30}[a_{10}][CO_2][M] + k_{35}[a_{11}][CO_2]$	$k_{36}[e]$
a13	NaO$^+$	$k_{31}[a_{11}][O]$	$k_{32}[N2] + k_{33}[O] + k_{34}[O_2] + k_{36}[e]$
a14	NaH2O$^+$	$k_{35}[a_{11}][H_2O]$	$k_{36}[e]$

*In Species 1, as of the current state of the model, all Na(2p) atoms return to their ground state immediately, so the loss term is set to 1. The [hv] is the term that represents loss via photon emission, which follows a sinusoidal function based on the zenith angle of the respective local time.

3. CSU and ALO Sodium Lidar Observations and data processing

3.1 Observations

Several aspects of the current research, i.e., the presence of sodium in the MLT, require cross-validation with the measurements. One primary objective of the present model is to match the observed seasonal variation of the sodium layer. Measurements by the Colorado State University (CSU, 41.4°N, 111.5°W) Lidar, also known as Utah State University (USU) Lidar, and the lidar data acquired by the Andes Lidar Observatory (ALO, 30.3°S, 70.7°W), are used to facilitate the research in the current study. We are unable to acquire more ALO data after 2019 as the COVID situation disrupted the site operation. The CSU data comprises 27,930 hours of lidar observations between 1990 and 2020, whereas the ALO data consists of 1872 hours between 2014 and 2019.

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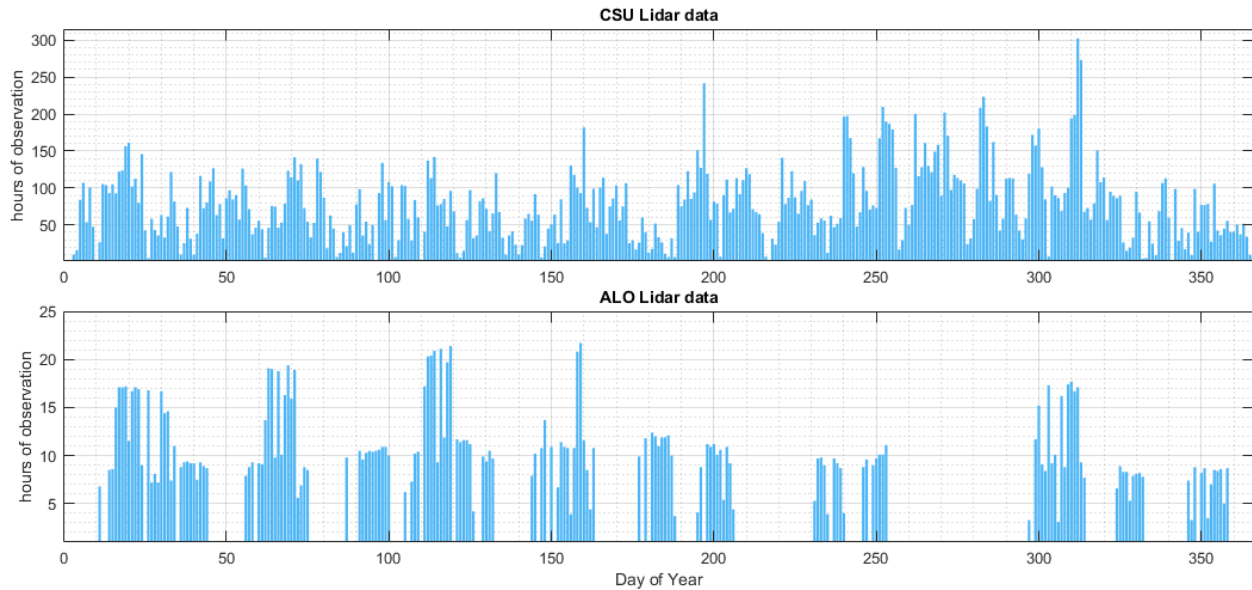
Figure 1. CSU lidar data from 1990 to 2020 (top plot) and ALO lidar data from 2014 to 2019 (bottom
plot).

The statistics of CSU and ALO available data are presented in Figure 1. The Lidar observations of both
sites consist of nocturnal observations only, and a typical nocturnal observation lasts between 8 and 11
hours. Note that in Figure 1, there could be as many as 300 hours of sodium observations on a single day
of year, which means the data of the date comprise observations of many years on that day in different
years. The CSU data almost covered every day of the year with only a few exceptions, whereas the ALO
data was much more sparse. As a result, due to the significantly larger number of CSU observations, the
statistical reliability of the seasonal variation in the sodium layer derived from ALO observations may not
be as strong as that of the CSU data. As depicted in Figure 2, the overall seasonal trend of the sodium
vertical profile derived from CSU lidar observations closely aligns with the simulation-based estimate by
Marsh et al. (2013). In contrast, the results of ALO lidar observations diverge from the findings reported
by Marsh et al. (2013).

3.2 Data processing

The sodium layer in atmospheric observations is often affected by perturbations of atmospheric
dynamics, which is why sodium is commonly used as a tracer in the study of the MLT dynamics (Plane et
al., 2015). However, studying the sodium layer itself can be complicated due to the underlying chemical
processes coupled with the dynamics. In order to mitigate the effects of atmospheric dynamics, we
process the sodium vertical profiles from observations in three steps. First, we average the profiles by
day of the year, meaning we take the average of the data from the same day of the year from different
years. Missing data are treated using linear interpolation. Next, we smooth the averaged profiles using a
15-day running average. Finally, we further smooth the profiles by fitting them with a skew-normal
distribution (Azzalini & Valle, 1996) using the least squares error method.

