

Evaluation and updates to the oxidized reactive nitrogen gaseous dry deposition parameterization from the GEOS-Chem model, including a pathway for ground surface NO₂ hydrolysis

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Abstract. Dry deposition is a major loss pathway for reactive nitrogen species from the atmospheric boundary layer. We
10 evaluate isolated components of the parameterization for species-specific gaseous dry deposition velocity $V_d(x)$ for HNO₃ and
NO₂ from the GEOS-Chem chemical transport model by running a stand-alone version of V_d code in single-point-mode to
enable more direct comparison to field observations. Improved measurement–model agreements result mainly from (i) updates
to the calculation of molecular diffusivities and (ii) representing ground surface NO₂ hydrolysis in the formulation of non-
stomatal uptake. We evaluate the parameterization for non-stomatal dry deposition of NO₂ by comparing to eddy covariance
15 inferred nocturnal $V_d(\text{NO}_2)$ over Harvard Forest. We address a large low bias (-80 %) in simulated nocturnal $V_d(\text{NO}_2)$ by
representing NO₂ heterogeneous hydrolysis on deposition surfaces, paying attention to chemical flux divergence, soil NO
emission, as well as canopy surface area effects. Finally, we evaluate the updated oxidized reactive nitrogen (NO_y) dry
deposition parameterization by comparing to eddy covariance inferred $V_d(\text{NO}_y)$ over Harvard Forest, finding a modest
nocturnal low bias (-19 %) remains in simulated $V_d(\text{NO}_y)$ due to the compensating effects of updates to the calculation of
20 molecular diffusivities (28 % reduction in nocturnal $V_d(\text{NO}_y)$) and representation of NO₂ heterogeneous hydrolysis (25 %
increase in nocturnal $V_d(\text{NO}_y)$). These developments are a first step towards a tractable representation of NO₂ hydrolysis in a
dry deposition scheme and have important implications for near-surface NO₂ lifetime through a mechanism involving HONO
emission.

1 Introduction

25 Chemical species comprising oxidized reactive nitrogen (NO_y) form a main component of atmospheric reactive nitrogen ($\text{N}_r \equiv$
NO_y + reduced nitrogen species) which together play a central role in atmospheric chemistry by modulating the oxidative
capacity of the atmosphere through nitrogen oxides (NO_x $\equiv$ NO + NO₂) (Crutzen, 1979), contributing to nitrogen loading of
natural ecosystems (Clark et al., 2018), and influencing air quality (Fields, 2004). Accurate knowledge of sources and sinks of
N_r is vital for understanding and modeling atmospheric chemistry, including the sensitivity of air quality to changes in

anthropogenic emissions. Dry deposition of N_r from the atmospheric boundary layer is an important removal process, typically contributing between one-third to two-thirds of total (wet + dry) deposition (Flechard et al., 2011; Hanson & Linder, 1991; Munger et al., 1998; Sparks et al., 2008; Walker et al., 2020), but questions remain about its representation in chemical transport models (CTMs).

The atmosphere–surface exchange of N_r may be measured directly via micrometeorological techniques (Businger, 1986; Walker et al., 2020) or under more controlled conditions via enclosure techniques (Breuninger et al., 2012; Hanson and Linder, 1991). Direct measurements of above-canopy air–surface exchange of N_r , including via the eddy covariance technique, are technically complex and resource intensive, resulting in a scarcity of flux observations across representative land types and seasons (Walker et al., 2020). Therefore, studies of above-canopy dry deposition tend to be intensive in nature and are typically designed to characterize exchange processes rather than to monitor long-term deposition patterns. Dry deposition budgets thus fall to the realm of inferential methods, where deposition fluxes F_x are inferred from parameterizations of above-canopy deposition velocity V_d —a first-order rate coefficient for heterogeneous surface reaction/uptake for a specific gas x to a specific bulk surface/land type from a specified height:

$$F_x = -V_d(x) [x], \quad (1)$$

By convention, downward fluxes toward the surface are negative values represented by positive deposition velocities. N_r component concentrations $[x]$ from which dry deposition budgets may be inferred have been obtained from (i) surface networks such as U.S. CASTNET (Clarke et al., 1997) and Canadian CAPMoN (Zhang et al., 2009), (ii) chemical transport models (Dennis et al., 2013; Zhang et al., 2012; Zhang et al., 2018), and (iii) satellite observations (Geddes & Martin, 2017; Kharol et al., 2018; Nowlan et al., 2014).

Deposition velocity represents a bulk quantity with contributions from complex processes including turbulent and molecular diffusion in air, meteorological influence on the physical, chemical, and biological state of surfaces, and species-specific interfacial chemistry. The most common parameterization of V_d in large-scale CTMs considers the deposition pathway as a series of three resistances (Baldocchi et al., 1987; Wesely and Hicks, 1977):

$$V_d(z, x) = \frac{1}{R_a(z) + R_b(x) + R_c(x)}, \quad (2)$$

where for bulk-canopy V_d above a projected ground area, $R_a(z)$ is the aerodynamic resistance to turbulent transport from a specified height z and is common for all species, $R_b(x)$ is the species-specific quasi-laminar boundary layer resistance to transport through the thin non-turbulent layer in direct contact with surfaces, and $R_c(x)$ is the bulk-canopy surface resistance for a specific land type. Expressions for R_a and R_b can be obtained from micrometeorological flux–gradient relationships (Garratt, 1992; Wesely and Hicks, 1977) and vary as a function of surface roughness, wind speed, diabatic stability, and molecular diffusivity in air. For highly soluble species such as HNO_3 and H_2O_2 , contributions from R_c are nominally small with resulting deposition varying between R_a - and R_b -limited depending on the state of turbulence (Nguyen et al., 2015). For species with low aqueous solubility or limited interfacial reactivity, R_c is the limiting term, except under very stable conditions

(Toyota et al., 2016). Given the complexity and variability of canopy types and species-specific surface reactivities, R_c is difficult to treat theoretically, with parameterizations relying heavily on empirical formulations.

The most common parameterization of R_c used by large-scale atmospheric models, including the widely utilized WRF-Chem and GEOS-Chem CTMs, is the Wesely 1989 algorithm (Wesely, 1989; hereafter referred to as W89), or modifications thereof (Hardacre et al., 2015). In this scheme, the bulk-canopy is treated as a single uniform surface or ‘big-leaf’ with stomatal and various non-stomatal deposition pathways acting in parallel. Trace gas specific component surface resistances are calculated following basic similarity relations, including solubility relative to SO_2 and oxidative potential relative to O_3 . Zhang et al. (2003a) present a parameterization of $R_c(x)$ for use in air quality models, including at the global scale, using similarity arguments to SO_2 and O_3 as done in W89, with updates including online computation of within canopy aerodynamic resistance, influence of leaf water vapour pressure deficit and water stress on stomatal resistance, and updated parameterizations of non-stomatal surface resistances for O_3 (Zhang et al., 2002b) and SO_2 (Zhang et al., 2003b). Zhang et al. (2003a) note that application of the algorithm (hereafter referred to as Z03) to compounds for which little to no deposition flux observations exist will continue to be a source of uncertainty. Bulk-canopy surface resistances deviating from W89 similarity to SO_2 and O_3 have been observed for NO_2 (Eugster and Hesterberg, 1996; Horii et al., 2004; Stocker et al., 1995), PAN (Shepson et al., 1992; Sun et al., 2016; Turnipseed et al., 2006), and many other species (Nguyen et al., 2015). Wu et al. (2012) compare observed $V_d(\text{PAN})$ over a coniferous forest to deposition velocities parameterized according to both the W89 (WRF-Chem) and Z03 (NOAH) schemes and find underestimates greater than a factor of 2, motivating effort to fit non-stomatal $R_c(\text{PAN})$ directly from above-canopy flux observations. Using the eddy covariance flux dataset from Nguyen et al. (2015), the Z03 scheme was extended by Wu et al. (2021) to additional species by fitting non-stomatal uptake of oxidized VOCs and hydrogen cyanide directly from observations. Wu et al. (2021) maintain the Z03 algorithm structure through similarity to SO_2 and O_3 ; however, they suggest that future developments to dry deposition schemes consider other species-specific processes and reactions affecting measured uptake, including below-sensor chemical flux divergence, enzymatic reactions, and other non-stomatal processes/reactions.

A main result of Horii et al. (2004) in their analysis of an extensive eddy covariance flux dataset of NO_2 over a Northeastern U.S. mixed forest (Harvard Forest) from April–November was that a persistent deposition process was active at night, yielding NO_2 deposition velocities on average of $\sim 0.2 \text{ cm s}^{-1}$, with values up to 0.5 cm s^{-1} noted under high NO_2 loads of $\sim 30 \text{ ppb}$. This observation is contrary to the widely used W89 parameterization which does not allow significant surface uptake of NO_2 at night when leaf stomata are assumed closed or during vegetatively dormant seasons. Geddes et al. (2014) monitored eddy covariance fluxes of NO , NO_2 , and NO_y above midlatitude ($\sim 45^\circ \text{ N}$) summertime mixed hardwood forests in Ontario (Canada) and Michigan (U.S.), finding on average NO_x fluxes indistinguishable from zero for these relatively low NO_y environments ($< 2 \text{ ppb}$ on average). However, infrequent nocturnal events with high NO_x/NO_y ratios and large downward NO_y fluxes could be interpreted as yielding NO_2 deposition velocities similar to the average values of Horii et al. (2004). Geddes et al. (2014) were careful to note that above-canopy fluxes of NO_x are influenced not only by deposition processes but also by within canopy emissions and chemistry, resulting in above-canopy fluxes of NO_x that are confounded by a combination

of counteracting mechanisms which render flux observations difficult to interpret. Horii et al. (2004) considered below-sensor chemical flux divergence of NO₂, due to formation and subsequent hydrolysis of N₂O₅, where the maximum rate of loss was insufficient to account for the observed downward nocturnal NO₂ flux. To reconcile this they proposed a non-stomatal hydrolysis pathway for uptake of NO₂ on ground and canopy surfaces—a reaction which has been suspected in the field to be of atmospheric relevance for some time (Harrison & Kitto, 1994; Harrison et al., 1996).

The hydrolysis of NO₂ on hydrated surfaces is a well-known heterogeneous reaction from lab investigations yielding adsorbed HNO₃ and evolved nitrous acid (HONO):



Despite the stoichiometry of reaction R1, first-order kinetics in NO₂ have generally been observed in the lab and field, with a rate having dependence on surface area density (as expected for collision-limited heterogeneous catalysis), surface water content, and other surface chemical properties (Finlayson-Pitts et al., 2003; Finlayson-Pitts, 2009; Lammel, 1999; Spataro and Ianniello, 2014). In addition to hydrated ground (Kurtenbach et al., 2001; Lammel, 1999; Ren et al., 2020; VandenBoer et al., 2013) and aerosol (Bröske et al., 2003; Burkholder et al., 2015; Crowley et al., 2010; Tan et al., 2016) surfaces, reaction R1 has been implicated on the sea surface (Wojtal et al., 2011; Yang et al., 2021; Zha et al., 2014), on snow and ice surfaces (Beine et al., 2001; Kim and Kang, 2010), as well as on indoor surfaces (Collins et al., 2018; Febo and Perrino, 1991; Spicer et al., 1993). Spicer et al. (1993) and Collins et al. (2018) both found an indoor lifetime of NO₂ to reactive loss (HONO producing) on residential interior surfaces on the order of one hour in well-mixed air—lower than typical ambient NO₂ chemical lifetimes on the order of hours in regional (Kenagy et al., 2018; Shah et al., 2020) or urban outflows (Laughner and Cohen, 2019) and remote forest environments (Browne and Cohen, 2012). Reaction R1 may be an especially important surface removal process during summertime nights or winter months when NO₂ is longer lived with lifetimes on the order of 10 h to more than a day (Browne and Cohen, 2012; Kenagy et al., 2018; Martin et al., 2003). Reaction R1 has also been implicated in the uptake of NO₂ through leaf stomata, where it may be an important contributor to NO₂ deposition within the moist and high surface area substomatal cavities (apoplast) of leaves (Ammann et al., 1995).

Despite the evidence for reaction R1 proceeding on nearly any environmental surface with adsorbed water, regional and global CTMs have yet, to our knowledge, to update dry deposition parameterizations of NO₂ to include this effect, potentially underestimating and/or misrepresenting $V_d(\text{NO}_2)$ at night and throughout vegetatively senescent periods when stomatal uptake would be weak or absent. In this study, we compare simulated dry deposition velocities from the GEOS-Chem CTM to above-canopy observations of $V_d(\text{NO}_2)$ and $V_d(\text{NO}_y)$ inferred from an extensive publicly available dataset of NO₂ and NO_y eddy covariance fluxes and speciated NO_y concentration measurements over Harvard Forest (Munger and Wofsy, 2023), paying attention to soil NO emission, chemical flux divergence, and canopy surface area effects. Prior to updating simulated $R_c(\text{NO}_2)$ to include reaction R1, we conduct sensitivity tests to evaluate the parameterization of R_a and R_b used in GEOS-Chem by comparing to daytime deposition velocities of rapidly depositing species inferred by the method of eddy covariance over a Southern U.S. temperate forest (Nguyen et al., 2015). Specifically, we comment on the effects that site-specific roughness

length, reference height, and the roughness sublayer have on the simulation of daytime R_a , followed by correction of a positive bias in calculated molecular diffusivities which greatly improves the simulation of daytime $V_d(HNO_3)$ via a large relative increase in R_b .

2 Reference Model and Measurements

2.1 Reference algorithms for computing gaseous dry deposition velocities

2.1.1 GEOS-Chem dry deposition module

To facilitate site-specific comparisons to measured deposition velocities, we use a stand-alone version of the gaseous dry deposition algorithm from GEOS-Chem v10-01 (www.geos-chem.org), implemented to run in single-point-mode with the option to use on-site meteorology and canopy characterizations (i.e., LAI, canopy height, and land type classification). Section S1 of the supplement details the formulations used in GEOS-Chem for the resistance-in-series components of Eq. (2), which were also detailed in recent literature (Wong et al., 2019). Briefly, aerodynamic resistance R_a to the turbulent transport of scalars from a reference height z down to the roughness length z_o of the surface is computed following standard surface layer flux-gradient relationships (Wesely and Hicks, 1977). The quasi-laminar boundary layer resistance R_b is estimated following the semi-empirical formulation from Wesely and Hicks (1977) and has dependency on both friction velocity u_* and species-specific molecular diffusivity D_x . Surface resistance R_c is computed following a big-leaf scheme based on the W89 algorithm, modified for application to the global scale (Wang et al., 1998).

