



# A two-dimensional volatility basis set – Part 3: Prognostic modeling and NO<sub>x</sub> dependence

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**Abstract.** When NO<sub>x</sub> is introduced to organic emissions, aerosol production is sometimes, but not always, reduced. Under certain conditions, these interactions will instead increase aerosol concentrations. We expanded the two-dimensional volatility basis set (2D-VBS) to include the effects of NO<sub>x</sub> on aerosol formation. This includes the formation of organonitrates, where the addition of a nitrate group contributes to a decrease of 2.5 orders of magnitude in volatility. With this refinement, we model outputs from experimental results, such as the atomic N : C ratio, organonitrate mass, and nitrate fragments in Aerosol Mass Spectrometer (AMS) measurements. We also discuss the mathematical methods underlying the implementation of the 2D-VBS and provide the complete code in the Supplement. A developer version is available on Bitbucket, an online community repository.

known as cloud condensation nuclei (CCN) (Seinfeld and Pandis, 2006). In order to track the production of aerosols, models predict the interactions between emissions from various sources and their effects on the formation of PM.

PM<sub>2.5</sub> consists of a rich mixture including both inorganic and organic compounds, commonly found within individual particles. Inorganics include sodium chloride (from sea spray); sulfate (mostly from coal combustion); nitrate (mostly from high-temperature combustion); ammonium (from animal husbandry); elemental carbon (from combustion of organic fuels); and trace metals such as nickel and manganese. Organics comprise 20–50 % of PM<sub>2.5</sub> mass in the continental midlatitudes and as much as 90 % in tropical forests (Andreae and Crutzen, 1997; Kanakidou et al., 2005). The organics are called organic aerosol (OA), and, while the inorganics consist of a relatively small set of compounds, OA consists of a rich mixture containing many thousands of different individual organic compounds (Goldstein and Galbally, 2007; Kroll et al., 2011).

OA comes from biogenic (naturally occurring) and anthropogenic (human-related) sources. Biomass burning (BBOA) is the largest contributor to OA worldwide (Bond et al., 2004). Some BBOA comes from natural carbon (woodsmoke), but the combustion is largely associated with human activity, and so BBOA is classified as anthropogenic. Overall, OA comprises roughly 10 % of the total flux of organic carbon into (or out of) the atmosphere (Goldstein and Galbally, 2007; Hallquist et al., 2009); thus only a fraction of organic compounds have the right properties to reside in the condensed phase. The requisite property is a low volatility (Pankow, 1994; Donahue et al., 2011, 2014); to stay in the organic phase under typical conditions, an organic molecule

## 1 Introduction

Aerosols, or particulate matter (PM), cause numerous cardiovascular and respiratory diseases, and chronic exposure to high concentrations can significantly reduce life expectancy (Dockery et al., 1993; Peng et al., 2005; Pope et al., 2009). Particles smaller than 2.5 micrometers (PM<sub>2.5</sub>) directly contribute to respiratory- and cardiovascular-related deaths. This is most concerning in major cities where air pollution can reach extremely hazardous levels. In addition, PM affects ecosystems and the atmosphere, exerting significant direct and indirect forces on climate. Direct forcing comes from the scattering or absorption of solar radiation by aerosols. Indirect forcing comes from the scattering of solar radiation by clouds, which is in turn controlled by hydrophilic particles

must have a vapor pressure lower than roughly  $10^{-10}$  atm ( $10^{-5}$  Pa,  $C^* < 1 \mu\text{g m}^{-3}$ ) (Donahue et al., 2011).

OA is also classified as primary or secondary (Murphy et al., 2014; Cronn et al., 1977; Turpin and Huntzicker, 1995). Primary organic aerosols (POAs) are directly emitted into the atmosphere on particles and are largely from anthropogenic sources such as automobile exhaust and biomass burning. Secondary organic aerosols (SOAs) are formed when chemical reactions cause condensation of organic compounds from the gas phase. Oxidants include ozone, hydroxyl (OH $\cdot$ ) radicals, and nitrate (NO $_3$ ) radicals (Turpin et al., 2000), and oxidation can occur in the gas phase (Pandis et al., 1991) or in the aqueous phase (Turpin et al., 2000). Oxidation can add functional groups to an organic backbone (functionalization), forming products with a lowered volatility; however, oxidation can also lead to C–C bond cleavage (fragmentation), often forming products with an elevated volatility (Kroll et al., 2011; Chacon-Madrid et al., 2010). In addition, association reactions between relatively volatile reaction products can lead to higher-molecular-weight, lower-vapor-pressure products (oligomers) (Kalberer et al., 2004).

Oxidation of an organic compound, even one generation, generally forms many products, each with a different volatility (Atkinson et al., 1997; Aumont et al., 2005; Lim and Ziemann, 2009). Thus SOA production has been described in terms of a volatility distribution of reaction products (Odum et al., 1996; Presto and Donahue, 2006; Donahue et al., 2012a). However, this also means that the oxidation products (and thus volatility distribution and overall SOA mass yields) often depend strongly on ambient conditions. The dominant oxidant can have a large effect (OH vs. ozone during the day, NO $_3$  vs. ozone at night), but so can the organic radical chemistry following the initial oxidation step.

One notable source of variability in SOA mass yields is the NO $_x$  level (Presto and Donahue, 2006; Zhang et al., 2006; Ng et al., 2007; Logan et al., 1981; Thompson and Cicerone, 1982). NO $_x$  (NO and NO $_2$ ) can react with organic radicals to change the product distribution from oxidation reactions (Atkinson et al., 1997). NO $_x$  is most commonly formed under high-temperature conditions such as combustion; NO $_x$  concentrations range from 10–1000 ppb in urban areas to 10 ppt in remote regions (Seinfeld and Pandis, 2006), and so NO $_x$  is one of the most highly variable reactive trace species in the atmosphere and also one of the most potent indications of human activity. Because NO $_x$  can alter the oxidation chemistry of organic compounds, even biogenic organic precursors (e.g., isoprene and terpenes) may in fact form “anthropogenically enhanced” SOA if the SOA mass yields are in some way enhanced by the presence of NO $_x$ .