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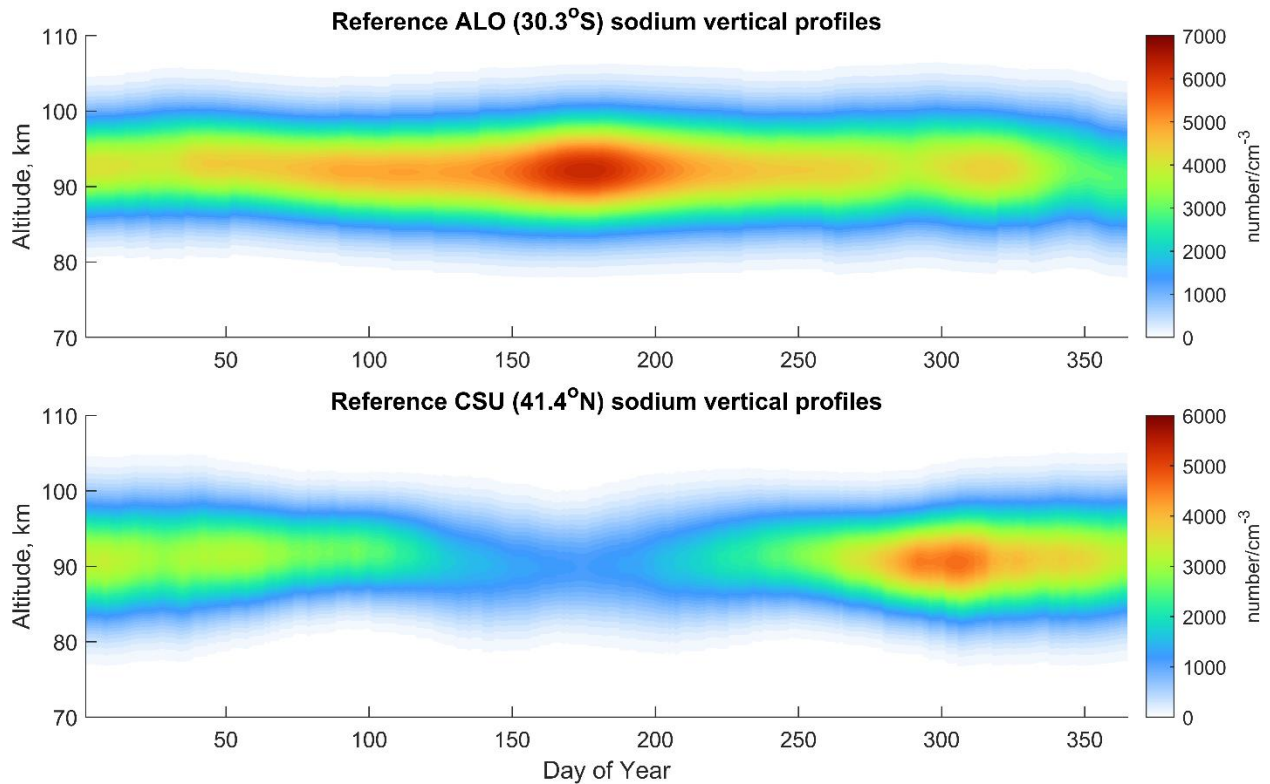
Figure 2. *The reference annual sodium vertical profiles at ALO (top plot) and at CSU (bottom plot). The reference profiles are the averages throughout all the available data on the same days at the respective site, then fitted by a skew-normal distribution that mitigates atmospheric dynamics. In essence, the reference profiles are measurements with small-scale dynamics removed via steps discussed in section 3.2.*

Figure 2 displays the processed annual sodium vertical profiles from the lidar measurements, referred to as reference profiles hereafter. These profiles serve as references to guide the numerical simulation of the NaChem model. The reference profiles are Na lidar measurements fitted using a skew-normal distribution, smoothed by a 15-day running average, and processed through linear 2-D interpolation across time and altitude. The lidar measurements have an altitude resolution of 500m for ALO and from 75m to 140m for CSU. These measurements are interpolated to a 100m resolution as inputs to the NaChem model. The time resolution is interpolated to 0.1 seconds. The reference profiles inherently include diffusion and other dynamic effects on the sodium species in the MLT, as these observational data represent snapshots of sodium diffusion at various times. By constantly matching the observed Na profile to the simulated Na profile, the diffusion is included implicitly in the model. The seasonal column densities of both ALO and CSU profiles are similar to a sinusoidal function, with ALO data peaking near June and CSU data peaking in November. The centroid height of the sodium layer is higher in the ALO data than in the CSU data.

4. Results

4.1 Sensitivity test

Sodium in the atmosphere could manifest in many forms, i.e., in sodium-bearing neutral chemicals and ionic chemicals. The sodium number densities are typically obtained via lidar measurements. Given the complexity of the sodium chemistry, the observed sodium is merely a subset, possibly not even a major constituent, of the total number of all the sodium-bearing species in the atmosphere. The total sodium content is defined as the total number of sodium atoms in all 14 sodium-bearing species, as listed in Table 2. In summary, the sodium that we can detect does not necessarily provide an accurate representation of the total sodium content or the overall count of sodium-bearing species, as unobservable species such as Na^+ and $NaHCO_3$ could constitute a substantial portion of the total sodium content.