2.1.2 Non-stomatal branch of Z03 dry deposition algorithm

The Z03 dry deposition algorithm includes several updates over the W89 scheme. Z03 is used in the Canadian Air and Precipitation Monitoring Network (CAPMoN) (Zhang et al., 2009) as well as in air quality models (e.g., Zhang et al. (2002a)), and was recently compared globally to the W89 scheme as implemented in GEOS-Chem for $V_d(O_3)$ (Wong et al., 2019). We implement a stand-alone version of the non-stomatal branch of the Z03 dry deposition algorithm for NO_2 to enable evaluation against nocturnal eddy covariance inferred $V_d(NO_2)$ at Harvard Forest. Leaf stomata are treated as fully closed at night in the Z03 scheme; accordingly, we treat stomatal resistance as infinite. The Z03 algorithm assigns scale factors (their α and β) for non-stomatal NO_2 uptake relative to inverse surface resistances (conductances) for SO_2 and O_3 of 0 and 0.8, respectively, resulting in parameterized NO_2 deposition velocities 10–20 % smaller than for O_3 (Zhang et al., 2002a). The canopy is flagged as wet from dew following the formalism adopted by Brook et al. (1999), with dependence on cloud fraction, temperature, dew point, and u_* . We estimate snow cover fraction from snow depth following Zhang et al. (2003a). Following a similar approach to Wu et al. (2018), we compute component surface resistances for the mixed forest as an average of deciduous broadleaf and evergreen needleleaf land type specific values from Z03, weighted by LAI-determined deciduous and coniferous

fractions for Harvard Forest of $\sim 80\%$ and 20% , respectively (Fig. S5 in the supplement). As with the stand-alone GEOS-Chem dry deposition algorithm, we use on-site meteorology and canopy characterizations, where available (Section 2.2.2).

160 2.2 Above-canopy dry deposition velocities inferred from eddy covariance measurements

We evaluate the gaseous dry deposition scheme from GEOS-Chem against eddy covariance inferred deposition velocities over two temperate forests in the U.S. First, we compare to deposition velocities from Nguyen et al. (2015) for species found to dry deposit with minimal surface resistance. Being able to neglect the complexities of a surface resistance scheme allows for a more direct evaluation of R_a and R_b components of the resistance-in-series pathway used in the parameterization of V_d (Wu et al., 2021). Second, for an in-depth evaluation of simulated $V_d(\text{NO}_2)$ and $V_d(\text{NO}_y)$, we use a publicly available long-term hourly dataset of eddy covariance flux observations of NO_2 and NO_y from Harvard Forest, supported with ancillary measurements including NO_y component concentrations, meteorological observations, and canopy characteristics.

2.2.1 Talladega National Forest: H_2O_2 , HMHP, and HNO_3

Nguyen et al. (2015) present a novel dataset containing eddy covariance inferred deposition velocities of 16 gaseous species, including species found to deposit with negligible surface resistance: H_2O_2 , hydroxy methylhydroperoxide (HMHP), and HNO_3 . Observations were taken at the Centreville (CTR) Southeastern Aerosol Research and Characterization Study (SEARCH) site (32.90289° N , 87.24968° W) near Brant, Alabama, U.S. in June 2013. The CTR site is situated in a grassy clearing in the Talladega National Forest—a mixed forest consisting of coniferous and deciduous tree species with a mean canopy height h_c of $\sim 10 \text{ m}$ and LAI of $4.7 \text{ m}^2 \text{ m}^{-2}$. Eddy covariance flux observations were measured at 22 m AGL . The analysis of Nguyen et al. (2015) includes daytime mean deposition velocities averaged between 10–15 Local Solar Time (LST) across five ideal days in June 2013 when winds had exclusively forest fetch. To compare with the reported daytime deposition velocities for H_2O_2 ($5.2 \pm 1.1 \text{ cm s}^{-1}$), HMHP ($4.1 \pm 1.1 \text{ cm s}^{-1}$), and HNO_3 ($3.8 \pm 1.3 \text{ cm s}^{-1}$), we average R_a and R_b components of the stand-alone GEOS-Chem dry deposition algorithm, applied at the location of the CTR site, between 10–15 LST on the aforementioned days. Meteorological inputs required to compute R_a and R_b components of the algorithm (u_* , T , P , and sensible heat flux) were obtained from NASA’s Goddard Earth Observing System (GEOS) Forward Processed (FP) assimilated meteorological fields (Lucchesi, 2013) at the native horizontal resolution of $1/4^\circ \times 5/16^\circ$, which Nguyen et al. (2015) note are in excellent agreement with values measured at the CTR site during this period.

2.2.2 Harvard Forest: NO_2 and NO_y

The utility of the Harvard Forest Environmental Monitoring Site (HFEMS) for evaluating parameterizations of atmosphere–surface exchange stems from the extensive datasets of meteorological and trace gas observations spanning many months to years at high temporal (hourly) resolution. The HFEMS is located in central Massachusetts, U.S. (42.54° N , 72.18° W ; 340 m ASL) and situated in a mature mixed forest ($h_c \sim 20 \text{ m}$) with a summertime LAI and deciduous LAI (DLAI) of 4.3 and 3.4, respectively (Fig. S5). Due to prevailing westerly winds, emissions from Boston (100 km to the east) rarely influence the site.

Cool, dry, and unpolluted air from the northwest and warm, moist, anthropogenically influenced air from the southwest are the predominant influences at this site (Horii et al., 2005).

Munger et al. (1996) have described the methodology of the long-term above-canopy (29 m) total nitrogen oxide (NO_y) concentration measurements for eddy covariance flux computation, as well as other details of the HFEMS. Measurements of above-canopy PAN concentrations were added in April 2000 (Horii et al., 2005). Eddy covariance fluxes of NO_2 along with above-canopy (22 m) measurements of HNO_3 concentrations were made at the HFEMS from April through November 2000 (Horii et al., 2004).

Trace gas data from the HFEMS used in this study, specifically, hourly eddy covariance fluxes of NO_y and NO_2 , and hourly concentrations of NO_y , NO , NO_2 , PAN, and HNO_3 are publicly available from the Harvard Forest Data Archive (Munger and Wofsy, 2004, 2023). Exchange velocities (V_{ex}) are computed herein by normalizing reported hourly NO_y and NO_2 eddy covariance fluxes by respective ambient hourly concentrations. Equating V_{ex} to V_d assumes that the observed flux is due to surface deposition only. Processes causing deviation from this assumption are discussed in later sections and include surface emission of NO , chemical flux divergence of NO_2 , and a potential non-zero canopy accumulation rate of NO_y . Eddy covariance fluxes have less error under conditions where turbulence is well developed (Baldocchi, 2003; Cherin et al., 2015; Goulden et al., 1996; Nguyen et al., 2015). Turbulent threshold u_* values in the range $0.15\text{--}0.35\text{ m s}^{-1}$ (median 0.23 m s^{-1}) have been found to be representative of multiple sites across many years (Cherin et al., 2015). Herein, following the approach of Wu et al. (2011), periods of low surface layer turbulence ($u_* < 0.2\text{ m s}^{-1}$) have been omitted from analysis, resulting in $\sim 25\%$ of hourly values of nocturnal $V_d(\text{NO}_2)$ and 18% of hourly values of $V_d(\text{NO}_y)$ being removed from the dataset. Outliers in the remaining hourly $V_d(\text{NO}_2)$ and $V_d(\text{NO}_y)$ timeseries were identified via the method of median absolute deviation (MAD) (Leys et al., 2013), where hourly values outside of the median $\pm 3\times$ MAD were removed from calculations of subsequent means; $\sim 20\%$ of hourly nocturnal $V_d(\text{NO}_2)$ and 10% of hourly $V_d(\text{NO}_y)$ were removed from the u_* filtered dataset. Overall, 60% of the nocturnal $V_d(\text{NO}_2)$ and 74% of the $V_d(\text{NO}_y)$ hourly timeseries were retained for analysis after application of these turbulence and outlier filters. Figure S6 in the supplement depicts monthly fractional coverage of hourly measurements of above-canopy trace gas concentrations and eddy covariance observed exchange velocities from 2000–2002, filtered for conditions of low turbulence.

Meteorological input variables required in the parameterization of V_d were taken from the HFEMS data archive; specifically, P , T , RH , u_* , and sensible heat flux (Munger and Wofsy, 2024) and incoming solar radiation (Fitzjarrald and Sakai, 2023) were available at hourly temporal resolution throughout the study period. Cloud fraction and snow depth were the only required meteorological variables not available and were instead taken from NASA’s Modern-Era Retrospective analysis for Research and Applications version 2 (MERRA-2) assimilated meteorological fields (Gelaro et al., 2017). Figure S7 in the supplement depicts comparisons of hourly observations of u_* , sensible heat flux, downward shortwave radiation, T , P , and RH made over Harvard Forest to coincident values from MERRA-2 assimilated meteorology, depicting good to excellent agreement. Canopy-specific inputs to the parameterization of V_d include roughness length z_o , displacement height d , LAI, and deposition land type. Values for z_o and d were estimated as $1/10^{\text{th}}$ and $2/3^{\text{rd}}$ of canopy height, respectively—values

representative of many vegetative surfaces (Garratt, 1992; Oke, 1987), including z_o for Harvard Forest (Wu et al., 2011). We estimate daily LAI values from a spline-fit to daily Plant Area Index (PAI) measurements from the HFEMS over April–
 225 December for years 1998–2015 (Matthes et al., 2024), corrected for the reported stem and twig area index (STAI) of $0.9 \text{ m}^2 \text{ m}^{-2}$ noted for this canopy (Horii et al., 2005). Estimated climatological daily LAI values range from $\sim 0.9 \text{ m}^2 \text{ m}^{-2}$ in winter to $4.3 \text{ m}^2 \text{ m}^{-2}$ in summer, in good agreement with MODIS-derived LAI at the location of Harvard Forest (Fig. S5).

2.3 Measured diffusion coefficients of atmospherically relevant molecules

A main result of Nguyen et al. (2015) was the importance of molecular diffusion in atmosphere–surface exchange of rapidly
 230 depositing compounds, where it was shown that maximum daytime dry deposition velocities scale with the inverse square root of molecular mass, as do gas phase diffusion coefficients (Poling and Prausnitz, 2004). To evaluate the calculation of molecular diffusivities used in the parameterization of gaseous dry deposition velocities in GEOS-Chem, we conducted a literature search to compile a list of measured diffusion coefficients for atmospherically relevant molecules, consisting of 23 inorganic and 17 organic species (Table S1). Diffusion coefficients (D) measured in either air or N_2 near STP were corrected to STP following
 235 Langenberg et al. (2020):

$$D = D_o \left(\frac{P_o}{P} \right) \left(\frac{T}{T_o} \right)^b, \quad (3)$$

where we set the temperature power dependence $b = 1.75$ following Fuller’s method, a semi-empirical technique for the estimation of binary gas-phase diffusion coefficients (Fuller et al., 1966), discussed further in Section 3.2.

2.4 Measurements of surface-specific deposition velocities for NO_2

240 Surface-specific NO_2 uptake coefficients (γ_{NO_2}) to both foliar and non-foliar forest elements facilitate bottom-up estimates of bulk-canopy $R_c(\text{NO}_2)$ and resulting $V_d(\text{NO}_2)$ to forest environments when corresponding surface area scale factors (i.e., LAI and STAI) and meteorological data are available. From literature values of surface-specific deposition velocities V_d^{surf} , we infer γ_{NO_2} for both non-foliar and foliar materials:

$$\gamma_{\text{NO}_2} = \frac{4 V_d^{\text{surf}}}{\bar{v}_t}, \quad (4)$$

245 where $\bar{v}_t$ is the mean thermal speed of NO_2 . Table S2 in the supplement contains literature values of V_d^{surf} and associated experimental temperatures which were used to infer values of γ_{NO_2} to foliar surfaces of deciduous and coniferous species under nocturnal/dark conditions, non-foliar forest materials (bark and forest floor), snow, and fabricated materials. With the exception of deposition to snow, literature values of V_d^{surf} are from chamber studies where mechanically mixed chamber air enables the direct estimation of γ_{NO_2} through Eq. (4), i.e., turbulent (R_a) and quasi-laminar (R_b) resistances may be neglected
 250 for species with slow surface uptake (such as NO_2) since $R_c \gg R_a + R_b$. Values of leaf-level V_d^{surf} for both deciduous and

coniferous species were averaged across periods of minimum stomatal conductance resulting from the absence of photosynthetically active radiation (PAR) and are interpreted herein for the purpose of computing the resulting γ_{NO_2} value as non-stomatal. Table S2 also includes corresponding surface areas used for flux normalization; care must be taken when comparing surface-specific V_d^{surf} and γ_{NO_2} , as various surface areas are used (i.e., planar, geometric, projected leaf area, and total leaf area). Hanson et al. (1989) report V_d^{surf} to coniferous species normalized to total leaf area, as stomata are distributed across the whole needle surface (amphistomatic). Often, studies normalize to projected leaf area, as is routinely done for deciduous leaves which generally have stomata on the lower (abaxial) leaf surface. Failing to recognize this difference would result in a misrepresentation of V_d^{surf} and inferred γ_{NO_2} by a factor of ~ 2.7 for coniferous species (Riederer et al., 1988).

Table 1 summarizes values of γ_{NO_2} , including those used in Section 3.3.4 to compute nocturnal bottom-up estimates of bulk-canopy $R_c(NO_2)$ and resulting $V_d(NO_2)$ over Harvard Forest. Also included are associated surface area scale factors α for which γ_{NO_2} is to be applied and relative humidities over which measurements were made. We suggest that NO_2 uptake to the surfaces listed in Table 1 may result from heterogeneous hydrolysis of NO_2 following reaction R1, with variability between surfaces primarily a result of differences in microscopic surface area supporting adsorbed water. Some of the studies measuring foliar uptake of NO_2 under conditions where stomatal aperture should be at a minimum conclude that uptake could occur to the interior of leaves via partially open stomata rather than non-stomatally to the exterior leaf surfaces (Breuninger et al., 2013; Chaparro-Suarez et al., 2011; Delaria et al., 2020; Rondón et al., 1993). Our assumption of nocturnal stomatal closure with deposition of NO_2 to the exterior of leaves via reaction R1 is discussed further in Section 4. To help contextualize values of γ_{NO_2} , Section S3 of the supplement provides a brief literature review of uptake coefficients for NO_2 to hydrated surfaces.

3 Measurement–model comparisons and updates

Table 2 summarizes modifications made herein to the offline gaseous dry deposition parameterization from GEOS-Chem, discussed in-turn throughout this section. Briefly, parameterization P1 is equivalent to the dry deposition scheme in GEOS-Chem, which references deposition from grid box centers (GBC) of the lowest model level (~ 60 m). Serial modifications to P1 include changes to the height in which dry deposition is referenced (P2), formulations to the calculation of aerodynamic resistance (P3) and molecular diffusivities (P4), updating non-stomatal surface resistance for NO_2 following the Z03 scheme (P5) and subsequent replacement with a scheme that represents heterogeneous hydrolysis on deposition surfaces (P6, P7), and finally, implementation of empirical updates to the non-stomatal uptake of PAN (P8). We begin by evaluating parameterizations P1–P4 by comparing to measured dry deposition velocities from Nguyen et al. (2015), where it was noted that above-canopy deposition velocities for H_2O_2 , HMHP, and HNO_3 corresponded to computed theoretical maximums (i.e., $V_{d,max} \sim [R_a + R_b]^{-1}$), thus enabling a more direct evaluation of the deposition pathway consisting of resistances R_a and R_b , as

280 **Table 1:** Surface-specific NO₂ uptake coefficients γ_{NO_2} inferred from literature values of surface-specific deposition velocities^(a) following Eq. (4). Also included are surface area scale factors α for which γ_{NO_2} is to be applied, and surface areas and relative humidities over which surface-specific deposition measurements were made.