There is evidence that a large source of SOA associated with CO emissions (and thus presumably anthropogenic activity) is required to explain global surface OA observations (de Gouw and Jimenez, 2009; Spracklen et al., 2011). At the same time, a large fraction of OA appears to consist of modern carbon (containing  $^{14}\text{C}$ ) (Weber et al., 2007; Aiken et al.,

2010; Minguillón et al., 2011; Zhang et al., 2012). This has led to speculation that interactions between biogenic precursors and urban plumes – possibly NO $_x$  – may be responsible for the anthropogenic enhancement (Shilling et al., 2013). If so, the dramatic decline in NO $_x$  levels in the southeast United States over the past decade (Russell et al., 2012) might be associated with a corresponding decrease in anthropogenically enhanced SOA.

Tracking any NO $_x$  effect is thus extremely important. If changed product distributions (and thus SOA mass yields) were the sole effect of increased NO $_x$ , then the NO $_x$  influence could be dealt with easily by adjusting SOA mass yields based on NO $_x$  levels (Presto and Donahue, 2006; Lane et al., 2008a). However, if there is a reason to track specific products, or if the subsequent, later-generation chemistry of the reaction products also differs substantially under high- and low-NO $_x$  conditions, then simply adjusting mass yields will not suffice. In the case of NO $_x$ , both of these conditions apply: there is strong evidence that the aging chemistry of SOA depends on its composition (Zhang et al., 2006; Henry and Donahue, 2012) and also that organonitrates produced under high-NO $_x$  conditions are independently measured as a diagnostic of ambient aging conditions.

High organonitrate concentrations are typically found in areas with high NO $_x$  levels (Garnes and Allen, 2002; Mylonas et al., 1991). The addition of a nitrate functional group to a carbon backbone reduces volatility by  $\sim 2.5$  orders of magnitude (Pankow and Asher, 2008), potentially contributing to SOA formation as a result. Organonitrates can be identified on filter samples by characteristic absorption features using Fourier transform infrared spectroscopy (FTIR; Russell et al., 2011) and also in bulk mass spectra from the Aerosol Mass Spectrometer (AMS, Aerodyne Inc.) (Zhang et al., 2006; Farmer et al., 2010). Detection in the AMS is challenging because organonitrates fragment almost completely to give NO $^+$  ( $m/z = 30$ ) and NO $_2^+$  ( $m/z = 46$ ). The same fragments arise from (often much more abundant) inorganic nitrate; however, the fragment signal ratio (30:46) can be indicative of organonitrates and is quite distinct from ammonium nitrate (Farmer et al., 2010).

Depending on the carbon chain length, organonitrates can have a gas-phase atmospheric lifetime ranging from days to months, and thus drive long-range NO $_x$  transport into remote marine regions (Atherton, 1989). Higher-carbon-number organonitrates will partition to the condensed phase (Ziemann and Atkinson, 2012). Organonitrates have been found in aerosol samples at numerous sites across North America (Russell et al., 2011). In particular, high concentrations of organonitrates have been observed in California and Texas (Garnes and Allen, 2002; Day et al., 2010). AMS data from photooxidation of  $\alpha$ -pinene, limonene, and longifolene under high-NO $_x$  conditions show mass fragments corresponding to the formation of organonitrates (Zhang et al., 2006; Ng et al., 2007). In some field observations, roughly 10–20 % of the organic aerosol mass is comprised of

organonitrates; however, organonitrates are not always found in urban areas even in high- $\text{NO}_x$  conditions. Field data from Pittsburgh and the US east coast revealed low concentrations of aerosol organonitrates (Wittig et al., 2004; Russell et al., 2011). This may be due to the hydrolysis of organonitrates, where water reacts in the condensed phase with the nitrate group to form an alcohol and nitric acid (inorganic nitrate) (Liu et al., 2012). The change in the functional group likely has little effect on the volatility of the compound; the -OH functional group replacing the - $\text{ONO}_2$  group has about the same effect on volatility (Pankow and Asher, 2008).

During daytime, organonitrates are formed through reactions of NO with organic peroxy radicals ( $\text{RO}_2\cdot$ ). As  $\text{NO}_x$  concentrations rise, progressively more  $\text{RO}_2\cdot$  react with NO instead of  $\text{RO}_2\cdot$  or  $\text{HO}_2\cdot$ . The reaction with NO produces an intermediate product, ROONO, that can either decompose into an alkoxy radical ( $\text{RO}\cdot$  and  $\text{NO}_2$ ) or isomerize to form an organonitrate ( $\text{RONO}_2$ ) (Atkinson et al., 1997; Zhang et al., 2004). The alkoxy radicals will further react, either forming stable organics that are more functionalized or fragmenting to produce lower carbon-number products. Organonitrate yields from ROONO rise with carbon number and decreasing temperature and vary somewhat with structure, reaching an asymptotic limit of roughly 0.3 at 300 K for most compounds relevant to SOA formation (Atkinson and Arey, 2003; Ziemann and Atkinson, 2012). Alkyl nitrates for species with > 20 carbons are predominantly in the particle phase (Lim and Ziemann, 2009). High- $\text{NO}_x$  products differ from low- $\text{NO}_x$  products; this can lead to comparatively higher or lower SOA concentrations (Presto et al., 2005; Kroll et al., 2006; Ng et al., 2007).

During nighttime, organonitrates are formed through reactions of  $\text{NO}_3$  with unsaturated organic precursors (Crowley et al., 2011; Rollins et al., 2009). This reaction can produce organonitrates on the same magnitude as daytime organonitrate formation (Fry et al., 2013). However, the concentration profile of  $\text{NO}_3$  is not well understood and can vary widely depending on boundary layer conditions (He et al., 2014; Fry et al., 2013). Daytime  $\text{NO}_3$  rapidly undergoes photolysis or reacts with NO to form  $\text{NO}_2$  and is thus typically unable to form organonitrates in the presence of light. Product distributions for  $\text{NO}_3 + \text{VOC}$  reactions have not been developed for the 2D-VBS, but its contribution to organonitrate production can eventually be incorporated within the current framework.

## 2 Background

### 2.1 Volatility basis set

The complexity of OA partitioning and the vast number of reaction products motivated the volatility basis set (VBS). The original (one-dimensional) VBS was designed to represent partitioning and aging by lumping organic molecules only by volatility, in bins separated by an order of magnitude

(at 300 K) in a logarithmic volatility space (Donahue et al., 2006). In principle the 1D-VBS can be implemented with as few as four transportable species (a single four-bin VBS with  $C^* = [1, 10, 100, 1000] \mu\text{g m}^{-3}$ ) at the cost of losing all information about the source of the SOA. SOA models that retain information (e.g., biogenic vs. anthropogenic SOA) have correspondingly more transportable species; Lane et al. (2008a) retained four VBS bins for each SOA precursor source. Other 1D-VBS implementations have used a wider volatility range, for example transporting seven species with  $0.01 \leq C^* \leq 10\,000$  for SOA formed via oxidation of evaporated primary emissions (Shrivastava et al., 2008).