Understanding the impact of each background species, i.e., species listed in Figure 3., on the total sodium content is essential to study the underlying mechanism of the chemical reactions. Therefore, we present a sensitivity test by isolating variables. The sensitivity test is done by altering the number density of background species in question by two orders of magnitude, i.e., with a factor of 0.1 and 10, while keeping the number densities of other background species and the atomic sodium fixed. The simulation is kept running until all the numbers are stable. The diurnal variations of the sodium and background species are not considered in sensitivity test as they introduce unnecessary complexity. The results of the sensitivity test of the 11 background species and temperatures involved in the numerical simulation are shown in Figure 3. Each panel contains three lines, where the red curve shows the unaltered vertical profile of the total sodium content. The results of the species altered by the factor of 0.1 and 10 are shown in light blue and yellow, respectively.

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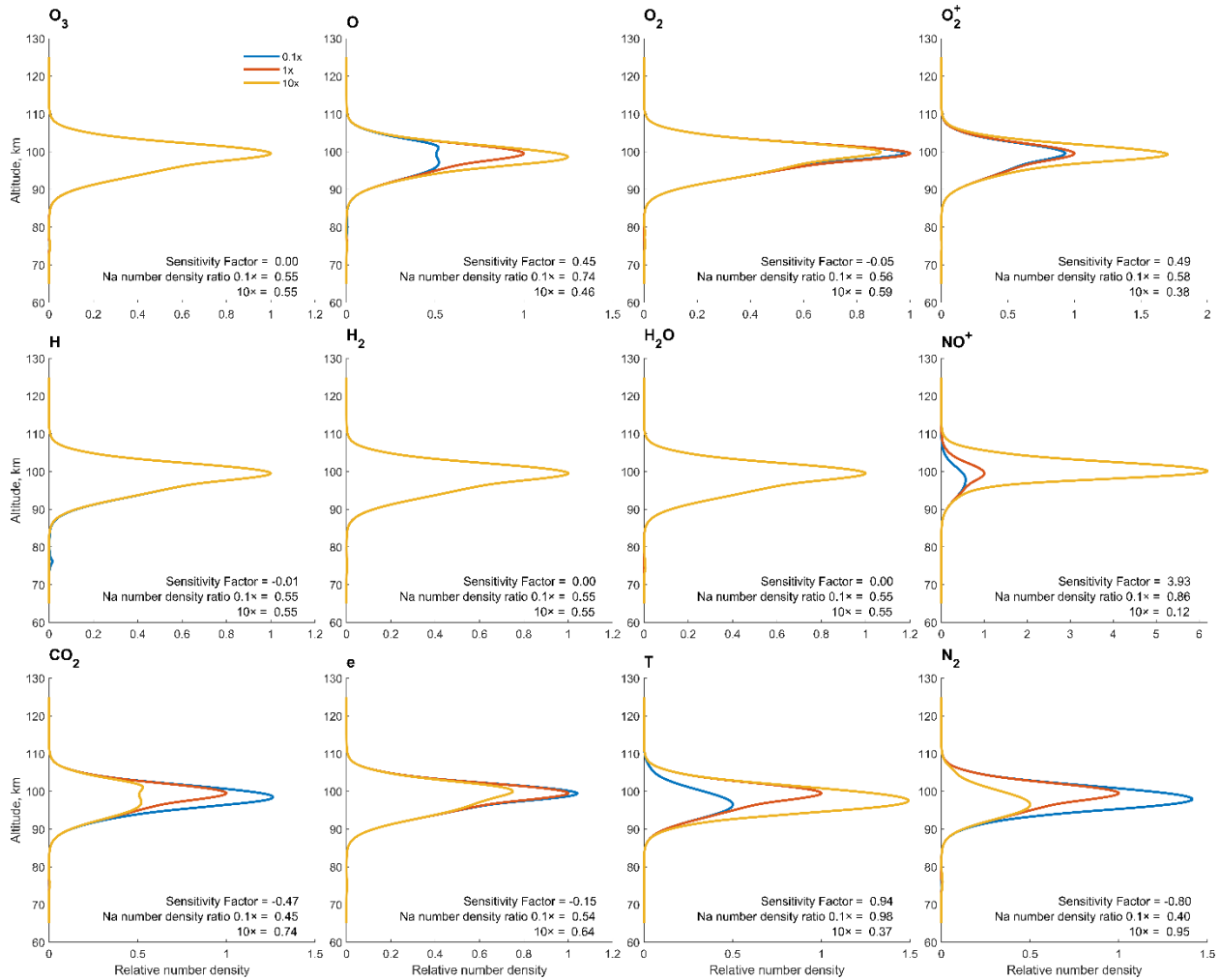


Figure 3. Sensitivity test of 11 background species and temperature on Na chemistry. The total sodium content vertical profile for the respective background species altered by 10x and 0.1x are shown in yellow and light blue. The reference sodium content vertical profiles are shown in red. Additionally, the sensitivity factor and the Na number density ratio to the concentration of all sodium species are presented on each panel.