Material	γ_{NO_2}	α	Surface Area ^(b)	RH	Ref. ^(g)
	[unitless]	[unitless]		[%]	
Non-Foliar surfaces					
distilled water	2.3 x 10 ⁻⁶	-	planar	N/A	1
wood board (untreated, hard, fine)	7.6 x 10 ⁻⁷	-	geometric	70	2
plywood (untreated)	1.6 x 10 ⁻⁶	-	geometric	90	
tree bark (dry) ^(c)	1.4 x 10 ⁻⁶	-	geometric	50	2
tree bark (wet) ^(c)	5.0 x 10 ⁻⁶	π STAI	geometric	unknown	1
forest floor ^(c)	1.0 x 10 ⁻⁵	π STAI	geometric	N/A	
snow ^(c)	4.3 x 10 ⁻⁵	1	planar	~ 60 +/- 20	3
	1.6 x 10 ⁻⁵	1	planar	N/A	4
Foliar surfaces^(d, e)					
deciduous leaves ^(c)	1.6 x 10 ⁻⁶	LAI	projected	50 to < 90	1, 6–8
coniferous leaves	4.1 x 10 ⁻⁶	LAI	projected	50 to < 90	1,3,5,6,9
coniferous leaves ^(c)	1.5 x 10 ⁻⁶	2.7LAI	total ⁽ⁱ⁾	50 to < 90	1,3,5,6,9

(a) Surface-specific deposition velocities were taken from chamber studies, with the exception of uptake to snow which was measured via the eddy covariance technique. Table S2 in the supplement contains study specific details.

285 (b) Surface area used to normalize surface-specific deposition fluxes in the computation of surface-specific v_d^{surf} (Table S2).

(c) Values used in Section 3.3.4 to compute bottom-up estimates of nocturnal bulk-canopy $V_d(NO_2)$ over Harvard Forest.

(d) Foliar uptake was measured under conditions of minimal stomatal aperture, i.e., dark conditions. We assume this uptake to be non-stomatal (Section 4).

(e) Multi-study mean values computed herein (Table S2).

290 (f) Reported v_d^{surf} normalized to projected leaf areas were scaled herein to reflect uptake to total leaf surface area—a factor of 2.7 for coniferous needles (Riederer et al., 1988).

(g) References for surface-specific $v_d^{surf}(NO_2)$

(1) Hanson et al. (1989)

(2) Grøntoft and Raychaudhuri (2004)

295 (3) Rondón et al. (1993)

(4) Stocker et al. (1995)

(5) Wang et al. (2020)

(6) Delaria et al. (2020)

(7) Delaria et al. (2018)

300 (8) Chaparro-Suarez et al. (2011)

(9) Breuninger et al. (2013)

discussed in Sections 3.1 and 3.2, respectively. Parameterizations P5–P7 are evaluated by comparing to both above-canopy nocturnal $V_d(NO_2)$ observed at the HFEMS (Section 3.3.3) and bottom-up estimates of nocturnal $V_d(NO_2)$ for Harvard Forest

305 from literature values of surface-specific deposition velocities $V_d^{surf}(NO_2)$ (Section 3.3.4). Parameterization P8 is evaluated in Section 3.4 in the context of effects on simulated $V_d(NO_y)$, including comparison to above-canopy diel $V_d(NO_y)$ observed at the HFEMS.

Table 2: Modifications to the offline dry deposition parameterization tested in this study. Parameterization P1 is equivalent to the gaseous dry deposition scheme in GEOS-Chem (GC). Modifications to P1 include changes to reference height z_{ref} (P2), formulation of aerodynamic resistance R_a (P3), molecular diffusivity D (P4), and non-stomatal surface resistances (R_c) for NO_2 (P5–P7) and PAN (P8).

Param.	$z_{ref}^{(a)}$	Aerodynamic Res. R_a	Diffusivity D	non-stomatal $R_c(NO_2)$	non-stomatal $R_c(PAN)$
P1	z_{GBC}	base GC (Eq. (S1))	base GC (Chapman–Enskog theory with constant mfp ^(b))	base GC (modified W89)	base GC (modified W89)
P2	z_{TNF} or z_{HFEMS}				
P3		RSL, $u(z_o) > 0 \text{ m s}^{-1}$ (Eq. (S13))	measured & Fuller’s method (Eqs. (3 & 5))	Z03	
P4					
P5					
P6					
P7				r_{hyd} with $\alpha = 1$ (Eq. (8))	
P8	r_{hyd} with $\alpha = 2$ (Eq. (8))			empirical ^(c)	

^(a) Dry deposition reference height: $z_{GBC} \sim 60 \text{ m}$, $z_{TNF} = 22 \text{ m}$; $z_{HFEMS} = 29 \text{ m}$.

^(b) Mean Free Path (mfp) held constant across depositing gases.

^(c) Empirical fit of non-stomatal cuticular deposition (Turnipseed et al., 2006), modified herein for LAI (Section S5 of the supplement).

3.1 Updates to the calculation of aerodynamic resistance

Table 3 contains an evaluation of simulated atmospheric resistances (R_a and R_b) in parameterizations P1–P4 by comparing to measured daytime deposition velocities for rapidly depositing species from Nguyen et al. (2015). In contrast to the findings of Wu et al. (2021) showing excellent model-measurement agreement to peak daytime $V_d(\text{HNO}_3)$ and $V_d(\text{H}_2\text{O}_2)$ between the Z03 scheme and the dataset from Nguyen et al. (2015), parameterization P1 overestimates daytime mean deposition velocities for these rapidly depositing species: H_2O_2 (+15 %), HMHP (+41 %), and HNO_3 (+52 %). Nguyen et al. (2015) found excellent agreement between hourly GEOS-FP assimilated meteorology at this site (used herein for computation of V_d in Table 3) and measured values, including u_* and sensible and latent heat fluxes. GEOS-FP fields report a summertime $z_o = 2.2 \text{ m}$ for the $0.25^\circ \times 0.3125^\circ$ grid cell that includes the CTR site—greater than would be expected at this site given the local 10 m canopy height. Prescribing z_o to be 10 % of h_c in parameterization P1b, in accordance with conventionally used values for natural vegetation and in agreement with an updated land-use module developed for GEOS-Chem (Geddes et al., 2016), results in a 35 % increase in R_a and notable reductions in V_d high biases. However, following the computation of R_a in GEOS-Chem, P1b computes R_a from a reference height z of $\sim 60 \text{ m}$ despite a measurement height of 22 m at the CTR site, while neglecting to include a displacement height d . Neglecting d from the computation of R_a in Eq. (S1) increases daytime R_a in parameterization P1b by 1 % when referenced from 60 m and 9 % when referenced from 22 m . Although the greatest sensitivity of R_a to z occurs in proximity to z_o (Fig. S2), the difference between R_a computed from an above-canopy measurement height vs. typical heights from which global CTMs reference dry deposition can be significant (Figs. S1 & S2). Referencing R_a from the CTR

measurement height of 22 m in parameterization P2 results in a 23 % decrease in daytime R_a , returning V_d biases to P1 levels (Table 3).

Table 3: Effects of updates to the calculation of aerodynamic resistance R_a and quasi-laminar sublayer resistance R_b on simulated daytime (10–15 LST) dry deposition velocities over Talladega National Forest (temperate, mixed) for three rapidly depositing species. Serial modifications to base parameterization P1 are highlighted, i.e., PX (update). Shown are mean quantities $\pm$ standard deviations about the hourly timeseries^(a).

Parameterization ^(b)	H ₂ O ₂				HMHP			HNO ₃		
	R_a	R_b	V_d	NMB ^(c)	R_b	V_d	NMB ^(d)	R_b	V_d	NMB ^(e)
	[s/m]	[s/m]	[cm/s]	[%]	[s/m]	[cm/s]	[%]	[s/m]	[cm/s]	[%]
P1 (base sim.)	9.5±2.7	7.1±2.1	6.0±1.2	15	7.7±2.9	5.8±1.1	41	7.6±2.9	5.8±1.1	52
P1b ($z_o = 0.1h_c$) ^(f)	12.8±3.4		5.0±1.0	-3		4.9±1.0	19		4.9±1.0	29
P2 ($z_{ref} = 22$ m - d)	9.7±2.5		5.9±1.3	14		5.8±1.2	40		5.8±1.2	51
P3 (RSL, $u(z_o) > 0$)	10.2±2.6		5.8±1.2	11		5.6±1.2	36		5.6±1.2	47
P4 (D update)		12.9±4.9	4.4±1.0	-15	14.9±5.7	4.1±1.0	-1	15.7±6.0	4.0±0.9	4

^(a) Mean quantities are averaged across the five daytime periods in June 2013 that Nguyen et al. (2015) use in their analysis of eddy covariance observed deposition velocities (Section 2.2.1).

^(b) Table 2 contains a list of parameterization updates. R_c set to 1 s m⁻¹ following minimum allowed in GEOS-Chem (Section S1 of supplement).

^(c) To measured (eddy covariance) daytime (10–15 LST) $V_d(H_2O_2) = 5.2 \pm 1.1$ cm s⁻¹ (Nguyen, 2015).

^(d) To measured (eddy covariance) daytime (10–15 LST) $V_d(HMHP) = 4.1 \pm 1.1$ cm s⁻¹ (Nguyen, 2015).

^(e) To measured (eddy covariance) daytime (10–15 LST) $V_d(HNO_3) = 3.8 \pm 1.3$ cm s⁻¹ (Nguyen, 2015).

^(f) z_o set to 10% of canopy height (h_c) for parameterizations P1b–P8.

Considering that the CTR and HFEMS measurement heights, $\sim 2 h_c$ and $1.5 h_c$, respectively, are at the upper limits of the roughness sublayer (RSL), a region where turbulent mixing in the wake of roughness elements is enhanced above that predicted by M–O similarity theory by a factor of 2 to 3 (see Section S2.4 of the supplement for a review of RSL mixing), R_a computed according to M–O similarity theory following Eq. (S1) may be in slight underestimate due to non-zero horizontal winds at z_o resulting from enhanced downward mixing of momentum. To quantify this effect, parameterization P3 computes R_a corrected for RSL mixing which allows $u(z_o) > 0$ m s⁻¹ (Eq. (S13)), resulting in a small (5 %) increase in R_a at the CTR measurement height under the daytime conditions of Table 3, and even smaller changes to V_d given the influence of R_b (Section 3.2). As demonstrated herein and in agreement with previous work (Simpson et al., 1998), it may be appropriate to neglect the effects of the RSL on depositing species when referenced from a height of at least $1.5\text{--}2 h_c$. However, studies endeavouring to understand bidirectional exchange or the dispersion of near-surface emissions should consider the effect of asymmetrical R_a that the RSL imposes (Sections S2.2 & S2.4).

3.2 Updates to the calculation of molecular diffusivities

As seen in Table 3, updates to the calculation of R_a failed to address high biases in simulated deposition velocities of rapidly depositing species. Larger molecular weight species HMHP and HNO₃ exhibit a greater high bias in V_d , 36 % and 47 %, respectively.

respectively, than the lower molecular weight species H_2O_2 (11 %). Given the dependence of $V_{d,max}$ on molecular diffusivity through influence on R_b (Eq. (S2)) (Meyers et al., 1989), we evaluate the calculation of molecular diffusion coefficients in GEOS-Chem against measured values for atmospherically relevant molecules.

Figure 1 depicts a large high bias in calculated diffusion coefficients from the dry deposition module of GEOS-Chem, which uses the Chapman–Enskog theory for binary diffusivity (Seinfeld, 1986). The bias results from the use of a constant collision diameter $\sigma = 2.7 \text{ \AA}$ for all species with air—an underestimate for many atmospherically relevant molecules, i.e., σ for O_3 with air is $3.793 \text{ \AA}$ (Massman, 1998; Poling and Prausnitz, 2004). The collision diameter σ is a pairwise characteristic length scale of the Lennard–Jones intermolecular force, which is not readily available for many atmospheric trace gasses (Tang et al., 2014). Several semi-empirical methods have been proposed for the estimation of D in low pressure binary systems (Poling and Prausnitz, 2004). Fuller et al. (1966) developed a simple and generalized semi-empirical correlation equation for the estimation of binary gas phase diffusion coefficients using additive atomic diffusion volumes V_i for each species $\sum_A V_i$ and $\sum_B V_i$. The diffusion coefficient $D [\text{cm}^2 \text{ s}^{-1}]$ for trace gas A in bath gas B is given by:

$$D = \frac{10^{-3} T^{1.75} (1/M_A + 1/M_B)^{1/2}}{P [(\sum_A V_i)^{1/3} + (\sum_B V_i)^{1/3}]^2}, \quad (5)$$

where P is the pressure [atm], T is the temperature [K], and M is the molecular mass [g mol^{-1}]. Atomic, and in some cases molecular, diffusion volumes were obtained from regression analysis of 153 binary systems across 340 T–P states and are summarized in Poling and Prausnitz (2004), Tang et al. (2014), and Tang et al. (2015). As seen in Fig. 1, diffusion coefficients computed using Fuller’s method result in a much-improved comparison to measurements, with better agreement to organic species ($R^2 = 0.99$ and $\text{NMB} = -3 \%$) than to inorganics ($R^2 = 0.88$ and $\text{NMB} = 13 \%$), consistent with the findings of Tang et al. (2014 & 2015).

Figure 1 also depicts molecular diffusion coefficients approximated by Graham’s law of effusion, i.e., $D_{1k} = D_{2k} \sqrt{M_2/M_1}$ (Mason and Evans III, 1969), where (continuum) diffusion coefficients are approximated by Knudsen diffusion coefficients D_k —an oversimplification of Eq. (5) and a strategy commonly used in the atmospheric science community (Nguyen et al., 2015; Weber and Renenberg, 1996; Wesely, 1989). Resulting diffusion coefficients scaled from measured D_{CO_2} correlate well with measured values ($R^2 = 0.91$), with NMB to inorganic and organic species of 3 % and 20 %, respectively. In a review of molecular diffusivities of atmospherically relevant molecules, Massman et al. (1998) note misapplication of Graham’s law to molecular diffusivities can lead to errors of up to 23 %. Measured and computed diffusion coefficients from Fuller’s method assuming air as the bath gas are presented in Fig. 1 and tabulated in Table S1. We do not differentiate between diffusivity measurements carried out in air or N_2 , as differences are expected to be small, i.e., a 2 % difference in D_{O_3} at STP in air vs N_2 .