Aging schemes within the 1D-VBS employ a single coupling matrix (Donahue et al., 2006) that can vary with ambient conditions to represent changes in chemistry (Lane et al., 2008a). However, individual VBS bins can include such different species as  $\text{C}_{23}\text{H}_{48}$  (tricosane) and  $\text{C}_6\text{H}_{10}\text{O}_5$  (levoglucosan). There is good reason to believe that the chemistry of these species is quite different – most notably, fragmentation is likely to become more important with increasing oxygenation (Kroll et al., 2009, 2011). Furthermore, the degree of oxidation is an important diagnostic of ambient (Zhang et al., 2007; Jimenez et al., 2009) and chamber (Shilling et al., 2009; Chen et al., 2011) OA measurements. While fragmentation is described in the original 1D-VBS implementation (Donahue et al., 2006), the oxygen content is not tracked, cannot be used as a diagnostic, and cannot inform the aging chemistry.

The 2D-VBS adds a second dimension of oxygenation ( $\text{O}:\text{C}$ ) or average carbon oxidation state ( $\overline{\text{OS}}_{\text{C}} = 2\text{O}:\text{C} - \text{H}:\text{C}$ ) (Donahue et al., 2011; Donahue et al., 2012a). It was developed to capture these important defining properties of OA and track and predict changes over time.

The volatility and  $\overline{\text{OS}}_{\text{C}}$  of organic material are represented explicitly and evolve with aging chemistry, enabling explicit description of the production and evolution of SOA over time. In addition, rate constants for reactions with OH radicals are assigned based on typical chemical structures throughout the two-dimensional space (Donahue et al., 2013). While adding complexity, this expanded representation enables much more systematic exploration of different aging mechanisms in both box models (Donahue et al., 2012a) and Lagrangian chemical transport models (Murphy et al., 2012). It does not, however, enable explicit tracking of product categories such as organonitrates.

Here we describe a refinement to the 2D-VBS by introducing layers to hold key product classes – organonitrates in this case. This allows us to more accurately account for the effect of  $\text{NO}_x$  on the production and evolution of SOA and to capture a more complete picture of organic aging by incorporating the production and removal of organonitrates. The model replicates experimental data on the effect of  $\text{NO}_x$ , where high- $\text{NO}_x$  conditions increase SOA formation for certain precursors (longifolene) and decrease in oth-

ers ( $\alpha$ -pinene) (Presto et al., 2005; Pathak et al., 2007; Kroll et al., 2006; Ng et al., 2007).

Mathematically, all implementations of the VBS are formally one-dimensional: an array of species is connected via a chemical coupling matrix that describes transformations due to aging reactions. However, the physical properties represented by that array of species can be cast into multiple dimensions – volatility, oxidation state, and now levels of key molecular classes. As part of the Supplement for this paper we have released a full implementation of the VBS, coded in MATLAB, providing full graphical diagnostics, access to the different levels of complexity (1-D, 2-D, layered), and instructions for alteration and expansion (e.g. changing volatility or oxidation state ranges, adding layers). In addition, more recent and updated versions of the code are stored in the on-line repository Bitbucket; instructions are detailed within the Supplement.

## 2.2 Mathematical overview

The 2D-VBS follows the movement of carbon mass throughout the volatility and  $\overline{\text{OS}}_{\text{C}}$  space by modeling the basic aging processes of functionalization and fragmentation. Functionalization adds functional groups, producing more oxidized organics that have lower volatility than the parent compound. Fragmentation splits the carbon backbone of an oxidized compound, producing two smaller compounds of higher volatility and generally higher  $\overline{\text{OS}}_{\text{C}}$  (Donahue et al., 2012a). The 2D-VBS implements these aging processes as a series of matrix manipulations.

While the 2D-VBS is typically presented as a two-dimensional space, it is implemented as a one-dimensional concentration vector in the code.

The mass of carbon ( $C_{v,o}$ ) per cubic meter ( $\mu\text{g m}^{-3}$ ) is described in each cell ( $v, o$ ), where  $v$  is the volatility ( $x$ ) index and  $o$  is the oxidation ( $y$ ) index. This can be transformed into a single row vector  $\mathbf{C}$  with  $n$  elements, or cells,  $n = v \times o$ . The transformation method concatenates the rows (the volatilities of each O : C) together:

$$\begin{bmatrix} C_{1,1} & \cdots & C_{x,1} \\ \vdots & \ddots & \vdots \\ C_{1,y} & \cdots & C_{x,y} \end{bmatrix} \quad (1) \\ \rightarrow [C_{1,1} \dots C_{x,1} C_{1,2} \dots C_{x,2} \dots C_{1,y} \dots C_{x,y}].$$

The addition of dimensions, such as organonitrates, is implemented the same way by extending the one-dimensional array to track these compounds.

Chemical reactions can then be represented as transformations from an initial cell  $i$  to all potential final cells  $f$ ; each transformation  $\mathbf{T}$  is a column vector of length  $n$ , with  $\sum_f T_f = 1$  (the transformations conserve carbon). The  $n$  transformation vectors for each initial cell  $i$  can then be concatenated to form an  $n \times n$  transformation matrix,  $\mathbf{T} =$

$[\mathbf{T}_1 \dots \mathbf{T}_n]$ . The overall chemical transformation is then given by  $\mathbf{C}\mathbf{T}$ .

The rate at which components react with OH (or any other oxidant) depends on their composition. Molecules with more carbons typically have more hydrogens available for abstraction reactions. Additionally, the more oxygens a molecule has, the more it is destabilized, and the more likely it will react (Kwok and Atkinson, 1995). However, when enough functional groups become attached to the carbon backbone, the remaining hydrogen atoms are typically less reactive, and so reactivity decreases. A composition–activity relationship from Donahue et al. (2013) accounts for these factors and is used to approximate rate constants gas-phase reactivity for each cell in the VBS. These values form a gas-phase reactivity vector,  $\mathbf{k}^{\text{vap}}$ :

$$\mathbf{k}^{\text{vap}} \simeq 1.2 \times 10^{-12} (n_{\text{C}} + 9n_{\text{O}} - 10(\text{O} : \text{C})^2) \text{cm}^3 \text{molec}^{-1} \text{s}^{-1}, \quad (2)$$

where  $n_{\text{C}}$  and  $n_{\text{O}}$  correspond to the number of carbon and oxygen atoms, respectively.