In Figure 3, only the yellow curve is visible in some of the panels because the three curves are drawn on top of each other, indicating that the change of the respective background species bears little to no effect on the sodium chemistry. A sensitivity factor is defined to better quantify the weight of each background species in sodium chemistry. The factor is measured by the following equation:

$$Sensitivity\ Factor = \frac{NaT_c^{10} - NaT_c^{0.1}}{NaT_c} \tag{2}$$

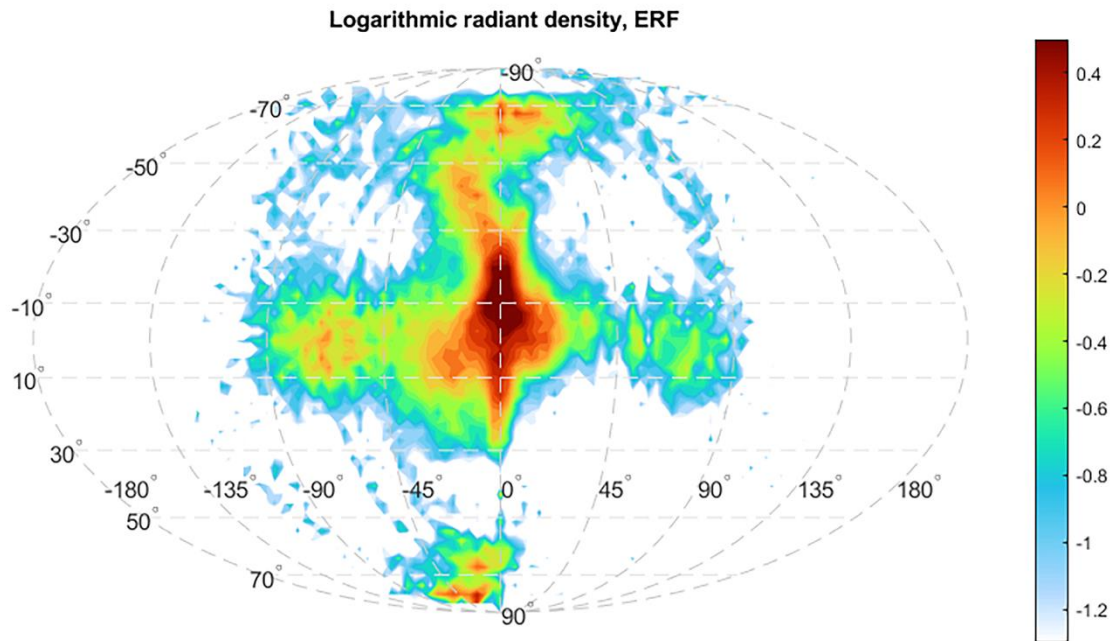
Where NaT_c^{10} is the column density of the total sodium content with the respective species altered by a factor of 10, and $NaT_c^{0.1}$ is the same operation as the previous one but altered by a factor of 0.1. The denominator, NaT_c , is the column density of the reference profile. For example, a Sensitivity Factor of 5 indicates that the total sodium content increases by five times when the respective background species

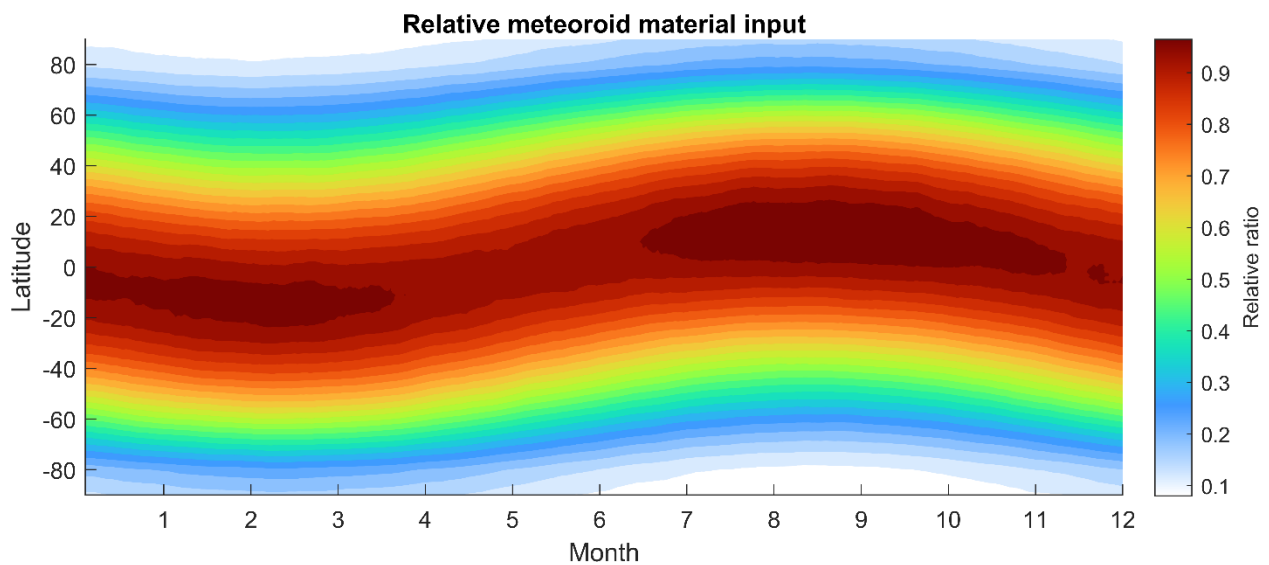
262   increases 100 times. A positive sensitivity factor indicates the total sodium content is positively
263   correlated to the respective species and vice versa. The reference profile is the total sodium content in
264   steady-state in the background condition of the midnight new year of 2002, giving a typical sodium
265   vertical profile similar to the one shown in Figure 5 of Plane (2004). In the simulation, a greater total
266   sodium content implies that a smaller percentage of the sodium chemicals are present as sodium atoms
267   as the altitude profile of the sodium atoms is fixed. In reality, instead of the sodium atoms, the total
268   sodium content should be more or less conserved. Hence a higher total sodium content in our
269   simulation suggests less sodium can be detected by the lidar.