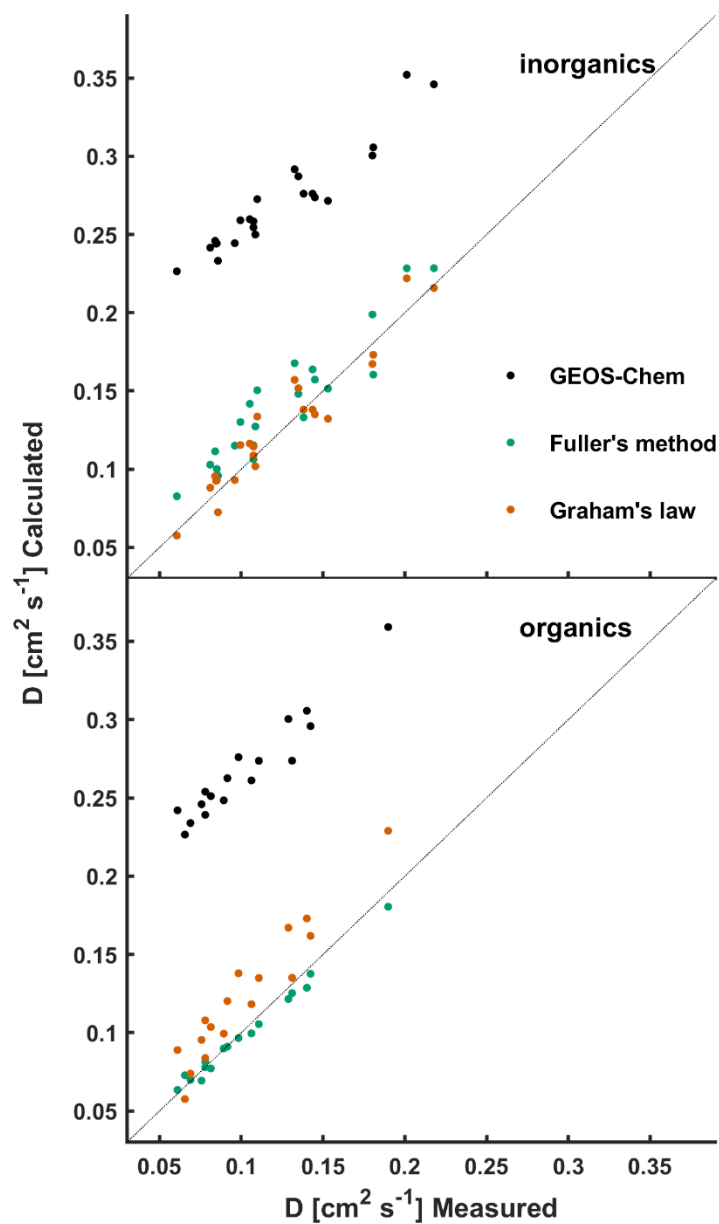


Figure 1: Measured diffusion coefficients of atmospherically relevant molecules in air or N₂ at STP are compared to calculated values. Molecular diffusivities calculated following the method used in GEOS-Chem (P1–P3) are compared to those calculated following Fuller’s method and Graham’s law (referenced from D_{CO_2}). Measured and computed (Fuller’s method) values are listed in Table S1 of the supplement.

Parameterization P4 computes R_b using measured diffusion coefficients when available and diffusion coefficients according to Fuller's method in the absence of measured values. Diffusion coefficients are adjusted to ambient T-P following Eq. (3) prior to calculating R_b . Eliminating the high bias in calculated molecular diffusivities resulted in a near doubling of R_b for the species in Table 3, and a much-improved comparison to the daytime deposition velocities for the larger molecular weight species HMHP (NMB -1 %) and HNO_3 (NMB 4 %). The increase in R_b for H_2O_2 results in a low bias of -15 %, which is well within the large relative uncertainty for R_b due to variations in canopy structures (Massman et al., 1994; Sievering et al., 2001).

Molecular diffusivity is also involved in the calculation of R_c via influence on stomatal resistance r_s , which is scaled by the ratio $D_{\text{H}_2\text{O}}/D_x$ in dry deposition parameterizations commonly used in chemical transport models (Wesely, 1989; Zhang et al., 2003a). The effect of updated molecular diffusivity on R_c in GEOS-Chem is significant for molecules which dry deposit under stomatal control, i.e., species with low aqueous solubility or surface reactivity, and is discussed in Section S5 of the supplement.

3.3 Nocturnal dry deposition of NO_2 over Harvard Forest

3.3.1 Eddy covariance inferred $V_d(\text{NO}_2)$

Nocturnal hourly eddy covariance NO_2 fluxes and resulting exchange velocities $V_{\text{ex}}(\text{NO}_2)$ over Harvard Forest from April–November 2000 are shown in Fig. 2 as a function of NO_2 concentration. We restrict our analysis to nighttime (20–04 LST), when above-canopy $\text{NO}_2 : \text{NO}_x \sim 1$ and photochemical flux divergence of the $\text{NO}-\text{NO}_2-\text{O}_3$ triad due to the presence of a vertical gradient in irradiance through the forest canopy (Gao et al., 1993) is absent. As seen in the top panel of Fig. 2, nocturnal fluxes of NO_2 over Harvard Forest are predominantly (~ 70 %) downward, especially at higher ambient NO_2 concentrations. Nocturnal mean (median) $\pm 1\sigma$ fluxes of NO_2 from April through November are -0.8 (-0.3) ± 2 ppb cm s^{-1} . These downward ($p < 0.01$) above-canopy aggregate fluxes of NO_2 are comparable in magnitude to counteracting summertime nocturnal soil NO emissions, estimated by Munger et al. (1996) through a mass-balance approach to be $\sim 0.9 \mu\text{mol m}^{-2} \text{h}^{-1}$ ($3.5 \text{ ng N m}^{-2} \text{s}^{-1}$, or $0.62 \text{ ppb cm s}^{-1}$) at the HFEMS. Munger et al. (1996) note that nocturnal NO is elevated near the forest floor and Horii et al. (2004) find decreasing within-canopy nocturnal NO profiles at Harvard Forest with above-canopy concentrations and fluxes indistinguishable from zero despite net downward fluxes of NO_2 , presumably due to titration of soil-emitted NO by O_3 on a timescale much shorter (minutes) than in-canopy vertical mixing, followed by nocturnal canopy loss processes for NO_2 . Previous studies have noted the importance of knowledge of local soil NO emissions and within-canopy processes involving NO_x when interpreting above-canopy NO_2 fluxes (Delaria and Cohen, 2020; Eugster and Hesterberg, 1996; Flechard et al., 2011; Min et al., 2014).

In an effort to isolate the contribution that dry deposition makes to above-canopy nocturnal eddy covariance fluxes (F_{EC}) of NO_2 , we infer $V_d(\text{NO}_2)$ following Eq. (6) to account for the effects of nocturnal chemical flux divergence (V_{chem}) and counteracting soil NO emissions assumed to rapidly titrate with O_3 and ventilate the canopy as NO_2 (F_{soil}). The resulting

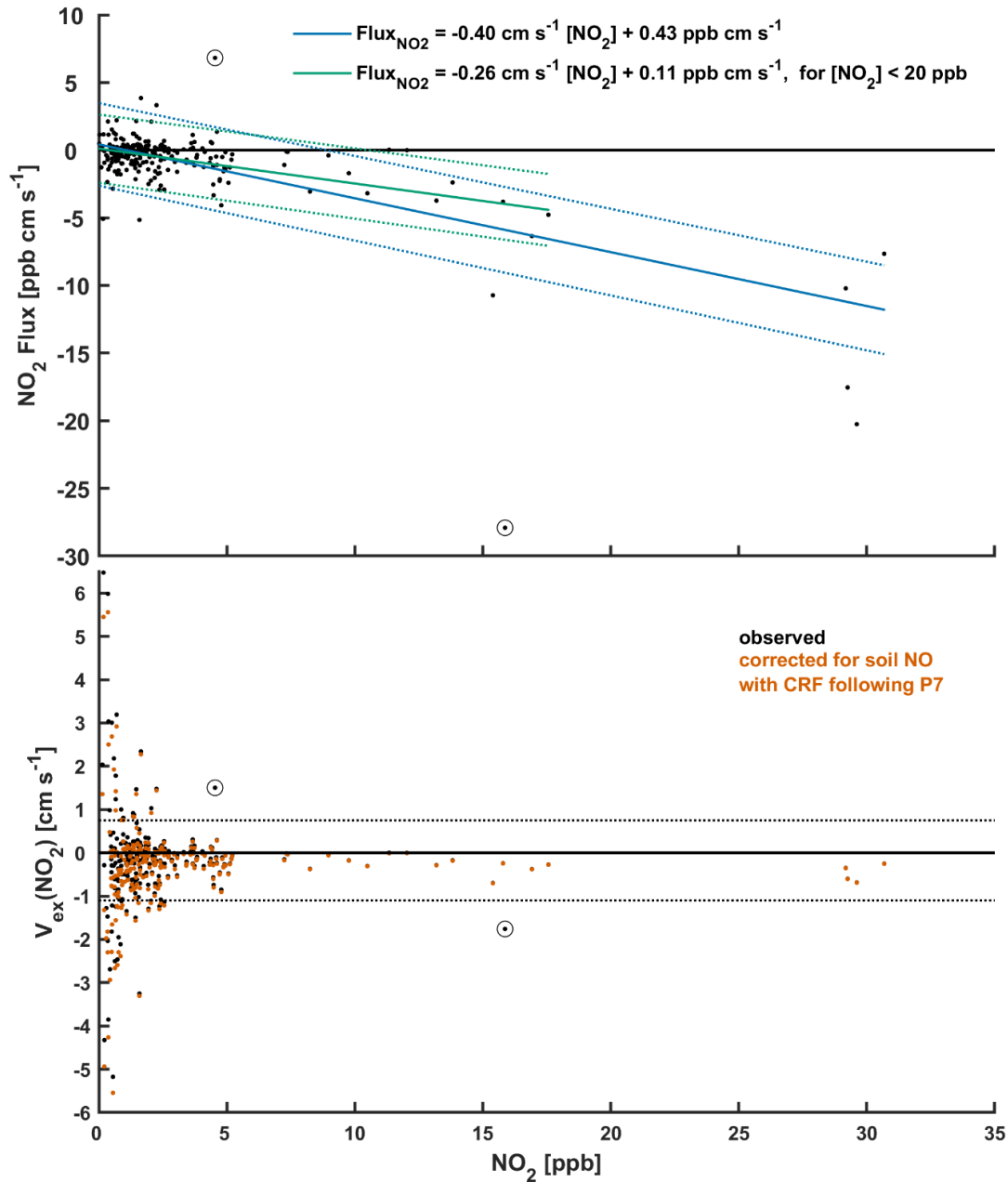


Figure 2: (TOP) Nocturnal (20–04 local solar time) hourly eddy covariance NO_2 fluxes and **(BOTTOM)** resulting exchange velocities $V_{\text{ex}}(\text{NO}_2)$ as a function of NO_2 concentration. These publicly available measurements (Section 2.2.2) were taken over an established mixed forest (Harvard Forest) from April–November 2000. Estimated soil NO flux (assumed to ventilate the canopy as NO_2) was subtracted from measured hourly NO_2 fluxes in order to estimate $V_{\text{ex}}(\text{NO}_2)$ corrected for soil NO (Eqs. (6–7)). Included in the top plot are linear fits and associated 95% prediction intervals. Dashed lines in the bottom plot depict boundaries of an outlier filter applied to hourly $V_{\text{ex}}(\text{NO}_2)$ prior to calculation of means (Section 2.2.2). Data points excluded from analysis based on visual inspection are circled. Hourly observations made under conditions of low turbulence ($u_* < 0.2 \text{ m s}^{-1}$) were excluded from this analysis.

observation-inferred $V_d(NO_2)$ is a best estimate of the nocturnal dry deposition pathway with which to evaluate parameterizations:

$$V_d + V_{chem} = -V_{ex} = -\frac{(F_{EC} - F_{soil})}{[NO_2]}, \quad (6)$$

where V_{ex} is the eddy covariance observed NO_2 exchange velocity which does not assume predominant deposition and therefore has sign convention analogous to F_{EC} ; V_{chem} represents an estimate of below-sensor nocturnal chemical loss of NO_2 via formation and loss of N_2O_5 , limited by the rate of oxidation of NO_2 with O_3 (Browne and Cohen, 2012; Jacob, 2000). We use an estimate of the maximum rate of nocturnal chemical loss of NO_2 proposed by Horii (2002) in their analysis of the dataset used herein, $V_{chem} \sim 0.05 \text{ cm s}^{-1}$, which translates to a below-sensor ($< 29 \text{ m}$) nocturnal chemical lifetime of NO_2 to oxidation by O_3 of $\sim 16 \text{ h}$. The bottom panel of Fig. 2 includes hourly values of $V_{ex}(NO_2)$, both uncorrected and corrected for soil NO . Values of F_{soil} used in Eq. (6) are less than the peak summertime forest floor estimate from Munger et al. (1996) due to seasonality and within-canopy loss processes. Hourly estimates of F_{soil} were calculated by scaling the reported summertime nocturnal soil NO emission flux at Harvard Forest $F_{NO,summer}$ ($0.62 \text{ ppb cm s}^{-1}$) by GEOS-Chem simulated seasonality κ and a parameterized canopy reduction factor CRF :

$$F_{soil}(hr) = F_{NO,summer} \kappa(month) [1 - CRF(hr)], \quad (7)$$

Month-specific κ scale factors were obtained by normalizing simulated monthly mean nocturnal soil NO emission, output at the location of the HFEMS from a high resolution ($0.25^\circ \times 0.3125^\circ$) GEOS-Chem simulation, by the peak monthly mean simulated emission (July at the location of HFEMS). GEOS-Chem simulated soil NO emission in the region of Harvard Forest exhibits significant seasonality, with a winter minimum a small fraction ($< 5 \%$) of the summertime maximum (Fig. S3). Section S4 of the supplement describes the parameterization of CRF used in GEOS-Chem and herein in Eq. (7). As NO_2 surface uptake resistance $R_c(NO_2)$ is used in the calculation of CRF , values are parameterization specific (Fig. S3), with larger values of CRF resulting from lower values of $R_c(NO_2)$.

Figure 3 depicts monthly nocturnal $V_d(NO_2)$ inferred from Eq. (6) over the HFEMS from April–November 2000 alongside coincidently-sampled simulated values from parameterizations P4–P7. Table 4 presents observation-inferred and simulated values of $V_d(NO_2)$ aggregated across all months, as well as associated NO_2 lifetimes to dry deposition from the 29 m measurement height. We begin discussion of eddy covariance inferred bulk-canopy $V_d(NO_2)$ below, followed by discussions in Section 3.3.2 of the (i) large low bias in simulated values stemming from the widely used W89 parameterization of surface resistances and (ii) reduced bias using the Z03 scheme. In Section 3.3.3, we evaluate a simple representation of non-stomatal NO_2 uptake following reaction R1 against eddy covariance inferred $V_d(NO_2)$.