We also parametrize the chemistry following the initial reaction with OH. The probabilities of fragmentation and functionalization are represented by vectors  $\mathbf{f}^{\text{frag}}$  and  $\mathbf{f}^{\text{func}}$ , respectively, where  $\mathbf{f}^{\text{frag}} = (\text{O} : \text{C})^n$  and  $\mathbf{f}^{\text{func}} = 1 - \mathbf{f}^{\text{frag}}$ . The formula for the potential for fragmentation is determined by the higher likelihood of fragmentation when a molecule is highly oxidized, because of the destabilizing effect of functional groups on the carbon backbone. The exponent,  $n$ , expresses the strength of fragmentation reactions, and is currently estimated to be 1/4 (Donahue et al., 2012b).

Functionalization kernels are created using a distribution based on the degree of oxidation of the compound. The kernel is defined in terms of a probability distribution for added oxygen and decreased volatility (Donahue et al., 2012b). That distribution is then mapped onto the  $C^o$ , O : C space to form a functionalization transformation array. For each volatility cell  $i$ , functionalization ( $\mathbf{T}_i^{\text{func}}$ ) distributes carbon mass throughout the  $n$  VBS cells.

Fragmentation kernels describe the distribution of organic material throughout (mostly) higher-volatility bins and O : C following cleavage of the carbon backbone. Some fragmentation products are molecules and some are radicals, which in turn are rapidly functionalized; the fragmentation kernel describes the ultimate distribution of stable products from this process and thus distributes carbon over a much wider range of the VBS than functionalization (Donahue et al., 2012b). The overall transformation vector  $\mathbf{T}_i$  is found for functionalization and fragmentation by combining the two processes together:  $\mathbf{T}_i = \mathbf{f}_i^{\text{frag}} \cdot \mathbf{T}_i^{\text{frag}} + \mathbf{f}_i^{\text{func}} \cdot \mathbf{T}_i^{\text{func}}$ . Also,  $\sum_{f=1}^n T_{i,f} = 1$ , meaning carbon mass is conserved. Concatenated over all columns,  $i = 1 \dots n$ , the transformations form the transformation matrix  $\mathbf{T}$ . The aging process is then ex-

pressed as

$$\frac{\partial \mathbf{C}}{\partial t} = \mathbf{k}^{\text{vap}} \cdot [\text{OH}] \cdot \mathbf{C} (\mathbf{T} - \mathbf{I}). \quad (3)$$

This can be modeled with differential solvers in MATLAB.

The transformations just described apply to species in the vapor phase. The VBS separately tracks vapor ( $\mathbf{C}^{\text{vap}}$ ) and condensed ( $\mathbf{C}^{\text{cond}}$ ) phase concentrations. For the condensed phase, the OH rate constant is specified as an effective gas-phase rate constant,  $k_i^{\text{eff}}$ , which is determined by the diffusion-limited uptake of gas-phase OH by particles, based on the Fuchs corrected surface area of a given particle size distribution (Donahue et al., 2013). We generally assume this is independent of composition and that OH reacts with unit efficiency once it reaches the surface of a particle. Lacking other constraints, we assume that  $f^{\text{func}}$  and  $f^{\text{frag}}$  are the same in the gas and condensed phases, so  $\mathbf{T}^{\text{vap}} = \mathbf{T}^{\text{cond}}$ ; however, it is likely that even the functionalization and fragmentation kernels differ in the gas and condensed phases, so we anticipate that future refinements will remove this assumption.

Additional details on the distribution of material throughout the VBS may be found in Donahue et al. (2012b) and Murphy et al. (2012).

### 2.3 Treatment of organonitrates

The introduction of  $\text{NO}_x$  to the system brings additional chemical reactions and the potential to form organonitrates through the functionalization process. Organonitrates are accounted for in a separate layer of the 2D-VBS, called the “ $\text{N}_1$ ” layer for the inclusion of an atomic nitrogen. The addition of a nitrate group decreases the volatility by  $\sim 2.5$  orders of magnitude (Pankow and Asher, 2008) and increases the number of oxygens attached to the carbon backbone by 1 (the other oxygen atoms and the nitrogen constitute an  $\text{NO}_2$  group).

This has the potential to produce high concentrations of organonitrates in the condensed phase because of the large decrease in the volatility of the product compared to the reactant.

The formation of an organonitrate involves two branch points. The first is the probability  $(1 - \beta)$  that  $\text{RO}_2$  will react with  $\text{NO}$ ; the second is the yield,  $\eta$ , of organonitrates from  $\text{RO}_2 + \text{NO}$ . The nitrate yield  $\eta$  rises with increasing carbon number (Arey et al., 2001; Yeh and Ziemann, 2014) and depends on the functionality of the  $\text{RO}_2$  (Lim and Ziemann, 2009; Elrod, 2011); however, for organics large enough to partition to the condensed phase,  $\eta$  is at an asymptotic limit, and we assume a homogeneous distribution of  $\text{RO}_2$  structures within any given VBS cell. We thus assume that  $\eta = 0.30$  in all cases. Various ways of estimating  $\beta$  are possible, such as from the  $\text{VOC}:\text{NO}_x$  ratios obtainable in experimental data (Presto et al., 2005) to the comparison of the rates of reaction