270   Although the sensitivity factor could be different upon the change of the reference profile, it still gives
271   an insight into the significance of each background species to the sodium chemistry. Apparently, the
272   weight of some background species, namely O_3, H, H_2, and H_2O, is negligible in sodium chemistry,
273   meaning that removing these species and their associated reactions has no effect on the overall sodium
274   chemistry. Nevertheless, these species are still kept in our numerical model for completeness. The
275   impact of species that convert Na atom to Na^+, as listed in reactions 27 and 28 of Table 1, is generally
276   strong. The effect of NO^+, in particular, is the most significant according to the sensitivity factor, greater
277   than the combined effect of all the other species. Consequently, the number density of the sodium atom
278   by lidar observation is strongly correlated with the fluctuations of the NO^+. In a nutshell, more NO^+ will
279   directly lead to fewer observable Na atoms. That being said, the interaction between sodium and
280   background species is rather complex. The scope of the sensitivity factor in the present paper was
281   limited to column density. As a result of such, variations and behaviors of the sodium chemicals by
282   altitude are overlooked. The actual impact of the background species may differ at different altitudes.

283   **4.2 Meteor input function**

284   The estimation of meteoric influx is subject to many uncertainties among different techniques (Li et al.,
285   2022). Moreover, the meteor flux estimated by the sodium chemistry model also varies (Marsh et al.,
286   2014; Plane et al., 2015). The previous model of Plane (2004) and the following similar models indicate
287   that the rate of dimerization, or the speed of removing sodium from the system, is heavily correlated to
288   the vertical transport in the MLT. The NaChem model does not explicitly incorporate vertical transport,
289   but the vertical transport by diffusion is inherently embedded within the input of the observed sodium
290   vertical profile.

291   Unlike the previous models (Plane 2004; March et al. 2014; and references therein), the present
292   NaChem model took an indirect route to estimate the meteor mass input. During the simulation, the
293    $NaHCO_3$ dimerization and the uptake of the sodium species on meteoric smoke particles, which can be
294   turned on or off, create a deficit of sodium atoms. Meanwhile, a meteor input function injects an
295   appropriate amount of sodium atoms so that the present sodium vertical profile always matches the
296   reference profiles. This is carried out by finding the difference between the current sodium profile (with
297   the deficit) and the corresponding reference profile in every iteration and then replacing the former
298   with the latter. The diffusion coefficient is found to be highly correlated with the sink rate of the
299   dimerization reaction with large uncertainties (Plane, 2004). The simulation circumvents this uncertainty
300   by directly incorporating the observational sodium vertical profile, given that diffusion is already
301   inherently in the measurements.

302

303 **Figure 4.** *Meteor radiant source derived from the AO observations. The result is in the Earth Reference*
304 *Frame (ERF), equivalent to ground-based observations. The latitude of the ERF is centered on the ecliptic*
305 *plane. The longitude of the ERF is centered to the Apex direction, the moving direction of the Earth,*
306 *where the highest number of meteors encounter Earth. The radiant distribution is derived from the*
307 *number of meteor events. Figure reproduced from Li et al. (2022).*

308

309 **Figure 5.** *Relative seasonal and latitudinal meteoroid input, inferred from the radiant source distribution*
310 *shown in Figure 4.*