As previously mentioned, hourly values of observed $V_{ex}(NO_2)$ were subjected to an outlier filter (Fig. 2) prior to computation of mean values, whereas median and ‘mean flux-to-mean concentration’ ratios ($\bar{F}/[\overline{NO_2}]$) included in Table 4 were not, and instead computed directly from u_* filtered hourly data as the latter two statistics

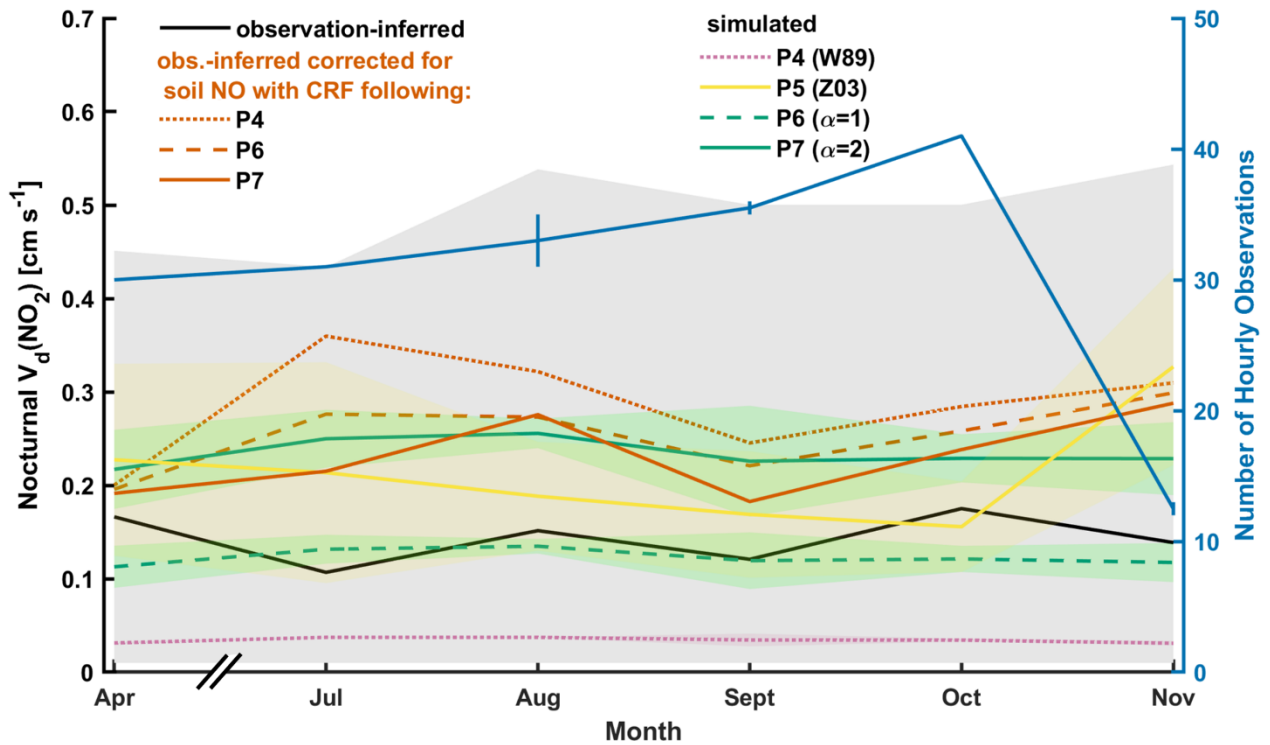


Figure 3: Observation (eddy covariance)-inferred and simulated monthly mean nocturnal (20–04 local solar time) NO_2 deposition velocities $V_d(\text{NO}_2)$ over Harvard Forest. Simulated values are coincidentally sampled with hourly observations prior to averaging. Also depicted is observation-inferred $V_d(\text{NO}_2)$ corrected for soil NO emission using simulated soil NO canopy reduction factors (CRF) from parameterizations P4, P6, and P7. Standard deviations about simulated monthly mean values, as well as observation-inferred monthly mean values uncorrected for soil NO, are depicted as shaded areas. Month-specific range in number of hourly observations used in calculation of monthly means are indicated as vertical lines, and result from parameterization-specific soil NO corrections causing differential outlier filter exceedance (Fig. 2 and Table 4). Insufficient data prevented analysis for May and June (Fig. S6).

are less influenced by outliers than arithmetic means. Aggregate values of $\bar{F}/[\text{NO}_2]$ in Table 4 are in the same units as $V_d(\text{NO}_2)$ [cm s^{-1}] and include corrections for F_{soil} and V_{chem} as do mean and median quantities computed from hourly values following Eq. (6). Assuming first-order dependence of NO_2 dry deposition with concentration (Eq. (1)), computing values of $\bar{F}/[\text{NO}_2]$ over long averaging times is a strategy to reduce the influence of random variability in deposition velocity estimates, especially under low NO_2 conditions as evident in Fig. 2.

As seen in Fig. 3, monthly mean values of observation-inferred $V_d(\text{NO}_2)$ uncorrected for the influence of soil-emitted NO are in the range of 0.1–0.2 cm s^{-1} . Although variability is large, with standard deviations greater than mean values, corrections for soil NO (venting the canopy as NO_2) results in a significant ($p < 0.03$) increase in nocturnal $V_d(\text{NO}_2)$, yielding monthly mean values in the approximate range of 0.2–0.3 cm s^{-1} . Both uncorrected and soil NO corrected nocturnal $V_d(\text{NO}_2)$ lack discernible seasonality. In Section 3.3.4, bottom-up estimates of nocturnal $V_d(\text{NO}_2)$ for Harvard Forest are developed in an effort to understand the apparent lack of seasonality in top-down observations. Large variability in eddy covariance observed NO_2 flux and resulting deposition velocities have been noted in other studies (Eugster and Hesterberg, 1996; Farmer et al.,

2006; Geddes and Murphy, 2014), wherein authors restrict analysis to average values in order to reduce the variability in these complex ecosystem-scale observations (Baldocchi, 2003). Herein, we restrict analysis to average values over at least one month.

Table 4: Nocturnal (20–04 local solar time) NO₂ deposition velocities over Harvard Forest aggregated from April–November. Observation-inferred values, with and without estimated soil NO corrections using simulated canopy reduction factors (CRF) corresponding to parameterizations P4, P6, and P7, are shown along with coincidentally sampled simulated values. Measurements under conditions of low turbulence (friction velocity $u_* < 0.2 \text{ m s}^{-1}$) were excluded from analysis, leaving 230 hourly observations in the timeseries (Section 2.2.2).

	CRF	Va(NO2) [cm s ⁻¹]			Lifetime to dry deposition ^(c)
	[%]	mean ^(a)	median	$\bar{F}/[\overline{NO_2}]^{(b)}$	[h]
Obs.-inferred					
no soil NO	N/A	0.15 ± 0.34 (185)	0.13	0.21	5.4
soil NO, CRF(P4)	31	0.28 ± 0.35 (181)	0.25	0.3	2.9
soil NO, CRF(P6)	47	0.25 ± 0.34 (181)	0.22	0.28	3.2
soil NO, CRF(P7)	59	0.23 ± 0.35 (183)	0.21	0.26	3.5
Simulated ^(d)					
P4 (R _a & D)	N/A	0.04 ± <0.01	0.04	-	20
P5 (Z03)	N/A	0.20 ± 0.09	0.18	-	4.0
P6 (α = 1)	N/A	0.12 ± 0.02	0.13	-	6.7
P7 (α = 2)	N/A	0.24 ± 0.04	0.24	-	3.4

^(a) Observation-inferred hourly values of $V_{ex}(NO_2)$, both uncorrected and corrected for soil NO, were subjected to an outlier filter (Fig. 2) prior to calculation of arithmetic mean $V_d(NO_2)$; remaining number of hourly observations are included in brackets adjacent corresponding mean values and standard deviations.

^(b) The ratio of ‘mean NO₂ flux-to-mean NO₂ concentration’ ($\bar{F}/[\overline{NO_2}]$) in units of [cm s⁻¹] is included for comparison (Section 3.3.1).

^(c) Calculated at 29 m (measurement height) from aggregate mean deposition velocities.

^(d) See Table 2 for serial updates. Briefly,
P4: updates to aerodynamic resistance (R_a) and molecular diffusivity (D).
P5: compute resistance to non-stomatal surface uptake of NO₂ following Zhang et al. (2003a).
P6 : compute resistance to non-stomatal surface uptake of NO₂ following reaction R1, with surface area scale factor $\alpha = 1$ in Eq. (8).
P7: analogous to P6 but with $\alpha = 2$.

3.3.2 Evaluation of nocturnal V_d(NO₂) from GEOS-Chem and Z03

Parameterization P4, which computes NO₂ surface uptake resistance $R_c(NO_2)$ following the W89 representation in GEOS-Chem, yields a simulated nocturnal $V_d(NO_2)$ that is biased low by nearly 4-fold compared to observations uncorrected for soil NO, increasing to a 7-fold low bias after correcting for soil NO from parameterization P4 (Table 4). This underestimate is driven by the large nocturnal $R_c(NO_2)$ of $\sim 2,700 \text{ s m}^{-1}$ in parameterization P4 (Fig. S8), which has been noted in previous studies comparing the W89 algorithm to eddy covariance observations over forest (Horii et al., 2004) and grassland (Eugster and Hesterberg, 1996) ecosystems. Wesely et al. (1982) reported a nocturnal eddy covariance observed $V_d(NO_2)$ of 0.05 cm s^{-1} over a summertime soybean field, similar to the P4 value in Table 4. The authors acknowledge that counteracting soil NO emissions may have resulted in low measured values of above-canopy NO₂ deposition. In their analysis of eddy covariance

515 fluxes of NO_2 over a managed grassland in central Switzerland, Eugster & Hesterberg (1996) found that accounting for counteracting fluxes of soil-emitted NO , oxidized to NO_2 below the height of the sensor (~ 2.7 to $3.6 \text{ ng N m}^{-2} \text{ s}^{-1}$), resulted in an increase in inferred nocturnal $V_d(\text{NO}_2)$ by up to a factor of 2, corresponding to an inferred median value for nocturnal non-stomatal $R_c(\text{NO}_2)$ of 700 s m^{-1} (range $500\text{--}950 \text{ s m}^{-1}$)—a surface resistance on the order of 4 times lower than predicted by the W89 algorithm.

520 In parameterization P5, $R_c(\text{NO}_2)$ is computed following the Z03 scheme, resulting in large increases in simulated nocturnal $V_d(\text{NO}_2)$ across all months, as well as increased intra- and inter-month variability (Fig. 3). The Z03 scheme increases non-stomatal NO_2 uptake relative to the W89 scheme. The representation in the W89 scheme results in stomatal control over uptake, with very low non-stomatal uptake (Section S5; Fig. S4). The diurnal behaviour of observed $V_d(\text{NO}_2)$ relative to $V_d(\text{O}_3)$ reported in Wesely et al. (1982) provided support for the W89 assignment. For use in algorithmic developments to gaseous
525 dry deposition, Zhang et al. (2002a) compiled an updated list of half-reaction redox potentials for species of interest in dry deposition models, noting greater oxidizing capacity for both NO_2 and NO over W89 assignments. Zhang et al. (2002a) justify setting non-stomatal conductance of NO_2 at 80 % that of O_3 —greater than the 10 % used in W89—from noted oxidizing capacity and contemporary field studies (Eugster and Hesterberg, 1996; Pilegaard et al., 1998; Rondón et al., 1993; Walton et al., 1997). As seen in Fig. 3, we find the Z03 scheme in parameterization P5 to be in good agreement with eddy covariance
530 inferred nocturnal $V_d(\text{NO}_2)$ at Harvard Forest, with monthly mean values intermediate between observation-inferred values uncorrected and corrected for emissions of soil NO . As seen in Table 4, parameterization P5 compares well to observation-inferred $V_d(\text{NO}_2)$ corrected for soil NO with CRF at greater levels of uptake (i.e., P7 levels).

Although nocturnal $V_d(\text{NO}_2)$ from parameterization P5 is in good agreement with observation-inferred values over Harvard Forest, it is difficult to justify the increase in parameterized non-stomatal uptake of NO_2 in Zhang et al. (2002a), a
535 predecessor to the Z03 scheme, from half-reaction redox potentials alone. Support for the level of non-stomatal NO_2 uptake in the Z03 scheme comes largely from field observations—many of which are referenced herein in subsequent sections as we discuss the plausibility of non-stomatal NO_2 uptake being a result of heterogeneous hydrolysis—rather than suspected NO_2 reduction reactions based on half-reaction redox potentials. Exploring this potential misrepresentation of non-stomatal NO_2 uptake may have important implications towards future representation of HONO surface sources in atmospheric CTMs.

540 3.3.3 Updates to parameterized $V_d(\text{NO}_2)$ by representing NO_2 hydrolysis on deposition surfaces

Horii et al. (2004) note that observed nocturnal dry deposition of NO_2 may result from a surface hydrolysis reaction following R1. In parameterizations P6–P8, we replace the non-stomatal components of the bulk-surface resistance scheme for NO_2 with a dry deposition pathway representing NO_2 hydrolysis $r_{hyd} [\text{s m}^{-1}]$, formulated as a collision-limited heterogeneous reaction with ground surfaces (Cano-Ruiz et al., 1993):

$$545 \quad r_{hyd} = \frac{4}{\gamma_{g,\text{NO}_2} \bar{v}_t \alpha}, \quad (8)$$

where γ_{g,NO_2} is a ground uptake coefficient for NO_2 resulting from heterogeneous hydrolysis on deposition surfaces, $\bar{v}_t$ the mean thermal speed of NO_2 , and α a dimensionless scale factor introduced herein to facilitate application of Eq. (8) across land types of varying surface area densities. Lammel and Cape (1996) recommend reaction R1 be parameterized in atmospheric chemistry models using field-derived uptake coefficients, as realistic conditions are difficult to reproduce in the lab. We use the field-derived ground uptake coefficient γ_{g,NO_2} for reaction R1 from VandenBoer et al. (2013), determined from production of HONO in a winter nocturnal boundary layer in an agricultural region of Colorado, U.S. Consistent with the heterogenous hydrolysis of NO_2 requiring adsorbed water to proceed, VandenBoer et al. (2013) parameterized γ_{g,NO_2} as a function of RH [%] according to Eq. (9) to capture the factor of 2 variability in γ_{g,NO_2} on either side of their best fit value (8×10^{-6}):

$$\gamma_{g,NO_2} = \frac{RH}{50} 8 \times 10^{-6}, \quad (9)$$

Parameterization P6 computes r_{hyd} with $\alpha = 1$, resulting in a simulated nocturnal aggregate mean $V_d(NO_2)$ of $0.12 \pm 0.02 \text{ cm s}^{-1}$ —a 3-fold increase over P4 and satisfactory agreement with observation-inferred $V_d(NO_2)$ uncorrected for soil-emitted NO (Table 4). However, this is an underestimate by $\sim 50 \%$ when soil NO emissions are accounted for. The larger nocturnal mean CRF of 47 % for parameterization P6 is due to reduced nocturnal $R_c(NO_2)$ (median value $\sim 750 \text{ s}^{-1}$, Fig. S8), resulting in a small (11 %) decrease in observation-inferred $V_d(NO_2)$ corrected for soil NO (Table 4). Increasing the rate of non-stomatal uptake of NO_2 by computing r_{hyd} with $\alpha = 2$ in parameterization P7 resulted in a simulated nocturnal aggregate mean $V_d(NO_2)$ of $0.24 \pm 0.04 \text{ cm s}^{-1}$ which compares well to the observation-inferred $V_d(NO_2)$ of $0.23 \pm 0.35 \text{ cm s}^{-1}$ after correction for soil-emitted NO using a 59 % CRF from P7 (Table 4). Although parameterization P7 results in satisfactory simulation of nocturnal $V_d(NO_2)$ at the HFEMS when averaged across all months, intra- and inter-month variability in observation-inferred $V_d(NO_2)$ is not captured in simulated values (Fig. 3).