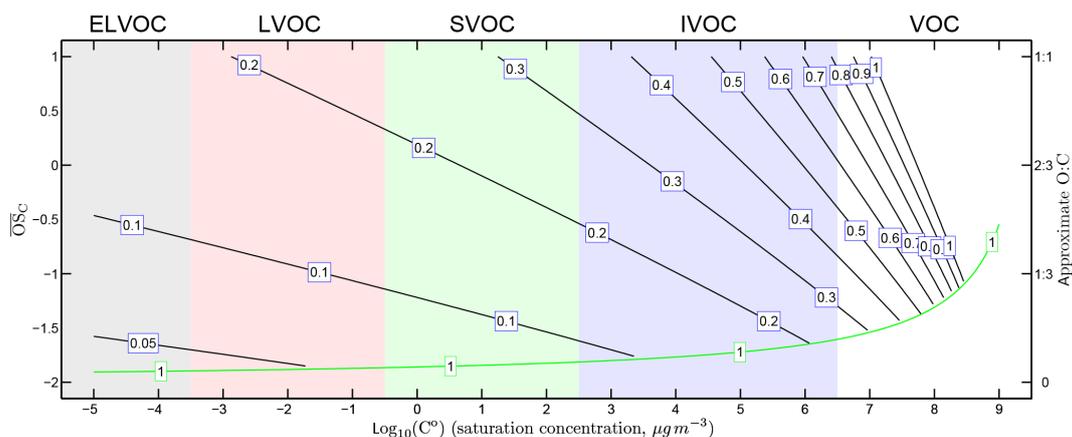
between  $\text{RO}_2 + \text{NO}$  and  $\text{RO}_2 + \text{HO}_2$  (Lane et al., 2008b). In this work, we will simply vary  $\beta$  parametrically.

There are now three different transformation pathways for organics in the  $\text{N}_0$  (non-organonitrate) layer: they can fragment, functionalize without forming organonitrates, or form organonitrates. As a simplification, we assume that there are two pathways for organonitrates in the  $\text{N}_1$  layer: they can fragment by eliminating the  $\text{NO}_2$  group and return to the  $\text{N}_0$  layer, or they can functionalize and remain in the  $\text{N}_1$  layer. Also as a simplification we assume that dinitrates with 2- $\text{ONO}_2$  functional groups are minor products, and so we do not add an  $\text{N}_2$  layer. The transformations for the organonitrate case are analogous to the non-organonitrate case, but with expansions to accommodate the production and loss of organonitrates. The concentration array doubles in size:  $\mathbf{C} = [\mathbf{C}_{\text{N}_0} \mathbf{C}_{\text{N}_1}]$ , where  $\mathbf{C}_{\text{N}_0}$  represents all the cells in the  $\text{N}_0$  layer and  $\mathbf{C}_{\text{N}_1}$  represents the cells in the  $\text{N}_1$  layer. More complex is the transformation matrix, which is now

$$\mathbf{T} = \begin{bmatrix} \mathbf{T}_{\text{N}_0 \rightarrow \text{N}_0} & \mathbf{T}_{\text{N}_0 \rightarrow \text{N}_1} \\ \mathbf{T}_{\text{N}_1 \rightarrow \text{N}_0} & \mathbf{T}_{\text{N}_1 \rightarrow \text{N}_1} \end{bmatrix}, \quad (4)$$

where  $\mathbf{T}_{\text{N}_0 \rightarrow \text{N}_0}$  represents the original functionalization and fragmentation of the non-organonitrate layer.  $\mathbf{T}_{\text{N}_0 \rightarrow \text{N}_1}$  is the formation of an organonitrate through functionalization.  $\mathbf{T}_{\text{N}_1 \rightarrow \text{N}_0}$  is the fragmentation of an organonitrate.  $\mathbf{T}_{\text{N}_1 \rightarrow \text{N}_1}$  is the functionalization of an organonitrate. The organonitrate ( $\text{N}_1$ ) layer of the 2D-VBS contains only one nitrate group, and thus each species contains a single nitrogen atom. Because  $n_{\text{N}}$  is constant,  $\text{N}:\text{C}$  drops as volatility decreases and the chemical species become larger. In addition, the oxygen attached to the carbon from the nitrate group contributes to the  $\text{O}:\text{C}$  of the overall compound. Therefore, the  $\text{O}:\text{C}$  of an organonitrate can never be 0. Figure 1 shows  $\text{N}:\text{C}$  for the nitrate layer ( $\text{N}_1$ ) of the 2D-VBS. In the regime where aerosols tend to form under atmospheric conditions,  $\text{N}:\text{C}$  ranges from 0.04 to 0.3. Semi-volatile organonitrates have  $\text{N}:\text{C}$  from around 0.05 to 0.3; extremely low volatility compounds have an  $\text{N}:\text{C}$  lower than 0.2.

$\text{N}:\text{C}$  for ambient aerosols are experimentally obtainable through atomic mass spectroscopy. Rollins et al. (2010) tested compounds that act as aerosol precursors in the atmosphere using the AMS, and the data from elemental analyses of these organic hydroxynitrates showed  $\text{N}:\text{C}$  ranging from 0.02 to 0.1. Other experiments of SOA production from isoprene, toluene, and naphthalene in high- $\text{NO}_x$  conditions show  $\text{N}:\text{C}$  ranging from 0.04 to 0.08, indicating the existence of low-volatility compounds with moderate levels of oxygenation (Chhabra et al., 2010). In addition to providing a sense of the range of  $\text{N}:\text{C}$  expected in aerosols, Fig. 1 also reveals the relationship between  $\text{N}:\text{C}$  and the size of the molecule itself. Molecules with very small  $\text{N}:\text{C}$  actually indicate high carbon content, meaning these organonitrates are, on a per-molecule basis, contributing significant mass to the overall organic aerosol.



**Figure 1.** Atomic N : C ratio isopleths in the organonitrate layer of the 2D-VBS. The volatility space is divided into regions of organic compounds: extremely low volatility (ELVOC), low volatility (LVOC), semi-volatile (SVOC), intermediate volatility (IVOC), and volatile (VOC). Most organonitrate aerosols will have an N : C ratio lower than 0.2. The green line represents the single oxygen isopleth under which no organonitrate exists, because organonitrates themselves contribute an oxygen to the O : C ratio.

## 2.4 Time evolution

Once the transformation matrix  $\mathbf{T}$  has been specified for any VBS configuration, the time evolution is trivially specified. A final detail is that we separate vapor-phase and particle-phase processes to treat the very different kinetics of homogeneous and heterogeneous oxidation (Donahue et al., 2013). This also facilitates dynamical treatments of aerosol processes (Trump et al., 2014; Trump and Donahue, 2014). We can also treat oligomerization within the condensed phase (Trump and Donahue, 2014), though we shall not address that here. Formally, this again requires that we double the concentration array to distinguish vapor- and condensed-phase concentrations:  $\mathbf{C} = [\mathbf{C}^{\text{vap}} \ \mathbf{C}^{\text{cond}}]$ . The transformation matrix becomes

$$\mathbf{T} = \begin{bmatrix} \mathbf{T}^{\text{vap}} & 0 \\ 0 & \mathbf{T}^{\text{cond}} \end{bmatrix}. \quad (5)$$

This is block diagonal because we do not treat particle microphysics – condensation and evaporation – at this stage. That is left to a later step via operator splitting, whether we assume equilibrium partitioning or specifically treat the dynamics.