311

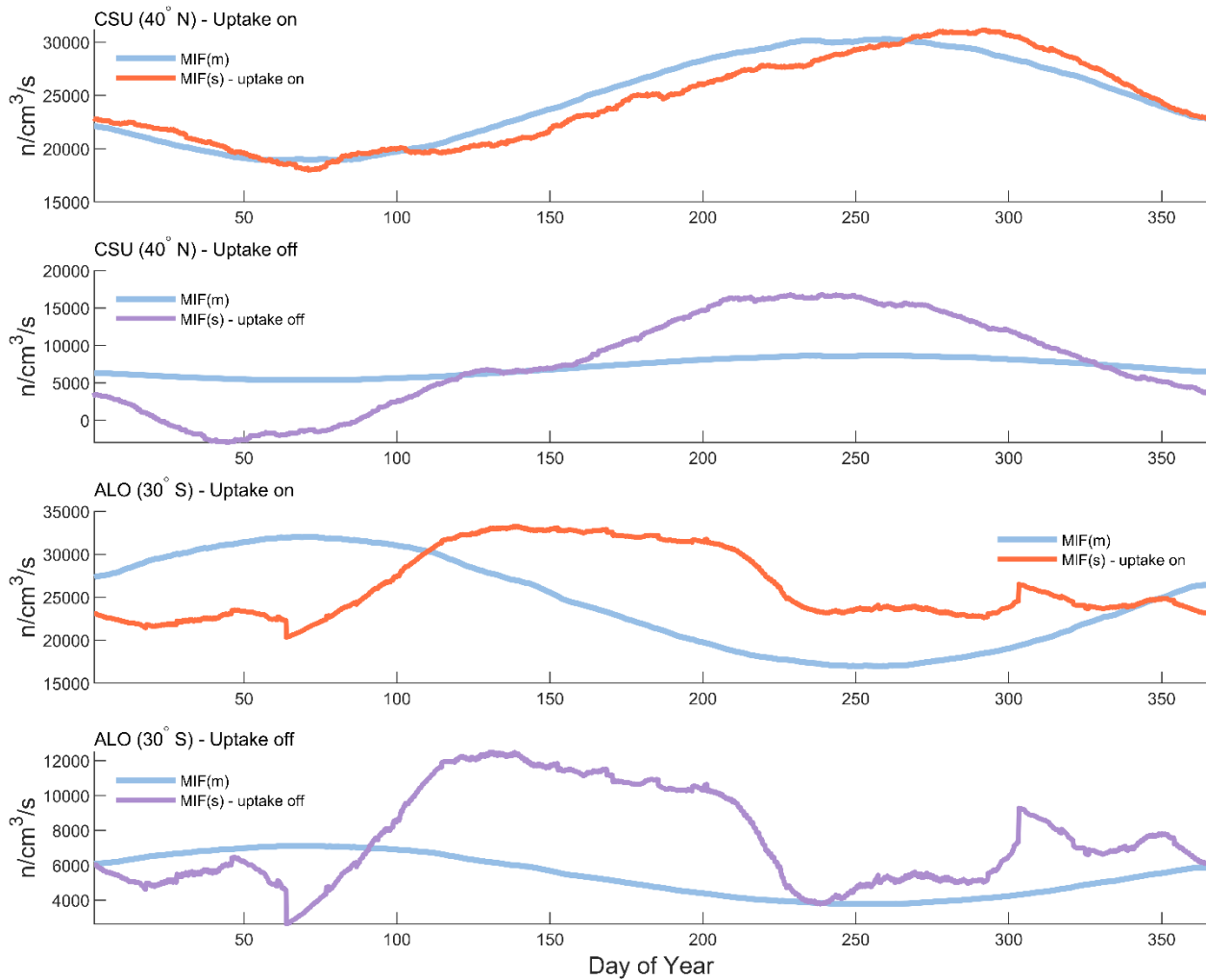



Figure 6. *A comparison between two meteor input functions: MIF(m), which is inferred from micro-meteor radiant distribution, and MIF(s), which is derived using a Na chemistry model. The purple line represents MIF(s) with uptake on, while the orange dotted line represents MIF(s) with uptake off. The MIF(m) is linearly scaled to match each curve.*

Figure 4 shows the high-resolution meteor radiant source distribution recently inferred from the AO observations (Li et al., 2022). The typical mass of the Arecibo meteors is estimated to be around 10^{-13} kg based on flux rate (Li and Zhou, 2019). Mathews et al. (2001) estimated the limiting meteor mass of 10^{-14} kg based on the meteor ballistic parameter. Limiting mass is the smallest mass a meteoroid must have to generate sufficient ionization to be detected by radar. Despite these estimations being based on various simplified assumptions that may lead to inaccurate results, the estimated mass is still at least two orders of magnitude smaller than the estimations of other facilities by similar means. More than 95% of the meteoroid population in the Earth's atmosphere is found to be sporadic meteors by HPLA radar observation (Chau and Galindo, 2007), which typically are low-mass meteors evolved from the outer Solar system due to the Poynting-Robertson drag (Nesvorný et al., 2011; Koschny et al., 2019).

328   That being said, the percentage of sporadic meteors, as well as the radiant source distribution, are both
329   estimated based on the occurrence. However, the occurrence of sporadic meteors may not be able to
330   represent their mass distribution. The relative seasonal and latitudinal meteoroid input by the number
331   of occurrences inferred from the new radiant distribution is depicted in Figure 5. The meteoroid input
332   generally follows a sinusoidal pattern and differs from the one used in the previous work, as shown in
333   Figure 1 of Marsh et al. (2013). Although the interplanetary dust (meteor) background on the Earth's
334   orbit could vary in different locations due to a variety of reasons, e.g., Jupiter resonance, it is still safe to
335   assume no change in the interplanetary dust background for our purpose. Taking a stable interplanetary
336   dust background, the MIF(m) 's seasonal sinusoidal pattern should follow the Earth's axis rotation.

337   Figure 6 shows a comparison between two types of meteor input function: MIF(m), which is inferred
338   from the micro-meteor radiant distribution, and MIF(s), which is derived using the Na chemistry model
339   with sodium input from the lidar observations. For the MIF(s) model simulations, we did two scenarios,
340   one with and one without uptake by smoke particles, for the ALO and CSU data. The MIF(s) with uptake
341   by smoke particles exhibit a good match with the MIF(m) on the CSU dataset, while it does not show as
342   good of a match on the ALO dataset. The MIF(s) with smoke uptake on is represented by a purple line,
343   while the MIF(s) with smoke uptake turned off is depicted by an orange dotted line. The MIF(s) could go
344   negative when the reference sodium vertical profile decreases faster than the removal rate by the
345   dimerization, as shown in the orange dotted line in Figure 6, indicating that the dimerization process
346   alone is not sufficient enough to account for all the sodium atom depletion in the MLT region. MIF(m) is
347   derived from a global micro-meteor radiant distribution model, as depicted in Figure 4 and Figure 5. The
348   smoke uptake of sodium species in this study is implemented using a methodology similar to Plane
349   (2004), but instead of applying smoke uptake solely to the three major sodium species, namely Na,
350   NaHCO3, and Na+, it is applied to all 14 sodium-bearing species. The optimal uptake factor to obtain the
351   best results was found to be 2×10^{-2}/km/s. The smoke uptake and $NaHCO_3$ dimerization account for
352   approximately 75% and 25% of the Na sink, respectively.