Physical justification for a scale factor value $\alpha > 1$ being necessary to reduce the bias between simulated nocturnal $V_d(NO_2)$ and observation-inferred values corrected for estimated soil NO could stem from a larger surface area available for NO_2 heterogeneous hydrolysis in a mature forest environment compared to the U.S. Midwest wintertime agricultural region over which VandenBoer et al. (2013) derived γ_{g,NO_2} . Heterogeneous reactions not limited by transport or diffusion to reaction surfaces are governed by a collision-limited rate which scales linearly with the surface area-to-volume ratio of the reaction vessel or environment (Jacob, 2000). Heterogeneous hydrolysis of NO_2 may proceed on any surface accommodating adsorbed water, including foliar surfaces, bark, or elements of the forest floor (i.e., rock, soil, and debris). Despite the hydrophobic nature of many foliar surfaces, thin aqueous films have been observed on coniferous needles (Altimir et al., 2006; Burkhardt and Eiden, 1994) and stomata-bearing surfaces of deciduous leaves (Burkhardt et al., 1999) at ambient humidities well below saturation. In addition to radiative cooling, elevated humidity within the thin laminar boundary layer surrounding leaves may result from stomatal transpiration (Burkhardt and Hunsche, 2013) and to a lesser extent the hydraulic activation of stomata (HAS) (Burkhardt, 2010), a process discussed further in Section 4. Surface area indices [$\text{m}^2 \text{ m}^{-2}$] for forest components at the HFEMS have been estimated (Fig. S5), including for stems and twigs (STAI = 0.9), coniferous needles (CAI = 0.8), and

deciduous leaves (DLAI = 3.4 summertime maximum). Assuming round stems and twigs (Sörgel et al., 2011) and oblate coniferous needles (Oren et al., 1986; Riederer et al., 1988), the total wintertime canopy surface area is estimated as $\pi\text{STAI} + 2.7\text{CAI} \sim 5 \text{ m}^2 \text{ m}^{-2}$. We estimate the summertime canopy surface area to be $\sim 12 \text{ m}^2 \text{ m}^{-2}$ accounting for both sides of deciduous leaves or $\sim 9 \text{ m}^2 \text{ m}^{-2}$ neglecting the non-stomatous adaxial (top) surface of deciduous leaves, in agreement with typical macroscopic surface area indices for temperate and boreal forest canopies of $12 \text{ m}^2 \text{ m}^{-2}$ (range $5\text{--}14 \text{ m}^2 \text{ m}^{-2}$) (Lammell, 1999). The surface area of the forest floor, including debris, would also be much greater than the planar ground area, and that of tree bark greater than the simple geometric surface area (Section 3.3.4).

The lack of seasonality in observation-inferred nocturnal $V_d(\text{NO}_2)$ depicted in Fig. 3 may reflect an inter-seasonal buffering of available surface area for reaction of above-canopy NO_2 due to increased air parcel mixing throughout the lower canopy in the absence of deciduous leaves (see Section 3.3.4). We did not attempt to parameterize non-stomatal deposition of NO_2 to upper and lower canopy elements separately in our top-down sensitivity analysis of $R_c(\text{NO}_2)$, as is currently the approach in the W89 and Z03 dry deposition schemes. Due to the lack of discernible seasonal variability in observation-inferred nocturnal $V_d(\text{NO}_2)$, above-canopy observations were insufficient to justify the additional variables. We acknowledge that the nocturnal canopy environment is under reduced turbulent mixing compared to daytime conditions when the forest would experience enhanced vertical exchange (Bannister et al., 2023; Sörgel et al., 2011; Thomas and Foken, 2007). Although daytime surface area available to above-canopy deposition is therefore likely to be greater than at night, nighttime sensitivity of $V_d(\text{NO}_2)$ to α is much greater than during the day when stomata are open and foliar uptake of NO_2 is a more substantial pathway to deposition than non-stomatal uptake (Fig. S8). Increasing α from 1 to 2 results in a 100 % increase in simulated $V_d(\text{NO}_2)$ at night (Table 4), but only a 10 % increase during mid-day (Section S5).

The canopy compensation point for NO_2 is the ambient above-canopy concentration at which point consumption (i.e., dry deposition) and production (i.e., soil emission) are in balance (Duyzer et al., 1995). Studies of above-canopy NO_2 exchange have observed aggregate fluxes to be upward (Min et al., 2014; Vaughan et al., 2016), downward (Coe and Gallagher, 1992; Horii et al., 2004; Walton et al., 1997), and not significantly different from zero (Geddes and Murphy, 2014)—highlighting the importance of knowledge of below-canopy NO_x emission and subsequent uptake and reaction prior to attempting interpretation of above-canopy fluxes. Although foliar compensation points for NO_2 —a concentration below which vegetation is proposed to become a net source of NO_2 —have been observed in the past via leaf-level chamber measurements to be generally $< 2 \text{ ppb}$ (Geßler et al., 2002; Sparks et al., 2001; Weber and Renenberg, 1996), recent chamber studies using highly specific NO_2 detection methods have failed to observe foliar emission (Breuninger et al., 2013; Chaparro-Suarez et al., 2011; Delaria et al., 2018, 2020; Wang et al., 2020). By restricting analysis to nocturnal conditions when RH is generally high (Fig. S7), stomata assumed closed (Section 4), and turbulence well established ($u_* > 0.2 \text{ m s}^{-1}$), we find monthly aggregate $V_d(\text{NO}_2)$ relatively constant in the range $0.2\text{--}0.3 \text{ cm s}^{-1}$ across April–November, with expected large variability on finer timescales (i.e., $\sigma = 0.35 \text{ cm s}^{-1}$ across the hourly dataset). Linear regression of hourly nocturnal NO_2 flux versus ambient NO_2 concentration (Fig. 2) yields a $V_{ex}(\text{NO}_2)$ of -0.40 cm s^{-1} ($p < 0.01$) over the entire NO_2 concentration range (up to $\sim 30 \text{ ppb}$) and -0.26 cm s^{-1} ($p < 0.01$) when the four outlying hourly observations beyond 20 ppb NO_2 are excluded—consistent with the findings of Horii

et al. (2004). An inferred $V_d(NO_2)$ of 0.21 cm s^{-1} is obtained after subtraction of $V_{chem} = 0.05 \text{ cm s}^{-1}$ —similar to aggregate values presented in Table 4. The y-axis intercept of $0.11 \text{ ppb cm s}^{-1}$ in Fig. 2, although not significant ($p > 0.1$), is in line with the estimated mean (April–November) above-canopy NO_2 flux of $0.13 \text{ ppb cm s}^{-1}$ resulting from soil NO emission and an average
615 CRF of 59% from parameterization P7. An empirical CRF of $\sim 70 \%$ is obtained from the ratio of y-axis intercept ($0.11 \text{ ppb cm s}^{-1}$, Fig. 2) to seasonal mean below-canopy soil NO flux ($0.39 \text{ ppb cm s}^{-1}$). An NO_2 canopy compensation point for Harvard Forest, likely due to soil NO emission, is approximated by the x-axis intercept of $\sim 0.4 \text{ ppb}$ in Fig. 2.

Given the dependence of r_{hyd} on surface area, land type specific α values evaluated across seasons would be desirable to improve confidence for use in global CTMs. As previously mentioned, Eugster & Hesterberg (1996) inferred a
620 nocturnal non-stomatal median value for $R_c(NO_2)$ of 700 s m^{-1} (range $500\text{--}950 \text{ s m}^{-1}$) over a managed grassland from soil NO corrected eddy covariance observations—similar to the median value of 750 s m^{-1} simulated herein following Eq. (8) with $\alpha = 1$ (Fig. S8). Pilegaard et al. (1998) report nocturnal $R_c(NO_2)$ of $771 \pm 111 \text{ s m}^{-1}$ inferred from eddy covariance observations over a harvested wheat field (with re-growth) in southern Germany during mid-September. Given the high nocturnal NO_2 concentrations of $10\text{--}30 \text{ ppb}$ at the location, soil NO most likely had a reduced relative effect on the resulting $V_d(NO_2)$
625 compared to the large influence noted by Eugster & Hesterberg (1996) where nocturnal NO_2 concentrations were less than 10 ppb during periods of soil NO emission. The effect of soil NO on $V_{ex}(NO_2)$ as a function of above-canopy NO_2 concentration can be seen in Fig. 2, where negligible influence is noted for concentrations above $\sim 5 \text{ ppb}$. Coe et al. (1992) used eddy covariance to estimate a non-stomatal $R_c(NO_2)$ of 548 s m^{-1} over a Heather moorland located in southern Netherlands. Plake et al. (2015) find a maximum median nocturnal bulk $R_c(NO_2)$ over a natural grassland site in Mainz, Germany of 560 s m^{-1} via
630 the dynamic chamber approach. The nocturnal $R_c(NO_2)$ values reported by Coe et al. (1992) and Plake et al. (2015) are intermediate between r_{hyd} values computed here using α of 1 or 2. Assigning $\alpha = 1$ for low roughness vegetative land types appears reasonable but may yield slight underestimates in nocturnal NO_2 uptake.

Reaction R1 has been observed to proceed efficiently on ice surfaces, even at low temperatures ($< 170 \text{ K}$) (Bang et al., 2015; Kim and Kang, 2010). Stocker et al. (1995) observed nocturnal deposition of NO_2 to a snow-covered grassland by
635 eddy covariance in northern Colorado, reporting a median resistance to surface uptake of 740 s m^{-1} , from which we estimate an uptake coefficient to snow $\gamma_{snow} \sim 1.6 \times 10^{-5}$ (Tables 1 & S2)—similar to γ_{g,NO_2} following Eq. (9) at 100% RH. If NO_2 dry deposition persists into winter months at levels observed for late fall in Fig. 3, this represents a significant depositional sink for wintertime NO_2 not currently represented in CTMs when both the lifetime and near-surface concentrations of NO_x are at a maximum.

640 3.3.4 Bottom-up estimates of nocturnal $V_d(NO_2)$

Simple estimates of bottom-up bulk-canopy $V_d(NO_2)$ provide a useful sanity check on top-down eddy covariance inferred values and are a starting point for a mechanistic explanation of bulk-canopy deposition. Bottom-up estimates of nocturnal $R_c(NO_2)$ for Harvard Forest were computed from parallel contributions of uptake to leaves, bark, and the forest floor:

$$R_c(NO_2) = [1/r_c \text{ leaf} + 1/r_c \text{ bark} + 1/(r_a + r_c \text{ floor})]^{-1}, \quad (10)$$

645 where in-canopy aerodynamic resistance r_a was computed according to the Z03 scheme. Component canopy surface resistances in Eq. (10) were computed at hourly resolution following Eq. (8) using surface-specific NO_2 uptake coefficients from Table 1. Component surface area scale factors α are material dependant, varying according to the surface area used to normalize deposition fluxes in corresponding measurement studies (Tables 1 & S2). Attention must be paid to the corresponding surface area for which a particular uptake coefficient is to be applied, and the understanding that application of

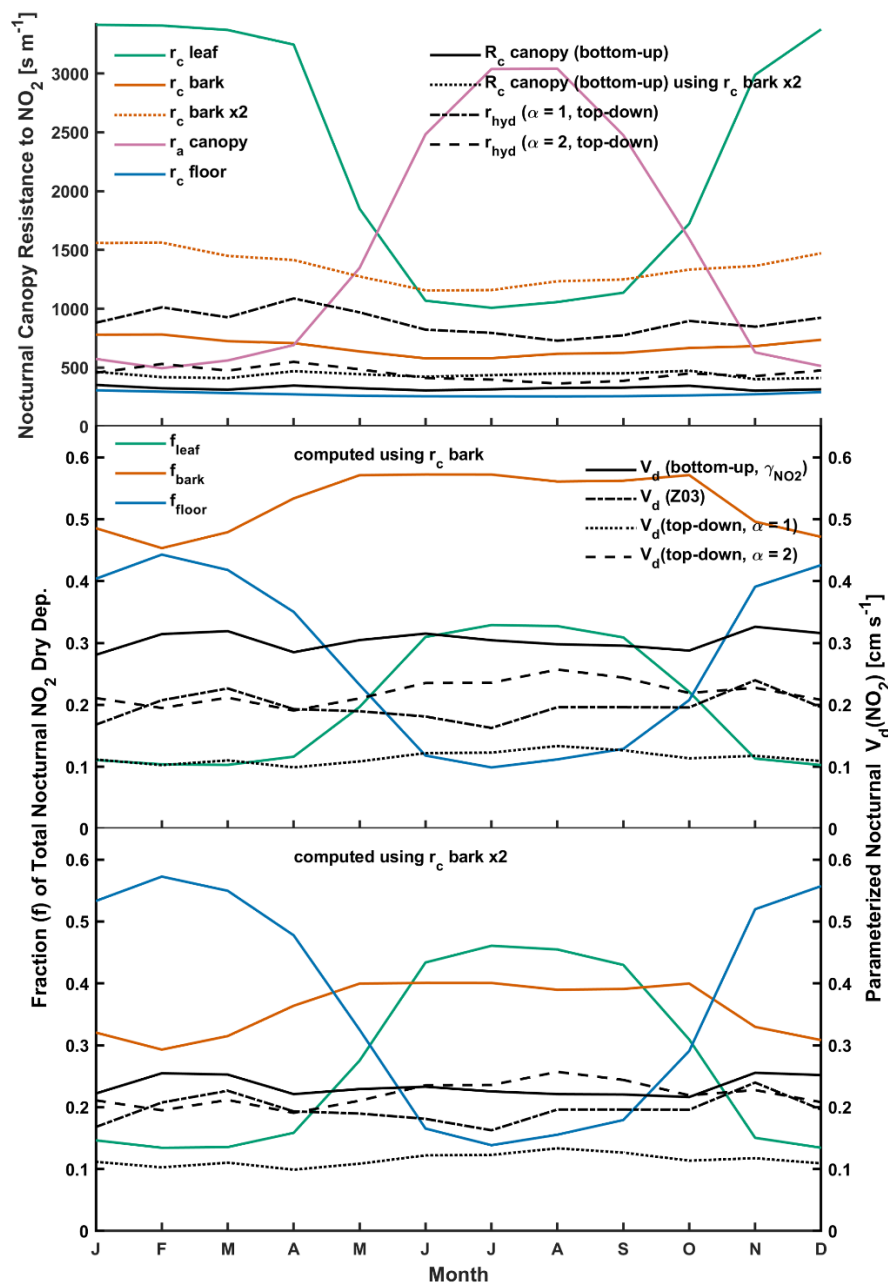
650 an uptake coefficient for a complex surface where planar or simple estimates of geometric surface areas were used during measurement (i.e., forest floor, snow, or bark) assumes surface area equivalence in subsequent applications. As seen in Table 1, uptake of NO_2 to coniferous foliage under conditions of minimal stomatal aperture and $RH < 90\%$ was found to be ~ 2.6 times greater than to deciduous leaves on a projected area basis. In addition to the greater total-to-projected surface area of coniferous (~ 2.7) compared to deciduous (~ 2) leaves, we attribute the reduced non-stomatal uptake of NO_2 to deciduous