The change in the mass within each cell is described by  $n$  differential equations. For each cell  $i$  out of a total of  $n$  cells in the VBS,

$$\frac{d\mathbf{C}}{dt} = C_{\text{OH}} [(\mathbf{k} \cdot \mathbf{C}) \mathbf{T} - (\mathbf{k} \cdot \mathbf{C})] = C_{\text{OH}} (\mathbf{k} \cdot \mathbf{C}) [\mathbf{T} - \mathbf{I}], \quad (6)$$

where the first term is the amount of mass that is reacted into the cell from all other cells, and the second term is the loss from reactions out of the cell.

Any differential equation solver can be employed to solve Eq. (6). We currently use the approximation  $\exp(-k\Delta t) \approx 1 - k\Delta t$  for small  $\Delta t$ .

As functionalized products shift into lower-volatility cells, they can condense into the aerosol phase. At equilibrium this

is given by

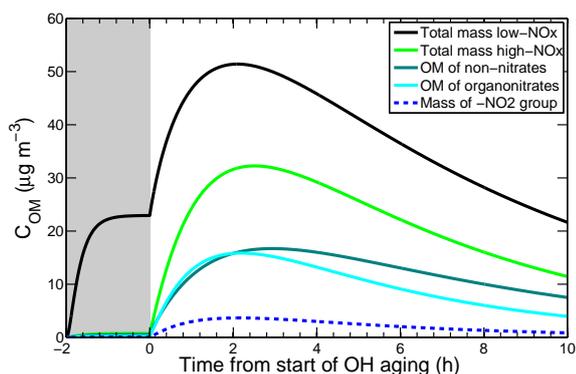
$$\xi_i = \left(1 + \frac{C_i^*}{C_{\text{OA}}}\right)^{-1}; \quad C_{\text{OA}} = \sum_i C_i^* \xi_i, \quad (7)$$

where  $C^*$  is the volatility (in  $\mu\text{g m}^{-3}$ ) of the compound,  $\xi$  is the fraction of organics in the condensed phase, and  $C_{\text{OA}}$  is the total organic aerosol mass. This is a constant balancing act – if more functionalized material with lower volatility forms and more mass condenses, some higher-volatility products will also condense. The reverse is also true – if fragmentation becomes dominant and higher-volatility products form, lower-volatility products may enter the vapor phase.

## 3 Results and discussion

Prior VBS model implementations have treated ozonolysis and subsequent OH multi-generational aging of  $\alpha$ -pinene in chamber studies. A 2D-VBS model reproduced aerosol mass well throughout the course of the Multiple Chamber Aerosol Chemical Aging Study (MUCHACHAS) experiments, which studied SOA aging under similar conditions in four different chambers (Donahue et al., 2012b). Building upon this, the current version (v1.0 in Bitbucket) of the 2D-VBS model can also model  $\alpha$ -pinene aging under high- $\text{NO}_x$  conditions with both ozonolysis aging and OH aging periods. While there are currently few high- $\text{NO}_x$  experimental data addressing multi-generational aging, mass yields of first-generation products from ozonolysis of  $\alpha$ -pinene are available (Pathak et al., 2007; Presto et al., 2005). Simulations for high- $\text{NO}_x$  cases are run based on these data.

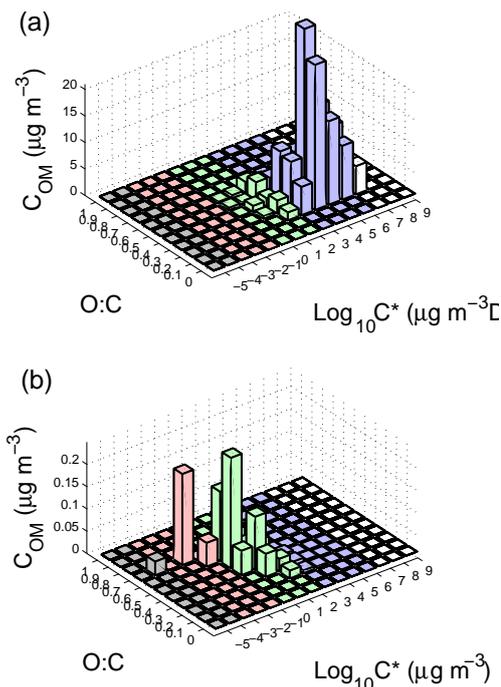
In the examples shown, we run the model under ideal chamber conditions, where there are no wall losses. A concentration of  $100 \mu\text{g m}^{-3}$  of  $\alpha$ -pinene is introduced into the chamber. For the first 2 h, we model dark conditions with a



**Figure 2.** Organic mass concentration over 2 h of dark ozonolysis followed by 10 h of OH aging for low- $\text{NO}_x$  and high- $\text{NO}_x$  conditions. The OM for non-nitrates and organonitrates plots is under high- $\text{NO}_x$  conditions. High- $\text{NO}_x$  conditions decrease the overall organic aerosol mass produced throughout the course of a model run. At  $\beta = 0.15$  (85%  $\text{RO}_2 + \text{NO}$ ), the mass concentration of organonitrates is roughly equivalent to the mass concentration of non-organonitrates. While the contribution of the  $-\text{NO}_2$  group itself is small, the organonitrate compounds can comprise a significant portion of the overall mass.

constant ozone level of 300 ppbv. The ozone reacts with the unsaturated carbon bonds to form first-generation products. At the end of 2 h, after all of the precursor material has been reacted to form first-generation products, OH is introduced into the chamber at a constant rate of  $10^7$  molecules  $\text{cm}^{-3}$  for 10 h. This is the equivalent of turning on the UV lights in a chamber with a strong OH precursor such as HONO. The continued reaction of organic compounds creates multi-generational products, many of which are low volatility and condense into the aerosol phase. We have taken a low- and high- $\text{NO}_x$  case, where  $\beta = 1$  and  $\beta = 0.15$ , respectively, and examined the differences in aerosol production during first-generation chemistry and subsequent multi-generation chemistry.