353   According to the global meteoroid orbital model outlined in Li et al., (2022), the latitudes spanning 29.5°
354   S to 30.5° S (ALO) account for 0.52% of the total meteor input, while those between 39.5° N and 40.5° N
355   (CSU) represent 0.67%. The CSU site shares more meteor input due to its closer proximity to one of the
356   Apex meteor radiant sources. The global total sodium injection rate inferred from the ALO data-based
357   simulation is 2.01×10^{23} atoms per second, and the CSU-data-based simulation suggests a global sodium
358   injection rate of 1.28×10^{23} atoms per second. Assuming the relative sodium elemental abundance in
359   meteoroid material is 0.8% (Vondrak et al., 2008), the deduced total meteoroid material input of ALO-
360   based simulation was 83 t d^{-1}. From CSU-based simulation, the rate is 53 t d^{-1}. Both estimations are close
361   to 80-130 t d^{-1}, the value reported by the Long Duration Exposure Facility (Love and Brownlee, 1993;
362   McBride et al., 1999). It is worth noting that the estimated total daily input of meteoroid materials
363   varies among previous studies, ranging from 4.6 t d^{-1} (Marsh et al., 2013) to 300 t d^{-1} (Nesvorný et al.,
364   2009), with an intermediate value of 20 t d^{-1} reported by Carrillo-Sánchez et al. (2020). While these
365   estimates seem quite disparate, the variance is relatively small given that the daily input rate is derived
366   from combinations of chemicals that can fluctuate by several orders of magnitude. For example, the
367   NO^+, which exhibits the highest sensitivity factor according to the sensitivity test, undergoes diurnal
368   variations of approximately three orders of magnitudes.

369

5. Discussion

The sodium concentration in the sodium layer in the MLT region is governed by several factors, including chemistry, dynamics, and the MIF. It's difficult to discern which of these three components is more important than the others. In this section, we discuss various factors that may contribute to modeling the sodium concentration in the MLT.

The mass of the meteoroids has been estimated and measured using various methods. These include the ballistic parameter derived from meteor deceleration (Mathews et al., 2001), estimation of meteor head echo plasma distribution through a combination of meteor ablation models and radar cross-section measurements (Close et al., 2005; Sugar et al., 2021), flux rate determination (Zhou and Kelley, 1997), as well as spacecraft in-situ measurements (Leinert and Grun, 1990), among others. The mass estimated by the meteor ballistic parameter is commonly referred to as momentum or dynamical mass. The mass estimated by the meteor ablation model is usually called the scattering mass. The meteor momentum mass from Arecibo Ultra-High-Frequency (UHF) radar observation is estimated to be 10^{-14} – 10^{-7} kg, with the typical mass being 10^{-13} kg. On the other hand, the meteor scattering mass is estimated to be 10^{-9} – $10^{-5.5}$ kg by data from EISCAT UHF radar (Kero et al., 2008) and 10^{-7} – $10^{-4.5}$ kg by data from ALTAIR UHF radar (Close et al., 2005). While the detection sensitivity among different facilities differs, these estimations are still off by many orders of magnitude. The assessments of either momentum mass or scattering mass are based on a variety of simplified assumptions. They are subject to errors due to the complexity of radar beam patterns, background atmosphere conditions, aspect sensitivity, meteor radiant sources, and many other possible factors. For example, radar meteor observation is subject to bias against low-mass, low-velocity meteors (Close et al., 2007; Janches et al., 2015).

Another aspect that may contribute to the MIF(m)'s uncertainty is the meteor radiant distribution. The meteor radiant distributions shown in Figure 4 and many others (Chau et al., 2004; Campbell-Brown and Jones, 2006; Kero et al., 2012) are inferred or measured by meteor occurrence instead of mass input. Currently, retrieving a more accurate estimation of the meteor mass input is still a topic under active research, and there is no quantitative study on the disparities between meteor occurrence and meteor mass input. The radiant sources of the meteors are expected to differ by mass as their orbital evolution is highly correlated to their mass. The interplanetary dust interacts with the solar wind while in the Solar System, losing its momentum in the process and evolving into orbits with a smaller semi-major axis and lower eccentricity. The effect is called the Poynting-Robertson effect (Robertson and Russell, 1937), which behaves like a drag force and defines the evolution of interplanetary dust, and it could be the major reason for the existence of sporadic meteors (Li and Zhou, 2019; Koschny et al., 2019). The importance of the Poynting-Robertson effect is highly dependent on the density and mass of the object. By and large, the orbits of the smaller particles evolve exponentially faster. The orbital dynamics of interplanetary particles have been very well summarized in section 2.2 of (Koschny et al., 2019). For the reasons above, the meteor radiant distribution of mass could deviate from the radiant distribution of occurrence. Therefore, the meteor input rates as shown in the blue curves of Figure 6 could be different from those derived from the meteor radiant distribution of mass since they were derived from the meteor radiant distribution by occurrence.

In the sodium chemistry model presented in this work, the MIF is the sole source of sodium, while the sodium sink comprises $NaHCO_3$ dimerization and smoke uptake. The MIF(s) is determined by matching the sink rate of the sodium atoms with the rate of sodium injection. In other words, MIF(s) represents

412 the amount of sodium injection needed to keep the sodium concentration equal to the reference
413 sodium profiles. If the chemical lifetime of sodium in the MLT is short, then the seasonal variation of
414 both the MIF and sodium concentration in the MLT should be similar. After examining Figures 3, 5, and
415 6, it can be observed that the averaged seasonal variation of sodium over the years at both sites (ALO
416 and CSU) does not correspond to the trend of the MIF(m) at their respective latitudes. This may indicate
417 that the chemical lifetime of sodium in the MLT should be relatively long, as there is no immediate effect
418 of MIF(m) on the sodium concentration. The MIF displays a sinusoidal pattern which peaks in March at
419 the ALO's latitude and in August at the CSU's latitude, whereas the sodium layer shows dual peaks in the
420 CSU's lidar observations and one peak in June in the ALO's lidar observations.