655 leaves as a consequence of the hydrophobic adaxial (top) surface of deciduous leaves which lack the elevated water vapor concentrations from stomata to support thin water films for reaction R1 to proceed. We treat the leaves and bark of the forest canopy as wet following the dew flag from the Z03 scheme (Section 2.1.2). Under wet conditions, both top and bottom faces of deciduous leaves would be wetted and we increase the α value used to scale uptake to deciduous leaves from LAI to 2LAI. The α value used to scale uptake to bark is computed as $\pi STAI$, where $STAI = 0.9$ is the projected area of tree branches (Horie et al., 2005). We compute forest floor surface resistances $r_c \text{ floor}$ as parallel contributions from uptake to snow and the snow-free forest floor, weighted by snow cover fraction following the Z03 scheme (Zhang et al., 2003a). As seen in Table 1, NO_2 uptake to wet bark is twice that of dry bark, and uptake to snow is approximately one-third that of the snow-free forest floor.

The top panel of Fig. 4 depicts monthly mean estimates of nocturnal component canopy resistances used in Eq. (10) to compute bottom-up bulk-canopy $R_c(NO_2)$. Also included for comparison are monthly mean estimates of top-down bulk-canopy r_{hyd} for α values of 1 and 2. The middle and bottom panels of Fig. 4 depict parameterized $V_d(NO_2)$ following Eq. (2)

665 using both bottom-up and top-down estimates of bulk-canopy surface resistance. Also included are the fractional contributions that leaf, bark, and forest floor surfaces make to total bottom-up NO_2 uptake. Due to the compensating seasonal contributions of $r_c \text{ leaf}$ and $r_c \text{ floor}$ to total NO_2 deposition, bottom-up nocturnal $V_d(NO_2)$ shows little seasonality, in accordance with eddy covariance inferred $V_d(NO_2)$ from spring through fall (Fig. 3). As depicted in the middle panel of Fig. 4, bottom-up estimates

670 of $V_d(NO_2)$ computed using uptake coefficients from Table 1 are greater than top-down optimized values across all seasons. Computing NO_2 uptake to dry and wet bark using the uptake coefficients from Table 1 of 5.0×10^{-6} and 1.0×10^{-5} , respectively, results in bark surfaces being the predominant nocturnal dry depositional sink for NO_2 at Harvard Forest. Uptake coefficients



675 **Figure 4: (TOP)** Computed estimates of monthly mean component canopy resistances for nocturnal NO₂ uptake to leaves, bark, and the
680 forest floor at Harvard Forest, and resulting ‘bottom-up’ bulk-canopy R_c following Eq. (10). Also included are ‘top-down’ bulk-canopy
resistances r_{hyd} following Eq. (8). **(MIDDLE)** Resulting bottom-up and top-down canopy-scale nocturnal deposition velocities $V_d(NO_2)$,
including fractional contributions of leaf, bark, and forest floor surfaces to bottom-up $V_d(NO_2)$. Also depicted is nocturnal $V_d(NO_2)$ computed
following the Z03 parameterization. **(BOTTOM)** As above (middle), with deposition fractions and bottom-up $V_d(NO_2)$ computed with
uptake to bark reduced by a factor of two. Monthly values of resistances, deposition velocities, and meteorological inputs are included in
Table S3 of the supplement. Surface-specific NO₂ uptake coefficients used to calculate bottom-up R_c are included in Table 1.

to dry and wet bark in Table 1 are from Hanson et al. (1989), where chamber-measured NO₂ uptake was to Teflon end-capped cylindrical branch or trunk samples with diameters of ~ 15 cm. It is expected that the calculated cylindrical geometric surface area used to normalize uptake to exposed bark samples in Hanson et al. (1989) would be less than the surface area available for reaction due to bark roughness and shape complexity for samples of this size. From Table 1, NO₂ uptake to smoother wood surfaces (i.e., wood board or plywood) at RH < 90 % is at least 3 times less than uptake to dry bark, which we speculate could result from a greater available surface area for the bark samples tested. Bark surface area for the trunk and branch samples examined in Hanson et al. (1989) may not be representative of the average bark surface area for the canopy at Harvard Forest. As depicted in the bottom panel of Fig. 4, reducing NO₂ uptake to bark by a factor of 2 (equivalent to a 2-fold increase in r_c bark) results in bottom-up $V_d(NO_2)$ within the range of eddy covariance inferred monthly values corrected for soil NO (0.2–0.3 cm s⁻¹) and within 6 % of top-down $V_d(NO_2)$ for $\alpha = 2$ (parameterization P7) over a 12-month period. At this uptake level, bark is no longer the predominant sink for nocturnal dry deposition of NO₂, instead taking on a secondary role where predominant uptake alternates between canopy foliage during summer months and the forest floor during late fall, winter, and early spring. On an annual basis, nocturnal NO₂ uptake to forest surfaces are within 30 % of one another for the reduced bark uptake case; specifically, the forest floor accounts for 37 % of uptake, bark 36 %, and foliage 27 %. Bottom-up modelling estimates of canopy-scale dry deposition of NO₂ would benefit from future chamber studies detailing NO₂ uptake to a variety of bark surfaces over a range of humidities and temperatures.

Figure 4 also depicts nocturnal $V_d(NO_2)$ computed using $R_c(NO_2)$ parameterized following the Z03 scheme (Table S3). As with top-down and bottom-up estimates of $V_d(NO_2)$, little seasonality is seen across monthly mean values. Over a 12-month period, $V_d(NO_2)$ following Z03 is within 12 % and 17 % of top-down ($\alpha = 2$) and bottom-up (r_c bark x 2) values, respectively. The Z03 scheme formulates non-stomatal uptake of NO₂ by scaling inverse surface resistances optimized for O₃ by 0.8 (Zhang et al., 2002a); herein, we explored the plausibility of non-stomatal NO₂ uptake being a result of R1, with negligible contributions from other reactions. Although additional pathways for NO₂ uptake may exist in parallel to R1 in some instances—particularly in leaf interiors (Section S6 of the supplement)—non-stomatal NO₂ uptake to the natural land types examined and reviewed herein can plausibly be understood as occurring via R1 when surface area effects are considered. Future work investigating the relative roles of R1 and reduction reactions in NO₂ uptake to a variety of natural land type surfaces would aid mechanistic realism in future model developments.

3.4 Evaluation of parameterized $V_d(NO_y)$ over Harvard Forest

Improved representation of V_d for NO_y component species HNO₃ and NO₂ via updates to molecular diffusivity (Section 3.2) and non-stomatal NO₂ uptake (Section 3.3.3), respectively, may be further evaluated at a broader scale through a full NO_y budget analysis. Simulated $V_d(NO_y)$ from base and updated parameterizations is evaluated against observations from the HFEMS, considering NO_y component species NO₂, NO, HNO₃, and PAN. Of particular interest is the period from June–

November 2000, when hourly observations of above-canopy HNO_3 concentration—a significant contributor to NO_y dry deposition at this location (Horii et al., 2005)—was added to the suit of measurements (Fig. S6). The top panel of Fig. 5 depicts the diel climatology of measured NO_y and constituent species over Harvard Forest, alongside inferred NO_y computed as the sum of the component species. The middle panel of Fig. 5 depicts species-specific fractional contributions to measured NO_y . On average, NO_y inferred from the sum of measured component species is $\sim 76\%$ of measured total NO_y , with contributions of 48 % (NO_2), 16 % (HNO_3), 8 % (PAN), and 4 % (NO).

To compare to $V_d(\text{NO}_y)$ inferred from measured fluxes, we compute simulated deposition velocities $V_{d,\text{sim}}(\text{NO}_y)$ from a linear combination of parameterized component deposition velocities $V_d(x_i)$ weighted by species-specific concentration fractions (Michou et al., 2005; Wu et al., 2011):

$$V_{d,\text{sim}}(\text{NO}_y) = \frac{\sum_i [x_i] V_d(x_i)}{\sum_i [x_i]}, \quad (11)$$

Due to the large number of coincident hourly observations required for the comparison, only 19 coincident hourly values of $V_d(\text{NO}_y)$ and $V_{d,\text{sim}}(\text{NO}_y)$ exist across the entire data set consisting of over 2000 hourly measurement points. For this reason, a gap-filling method is used to estimate date (d) and hour (h) specific missing NO_y component concentrations $[x_i]_{d,h}^{\text{infer}}$ for NO_2 , NO, HNO_3 , and PAN from measured NO_y :

$$[x_i]_{d,h}^{\text{infer}} = \left(\frac{[\overline{x_i}]_h^{\text{meas}}}{[\overline{\text{NO}_y}]_h^{\text{meas}}} \right)_{\text{clim}} [\text{NO}_y]_{d,h}^{\text{meas}}, \quad (12)$$

where the diel climatologies of component fractions $[\overline{x_i}]_h^{\text{meas}} / [\overline{\text{NO}_y}]_h^{\text{meas}}$ are depicted in the middle panel of Fig. 5. This method of gap-filling was used by Wu et al. (2011) to evaluate simulated NO_y deposition velocities from the WRF-Chem and NOAA-GEM dry deposition modules against this dataset. Here, we compute component fractions as the ‘ratio of smoothed means’ rather than the ‘mean of ratios’ to reduce the effect of outliers. The fraction of inferred species-specific hourly concentrations required for gap-filling over the study period is depicted in the bottom panel of Fig. 5 as the missing measurement fraction.

Diel profiles of simulated $V_d(x)$ used in Eq. (11) to compute simulated $V_d(\text{NO}_y)$ are depicted in Fig. S4 and reviewed in Section S5 of the supplement. Given the predominant contributions that NO_2 and HNO_3 make to the NO_y budget at Harvard Forest (Fig. 5), updates to the parameterization of $V_d(\text{NO}_2)$ and $V_d(\text{HNO}_3)$ discussed in previous sections will have a notable impact on simulated $V_d(\text{NO}_y)$.

3.4.1 Measurement–model comparison of $V_d(\text{NO}_y)$

Observations of hourly above-canopy eddy covariance fluxes of NO_y at the HFEMS are mostly downward ($> 99\%$; Fig. S9), regardless of adjustment for soil-emitted NO. Figure 6 depicts observation-inferred diel mean $V_d(\text{NO}_y)$ alongside simulated values from parameterizations P2, P4, and P8—selected from Table 2 to highlight the dominant effects of diffusivity updates

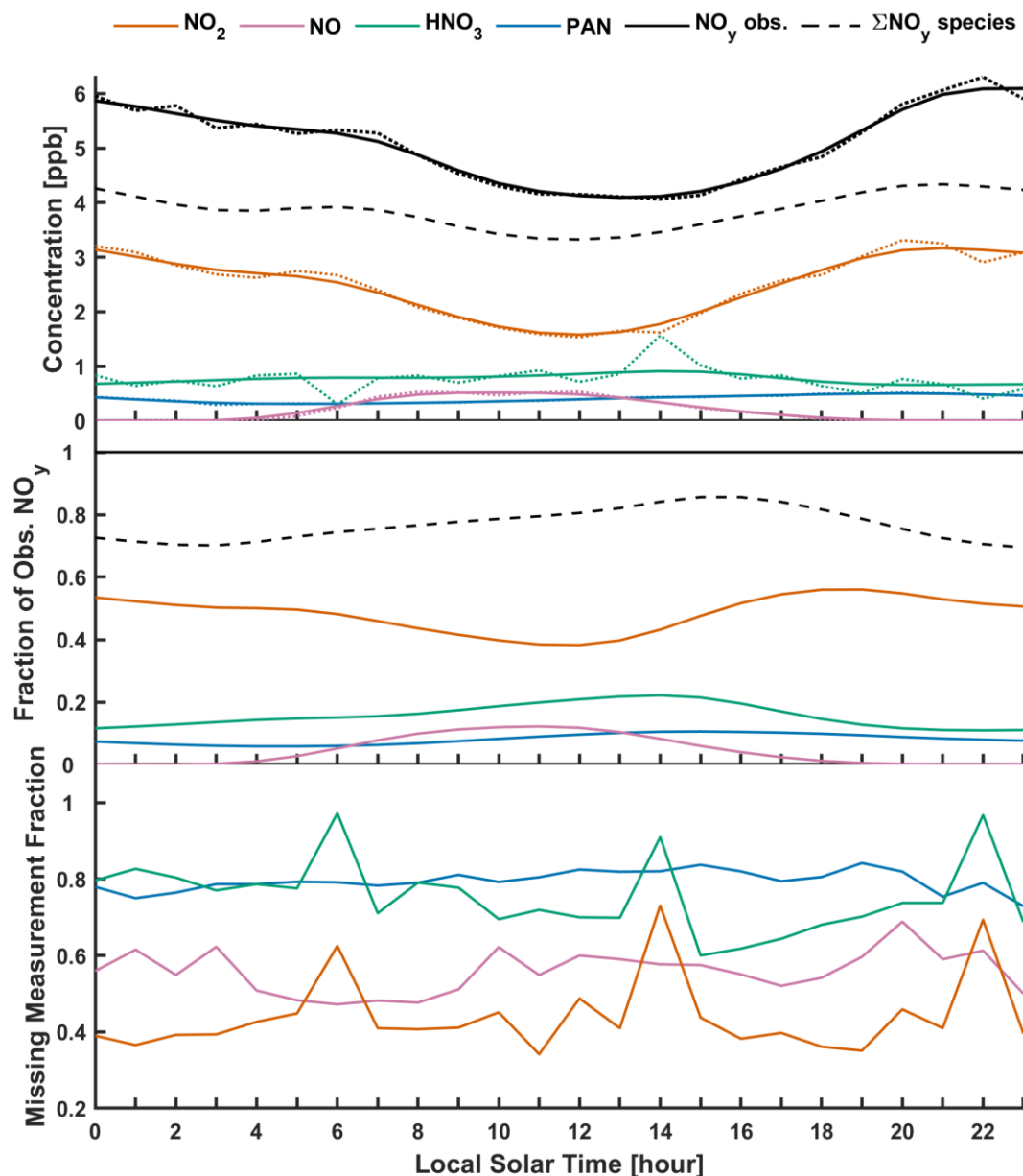


Figure 5: (TOP) Diel climatology of observed NO_y and component species NO_2 , HNO_3 , PAN , and NO measured above Harvard Forest (June–November 2000–2002). Solid lines depict a smoothing spline fit to hourly mean concentrations (dotted). Also shown is the sum of smoothed NO_y component species concentrations. **(MIDDLE)** Fractional contributions of NO_y component species to observed NO_y . These values were computed as ratios of the smoothed diel mean concentrations in the top panel. Shown are individual NO_y species fractions (colored as above) and their sum (dashed black). **(BOTTOM)** Fraction of hourly NO_y component species concentrations (colored as above) missing from the measured NO_y hourly time series spanning June–November 2000. As a gap-filling strategy in the calculated $V_d(\text{NO}_y)$ hourly time series, missing values were inferred using Eq. (12).