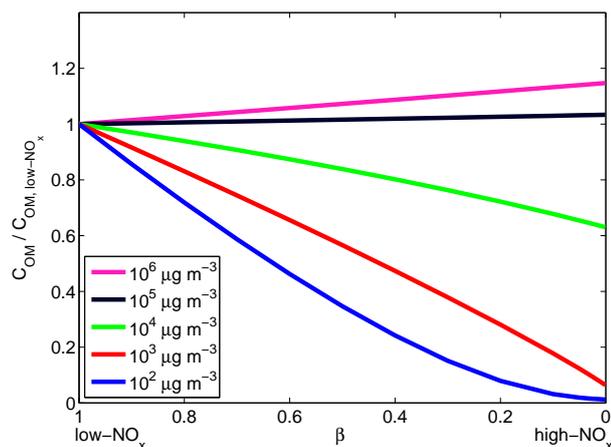
Figure 2 shows the differences in concentration of aerosol mass produced under low- and high- $\text{NO}_x$  conditions throughout the course of the chamber model run. During first-generation chemistry, high- $\text{NO}_x$  conditions form very little aerosol mass, while low- $\text{NO}_x$  conditions produce high SOA mass. This is due to higher-volatility compounds that are generated because of the presence of  $\text{NO}_x$ , instead of the semi-volatile compounds in the absence of  $\text{NO}_x$ . When OH aging begins at low  $\text{NO}_x$ , the aerosol mass more than doubles as volatile compounds react to form highly functionalized, low-volatility species. An even more dramatic effect is seen in the OH aging of high- $\text{NO}_x$  first-generation compounds. While most first-generation products were too volatile to condense, subsequent aging pushed a significant portion of these organics into the aerosol phase. Under these model conditions, even with the dramatic increase in concentration, the high- $\text{NO}_x$  condition produces less overall mass



**Figure 3.** Mass of first-generation products for each cell in the 2D-VBS, from  $100 \mu\text{g m}^{-3}$   $\alpha$ -pinene + ozone at  $\beta = 0.15$ . (a) Total mass of all (organonitrate and non-nitrate) organics in both the suspended and vapor phases. (b) Total mass of all organics in only the suspended phase. The suspended phase concentration axis is 2 orders of magnitude smaller than the vapor phase; most of the first-generation mass remains in the vapor phase.

during OH aging. This alternate pathway results in a significant contribution by organonitrates to the overall aerosol mass – organonitrates comprised nearly half of the total organic mass at one point during OH aging. While the mass of organonitrates is significant, if the AMS were to test this sample, it would measure only a small concentration of N-containing mass fragments (indicated with the dashed blue curve in the figure). In essence, these are very large organonitrate molecules that have significant impact on overall aerosol mass.

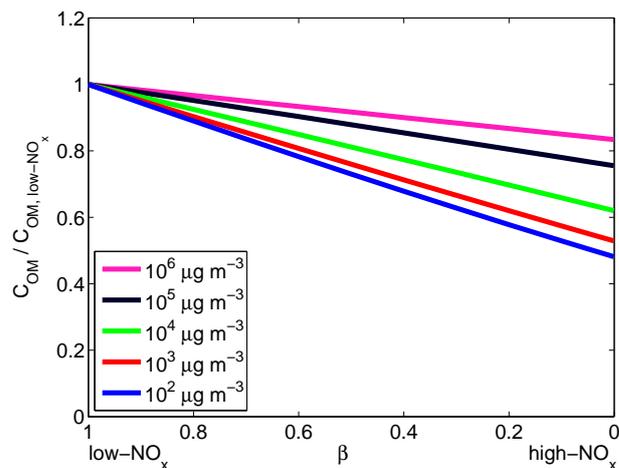
The distribution of mass under high- $\text{NO}_x$  conditions offers insight into the low mass production during first-generation aging. Figure 3 shows the first-generation distribution (at  $t = 0$  of Fig. 2) of organics when  $\beta = 0.15$ . This distribution is based on chamber experiments by Pathak et al. (2007) and Presto et al. (2005) of  $\alpha$ -pinene ozonolysis under high- $\text{NO}_x$  concentrations. Figure 3a shows the distribution of all organics (both aerosol and vapor phase). The majority of the organics reside in the IVOC range, which is too volatile to condense in the typical mass loadings of the atmosphere. Figure 3b shows the distribution of total suspended material, comprised of both organonitrates and non-nitrates. This mass contains more oxidized material than those in the vapor phase and ranges from LVOC to SVOC. Compared to the



**Figure 4.** The effect of peroxy-radical branching on the first-generation secondary organic aerosol concentration,  $C_{OM}$ . The  $x$  axis is the branching ratio for  $RO_2$  reaction ( $\beta$ ), where  $\beta = 1$  corresponds to low- $NO_x$  conditions dominated by  $RO_2 + HO_2$  and  $\beta = 0$  corresponds to high- $NO_x$  conditions dominated by  $RO_2 + NO$ . The  $y$  axis is the ratio of  $C_{OM}$  at a given value of  $\beta$  to the low- $NO_x$  value. Each curve represents a different amount of oxidized precursor, spanning 5 orders of magnitude from  $10^2$  to  $10^6 \mu\text{g m}^{-3}$ .

first-generation distribution in low- $NO_x$  conditions shown in Fig. 2, the high- $NO_x$  condition produces less aerosol mass because of the overall higher volatilities of resulting compounds.

The previous figures have shown that the presence of  $NO_x$  decreases aerosol mass production for  $\alpha$ -pinene under moderate loadings. Less volatile precursors like sesquiterpenes are affected differently by  $NO_x$ . A precursor with a lower volatility than  $\alpha$ -pinene, such as longifolene, produces products with correspondingly lower volatility (Ng et al., 2007). While first-generation distributions for these precursors are currently lacking, we can emulate this effect by increasing the mass loading of  $\alpha$ -pinene instead. A precursor compound that is an order of magnitude less volatile than  $\alpha$ -pinene can be modeled by an order-of-magnitude increase in mass loading. Figure 4 shows a comparison of first-generation aerosol concentration under various  $\beta$  and mass loadings to the mass concentration produced under low  $NO_x$ . This shows that, for lower mass loadings of  $\alpha$ -pinene such as  $100 \mu\text{g m}^{-3}$ , increasing  $NO_x$  levels decreases resultant mass. As the loadings increase in magnitude, the aerosol suppression effect of  $NO_x$  decreases. When loadings reach  $10^5 \mu\text{g m}^{-3}$ , the effect is reversed, instead producing more mass as  $NO_x$  levels increase. This has been shown in experimental data from Ng et al. (2007), where longifolene exhibits higher yields under high- $NO_x$  conditions than low- $NO_x$  conditions. Even though the presence of  $NO_x$  produces first-generation compounds from longifolene that are higher in volatility than first-generation compounds produced under low  $NO_x$ , these