421 In this study, the MIF(s) derived from the NaChem simulation, based on the CSU lidar measurements
422 with uptake turned on, was able to match the amplitude of MIF(m) obtained from the meteor radiant
423 distribution. Although the model does not directly incorporate any dynamical processes, the vertical
424 transport by diffusion is implicitly included. The model forces the sodium layer to be the same as the
425 data, which are averaged by the observations of many years, in which the diffusions are inherently
426 embedded. The combination of observational data with the numerical chemistry model in this paper is a
427 relatively straightforward application of data assimilation (Bouttier & Courtier 2002). The lidar data of
428 both sites (CSU and ALO) indicate that the sodium column density consistently increases by about 20%
429 from 22:00 to 4:00 LT the next day. This can be attributed to the fact that, during nighttime, the large
430 deposits of Na^+ formed by daytime reactions slowly neutralized to Na. As a result, the sodium column
431 density consistently increases throughout the night. The same effect can be reproduced in the NaChem
432 simulation, albeit with a smaller amplitude. The simulation shows the increase to be about 8%. The
433 value is obtained by maintaining a constant total number of sodium-bearing species through the
434 deactivation of the sodium sink.

435 While meridional transport or atmospheric dynamics both contribute to the seasonal variation of the
436 sodium layer in the MLT, the diurnal sodium profile is the mean of observations of thousands of days, of
437 which the variation by atmospheric dynamics should be much less prominent. The lack of explicit
438 dynamics in the model may be one of the sources of inconsistency when compared to the observations.
439 Further, the WACCM, which supplied the background species to the NaChem, is an older version that
440 does not fully incorporate the dynamics of each ion species. Despite our results showing good
441 agreement between the MIF(s) and the MIF(m), there might be several plausible factors that could lead
442 to potential errors. For example, the Na sink by $NaHCO_3$ dimerization varies by the diffusion rate or the
443 vertical transport of sodium atoms in the chemistry model (Plane, 2004). Likewise, the MIF(m) may also
444 differ if the meteoroid mass input differs from the radiant source distribution by the occurrence of
445 meteors, as discussed in the aforementioned paragraph.

446 **5. Conclusion**

447 This work introduced a new sodium chemistry model that simulates the time evolution of all sodium-
448 bearing species using the continuity equation without making any steady-state assumption. The model
449 employs an exponential integrator and runs in high-time resolution to maintain numerical stability. The
450 model is simple to maintain in such a configuration and can be scaled up to include additional
451 capabilities more easily. The model is highly optimized for processing efficiency and benefits from the
452 use of an exponential integrator. Therefore, within an acceptable total CPU time, the NaChem can afford
453 a time resolution of up to milliseconds, several orders of magnitude smaller than those used in other Na

454 models. During our testing, the CPU time to simulated real-time ratio is about 1 to 1000 using a 0.1
455 second time step.

456 The model simulation was able to reproduce the seasonal variation of the sodium layer in the MLT by
457 simulations of chemical reactions. The simulation results at the CSU's latitude capture the general trend
458 of the seasonal variation at the location. The MIF(s) based on the ALO data exhibited less conformity
459 with the corresponding MIF(m), which could be attributed to inadequate statistics of the observational
460 data. Comparably, the CSU dataset is more reliable as the insufficient lidar hours in the ALO dataset may
461 lead to inaccurate statistics. In the simulation, when forcing the sodium layer to be the observation-
462 based reference profile, the inferred MIF is estimated to be 83 t d^{-1} at ALO and 53 t d^{-1} at CSU. The
463 numerical simulation by NaChem could reproduce the general trend of diurnal and seasonal variation of
464 the sodium layer compared to the observations by the CSU Lidar. There are some inconsistencies in
465 MIF(m) and MIF(s) based on data obtained from ALO Lidar. These inconsistencies may have originated
466 from poor statistics resulting from insufficient observation hours.

467 In summary, a new sodium chemistry model has been developed in this work to investigate the
468 relationship between MIF and the sodium layer. We also compared the MIF(m) derived from radar
469 meteor observation to the MIF(s) derived from the chemistry model and lidar observations. Our results
470 indicate that the uptake of sodium species onto meteoric smoke particles removes approximately three
471 times more sodium than the dimerization of NaHCO3. Our future work will focus on incorporating the
472 plausible factors that may lead to potential errors discussed above into the chemistry model.

473

474

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482

483 Code/Data availability

484 The CSU lidar data is available through Utah State University data service (Yuan, 2023). The ALO data is
485 available through the ALO online database (ALO, 2023). The WACCM data used in this work are available
486 through Penn State Scholarsphere (Li, 2023b).

487

488 Author contribution

489 Conceptualization, Yanlin L., Tai-Yin H. and Julio U.; methodology, Yanlin L.; software, Yanlin L.;
490 validation, Yanlin L., Tai-Yin H., Fabio V., Julio U. and Wuhu F.; formal analysis, Yanlin L., Tai-Yin H. and
491 Julio U.; investigation, Yanlin L., Tai-Yin H., Julio U. and Wuhu F.; resources, Tai-Yin H., Julio U., Fabio V.,
492 and Wuhu F.; data curation, Yanlin L.; writing---original draft preparation, Yanlin L.; writing---review and

editing, Yanlin L., Tai-Yin H., Julio U., Fabio V., and Wuhu F.; visualization, Yanlin L.; supervision, Julio U. and Tai-Yin H.; project administration, Julio U. and Tai-Yin H.; funding acquisition, Julio U. and Tai-Yin H. All authors have read and agreed to the published version of the manuscript.

Competing interests

The authors declare no competing interests.

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