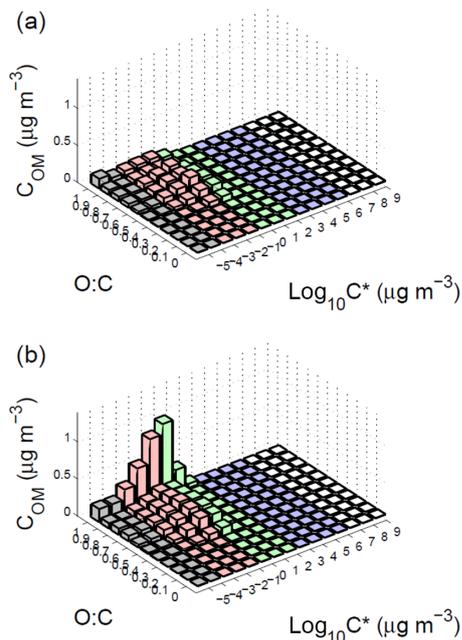


**Figure 5.** The effect of peroxy-radical branching on the aged secondary organic aerosol concentration,  $C_{OM}$ , after 10 h of oxidation by  $10^7 \text{OH cm}^{-3}$ . As in Fig. 4 the  $x$  axis is the branching ratio for  $RO_2$  reaction ( $\beta$ ), where  $\beta = 1$  corresponds to the low- $NO_x$  conditions dominated by  $RO_2 + HO_2$  and  $\beta = 0$  corresponds to high- $NO_x$  conditions dominated by  $RO_2 + NO$ . The  $y$  axis is the ratio of  $C_{OM}$  at a given value of  $\beta$  to the low- $NO_x$  value. Each curve represents a different amount of oxidized precursor, spanning 5 orders of magnitude from  $10^2$  to  $10^6 \mu\text{g m}^{-3}$ .

compounds are still LVOCs and SVOCs, and they partition into the aerosol phase. In addition, the organonitrate group contributes significant mass to the aerosol phase, resulting in the  $> 1$  high- $NO_x$  to low- $NO_x$  organic mass ratio. Depending on the volatility of the organic precursor, it is therefore possible for the presence of  $NO_x$  to increase or decrease the total first-generation aerosol mass.

As these first-generation products undergo multi-generational chemistry, becoming a more complex mix of organics, the stark differences between the mass loadings under first-generation chemistry are dampened. Figure 5 shows the comparison between different  $\beta$  and different initial mass loadings with respect to the low- $NO_x$  ( $\beta = 1$ ) case, after 10 h of OH aging. This corresponds to the end of the time period in Fig. 2. While higher loadings continue to produce more aerosol mass, an increase in  $NO_x$  now consistently produces lower mass. This comes from the tendency to cleave the nitrate group during fragmentation, decreasing the effect of the nitrate group on overall mass. In addition, the probability of fragmentation increases as functionalization continues throughout the course of aging, contributing to the loss of organonitrates and the loss of aerosol mass to the vapor phase.

The distribution of suspended mass throughout the VBS also changes over the course of OH aging. Figure 6 shows the distribution of mass in the 2D-VBS for the organonitrate layer (Fig. 6a) and the non-nitrate (Fig. 6b) after 10 h of OH aging. This corresponds to the end points of the “OM of organonitrates” and “OM of non-nitrates” lines in Fig. 2.



**Figure 6.** The distribution of mass is plotted over the 2D-VBS space for multi-generation products, after 10 h of OH aging of  $100 \mu\text{g m}^{-3}$   $\alpha$ -pinene + ozone at  $\beta = 0.15$ . (a) The 1N layer, where the mass of products is highest in the moderately oxidized and semi-volatile ranges. (b) The 0N layer, where the mass is highly oxidized and lower in volatility.

The organonitrate concentration is fairly small, and they are only moderately oxidized, while the non-nitrate organics are highly oxidized. Each cell of the organonitrate layer, with higher mass yet lower carbon numbers, is slower to react compared to the corresponding cell in the non-nitrate layer. However, as the organonitrates undergo repeated functionalization, fragmentation of the products becomes more preferable, leaving behind less oxidized material in the organonitrate layer. Fragmentation of these highly functionalized organonitrates cleaves the nitrate group, resulting in highly oxidized non-nitrate organics. As a result, the organonitrate layer tends to be less oxidized and semi-volatile, while non-nitrates are highly oxidized.

#### 4 Conclusions

We have added a layer to the 2D-VBS to account for organonitrates. In addition, we have developed a module to treat the formation and aging of organonitrates under high- $\text{NO}_x$  conditions, and we released a complete implementation of this code in MATLAB.

There are relatively few experimental constraints on the behavior of organonitrates in SOA, especially for multi-generational aging. Unknowns include differences between homogeneous and heterogeneous oxidation and the fate of nitrates after fragmentation. As a simplification we have as-

sumed that homogeneous and heterogeneous product distributions are the same and that fragmentation breaks the weakest bond in the organonitrate, assumed to be the O– $\text{NO}_2$  bond. However, it is certain that semi-volatile products (nitrates and non-nitrates alike) will react relatively rapidly in the gas phase and consequently that any first-generation semi-volatile products will not be long lived in the atmosphere.

We explored the aging chemistry of SOA under conditions typical of chamber experiments. During multi-generational aging under high- $\text{NO}_x$  conditions, the contribution of organonitrates to organic mass is similar to non-nitrate organics. This contribution of organonitrates to aerosol mass can be large even though the actual nitrate ( $-\text{ONO}_2$ ) mass is low, because the N : C of condensed-phase organonitrates can be low. We also reproduced the enhancement of aerosol production in the presence of  $\text{NO}_x$ , under conditions such as high mass loadings or low-volatility precursors. This showed that the VBS is capable of accounting for these effects on extensive organic chemistry and emphasized the important role that  $\text{NO}_x$  plays as part of the aging process.

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