(P4) and surface resistance updates to NO₂ and PAN (P8) on simulated $V_d(NO_y)$ referenced from the measurement height at Harvard Forest (P2). Inferred $V_d(NO_y)$ was calculated from eddy covariance observed $V_{ex}(NO_y)$ adjusted for estimates of soil-emitted NO, analogous to $V_d(NO_2)$ in Eq. (6). As evident from Fig. 6, soil NO corrections to observed $V_{ex}(NO_y)$ result in small increases to inferred $V_d(NO_y)$ —9 % for parameterizations P2 and P4, decreasing to 7 % for P8 due to an increase in the simulated canopy reduction factor (Fig. S3) from updates to non-stomatal NO₂ uptake. By far the largest contributor to simulated $V_d(NO_y)$ in Fig. 6 is HNO₃, contributing over 75 % to 24-hour NO_y flux for parameterization P2 (Table 5). Despite nocturnal mean NO₂ concentrations on the order of 3 ppb (Fig. 5), NO₂ makes near-negligible contributions to simulated nocturnal $V_d(NO_y)$ in parameterizations P2 and P4.

Updates to the parameterization of molecular diffusivity (P4) results in large reductions in simulated $V_d(NO_y)$ (-30 % day; -28 % night) and resulting 24-hour depositional flux (26 % reduction, Table 5) due to large reductions in simulated $V_d(HNO_3)$ (Fig. S4). This update exposes a morning peak in observation-inferred $V_d(NO_y)$ in Fig. 6, which simulated values fail to capture. Geddes et al. (2014) measured eddy covariance fluxes of NO_y, NO, and NO₂ above two midlatitude mixed hardwood forests, noting challenges in interpretation of NO_y fluxes between the hours of 07:30–11:00 LST due to suspected canopy storage contributions. Due to the absence of in-canopy NO_y measurements from which to evaluate canopy storage, Geddes et al. (2014) considered observations of above-canopy NO_y flux unrepresentative of the depositional flux over this timeframe and excluded observations during these hours from analysis. The anomalous morning peak in observation-inferred $V_d(NO_y)$ in Fig. 6 could, in part, be due to canopy storage contributions at Harvard Forest. It is conceivable that mixing down of rapidly depositing NO_y species (i.e., HNO₃ or organic nitrates) as growth of the morning boundary layer erodes the residual layer above could induce a downward spike in above-canopy NO_y flux. It is also possible that a morning low bias in simulated $V_d(NO_y)$ results from heavy reliance on the climatological diel profile of HNO₃ (Fig. 5) due to the paucity of hourly HNO₃ observations (Fig. 5 & Fig. S6).

As seen for parameterization P8 in Fig. 6, updates to non-stomatal deposition of NO₂ and PAN result in noticeable increases in simulated $V_d(NO_y)$ (16 % day; 30 % night), with updates to NO₂ deposition having a greater relative impact at night (25 % increase in $V_d(NO_y)$) than during the day (5 % increase in $V_d(NO_y)$). Parameterization P8 updates are associated with an 18 % increase in inferred 24-hour NO_y flux due to large increases in inferred dry deposition of NO₂ (56 % increase) and PAN (85 % increase) (Table 5).

Also depicted in Fig. 6 is the effect on observation-inferred $V_d(NO_y)$ from a maximum estimate of HONO emitted from the heterogeneous hydrolysis of NO₂ on deposition surfaces following reaction R1. Assuming unity for HONO surface emission to the gas phase and subsequent ventilation from the canopy produces a 4 % increase in inferred $V_d(NO_y)$. Uncertainties exist around the nature of the dynamic equilibrium that establishes between evolved and adsorbed/dissolved HONO (Collins et al., 2018; Harrison et al., 1996; Lee, 2012; Spicer et al., 1993; Wojtal et al., 2011) and subsequent implications of a nocturnal reservoir of deposited HONO as a daytime source of HONO to the atmospheric surface layer (He et al., 2006; Ren et al., 2020; VandenBoer et al., 2014, 2015; Wentworth et al., 2016). Lee (2012) monitored near continuous above-canopy NO₂ and HONO concentrations and eddy covariance fluxes at the HFEMS during 2011, finding nocturnal

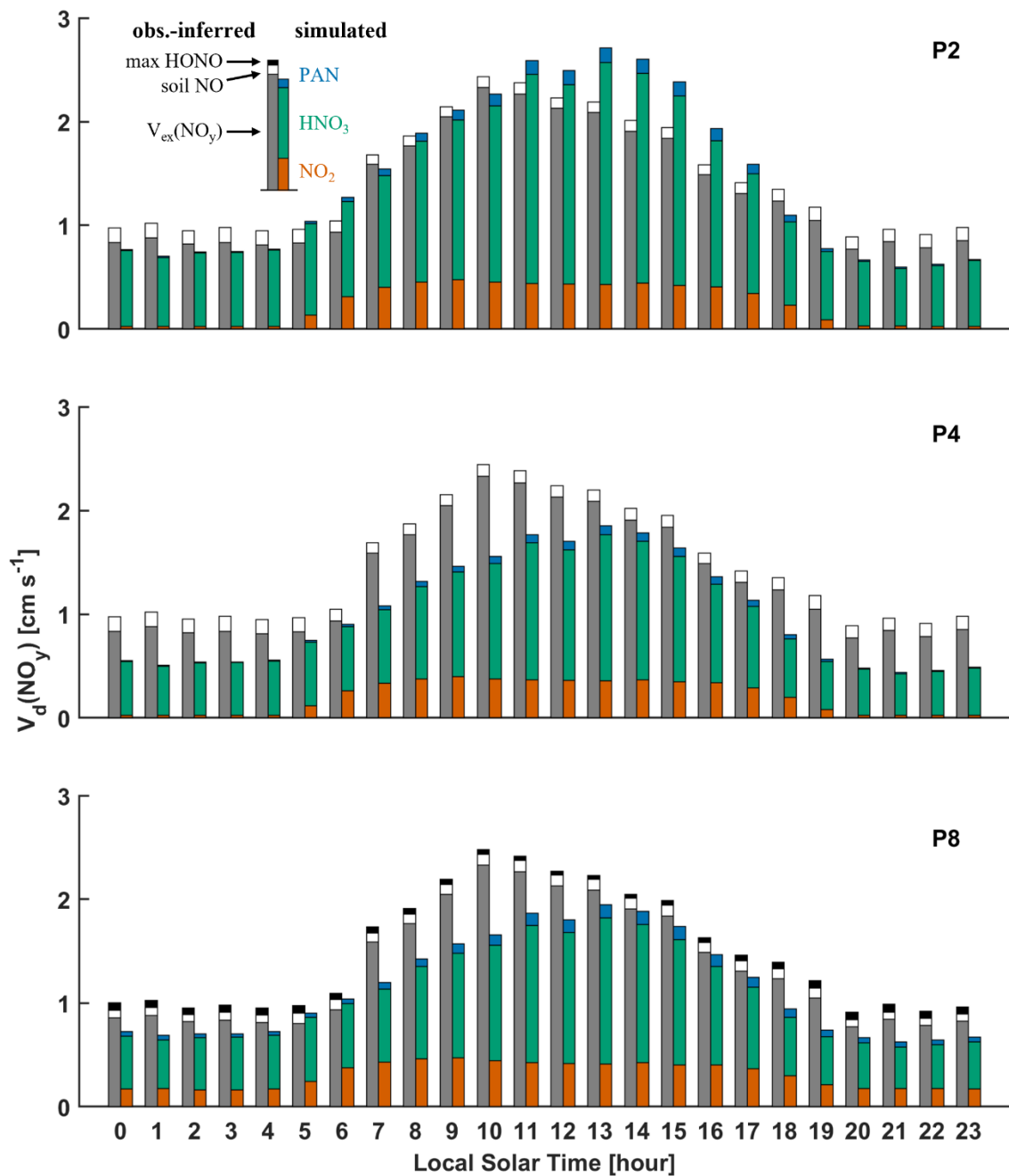


Figure 6: Simulated and observation-inferred diel mean NO_y deposition velocities over Harvard Forest (June–November). In addition to observed $V_{\text{ex}}(\text{NO}_y)$, inferred contributions to $V_d(\text{NO}_y)$ from estimated soil NO emissions (Eq. (7)) and maximum HONO emitted following reaction R1 are also depicted. Simulated $V_d(\text{NO}_y)$ is depicted as the weighted sum (Eq. (11)) of contributing NO_y component species NO_2 , HNO_3 , and PAN for three simulation types: **P2** (base), **P4** (updated R_a and diffusivity D), and **P8** (updated $R_c(\text{NO}_2)$ and $R_c(\text{PAN})$).

enhancements in HONO concomitant with NO₂. However, neither upward nor downward fluxes of HONO were observed, suggestive of establishment of dynamic equilibrium between HONO emission and deposition at Harvard Forest, where perturbation fluxes were below detection limits. Measurements of HONO and NO₂ fluxes over grassland and sugar beet surfaces have highlighted the bidirectional nature of HONO exchange; HONO emission was found to dominate the bidirectional flux under elevated NO₂ concentrations (> 10 ppb for the land types studied) while deposition was noted to be dominant at lower ambient NO₂ concentrations (Harrison et al., 1996; Harrison and Kitto, 1994). The diel flux behaviour of HONO is likely multifactorial, depending on land type, meteorology, and trace gas and particulate concentrations (Pusede et al., 2015; VandenBoer et al., 2015). The absence of significant fluxes of HONO over Harvard Forest as noted by Lee (2012), despite observed downward nocturnal fluxes of NO₂ and nocturnal enhancement of HONO, does not exclude reaction R1 as a source of HONO. Rather, this indicates the importance of deposition and re-emission processes from canopy surfaces which may dominate HONO behaviour at rural forest sites (Ren et al., 2011, 2020; Sörgel et al., 2011; Zhou et al., 2002) in the absence of strong pulses of ambient NO₂ perturbing the dynamic equilibrium between adsorbed/dissolved and gas phase HONO—as is routinely observed in laboratory studies (Finlayson-Pitts et al., 2003; Spicer et al., 1993).

Neglecting the possible effect of emitted HONO on $V_d(NO_y)$ given the findings of Lee (2012), we find a moderate underestimate of 20 % in simulated $V_d(NO_y)$ (24-hour) from parameterization P8, with similar daytime and nighttime biases of -20 % and -19 %, respectively; excluding the period from 07:00–11:00 reduces the daytime bias to -15 % and 24-hour bias to -17 %. As previously mentioned, the NO_y concentration budget at the HFEMS is closed to 76 % on average from observations of NO_x, HNO₃, and PAN. Horii et al. (2005) provide evidence of a rapidly depositing unidentified NO_y species at this site, especially under southwesterly flow, and speculate the unidentified NO_y species as organic nitrates.

Table 5: Inferred mean fluxes of NO_y and component species over Harvard Forest.

Parameterization ^(a)	Flux [ngN m ⁻² s ⁻¹] ^(b)			
	HNO ₃	NO ₂	PAN	NO _y ^(c)
P2 (base)	19 ± 24	4.4 ± 7.1	1.1 ± 1.4	25 ± 30
P3 (<i>R_a</i> update)	18 ± 22	4.3 ± 7.0	1.1 ± 1.5	23 ± 29
P4 (<i>D</i> update)	13 ± 16	3.7 ± 5.9	0.7 ± 0.9	17 ± 21
P7 (<i>R_c</i> (NO ₂), α = 2)	13 ± 16	5.8 ± 7.4	0.7 ± 0.9	19 ± 22
P8 (<i>R_c</i> (PAN))	13 ± 16	5.8 ± 7.4	1.3 ± 1.4	20 ± 23

^(a) See Table 2 for serial parameterization updates. Briefly,
P2: equivalent to base P1 referenced from Harvard Forest measurement height of 29 m.
P3: updates to aerodynamic resistance (*R_a*).
P4: updates to molecular diffusivity (*D*).
P7: update resistance to surface uptake of NO₂ (*R_c*(NO₂)) to include reaction R1 using surface area scale factor α = 2 (Eq.(8)).
P8: update resistance to surface uptake of PAN (*R_c*(PAN)).
^(b) 24 h mean (± 1σ) fluxes inferred from the product of simulated $V_d(x)$ and gap-filled measured concentrations over June–November 2000.
^(c) Inferred NO_y flux from the sum of inferred component (HNO₃, NO₂, and PAN) fluxes.

4 Assumptions of nocturnal stomatal closure & non-stomatal NO₂ uptake via R1

820 Despite longstanding uncertainty regarding nocturnal stomatal behaviour (Caird et al., 2007; Costa et al., 2015; Dawson et al., 2007), it is generally assumed that at night in response to elevated guard cell CO₂ concentration the stomata of C3 and C4 plants are nearly closed (Nobel, 2009), thereby shunting trace gas exchange. In both the widely used W89 and Z03 dry deposition schemes, stomata are assumed to be closed at night and therefore all nocturnal deposition is parameterized through non-stomatal pathways. Although chamber studies consistently find stomatal control dominates daytime foliar uptake of NO₂, 825 uncertainty remains regarding the importance of non-stomatal pathways at night. Some studies note negligible non-stomatal contributions to NO₂ deposition, attributing nocturnal uptake to partially open stomata (Chaparro-Suarez et al., 2011; Delaria et al., 2020; Geßler et al., 2000; Rondón et al., 1993; Sparks et al., 2001), while others find non-stomatal contributions to be non-negligible (Geßler et al., 2002; Hanson et al., 1989; Thoene et al., 1996; Wang et al., 2020; Weber and Renenberg, 1996). This amalgam of contradictory results warrants further consideration and is discussed in the context of a literature review in 830 Section S6 of the supplement.

Notwithstanding nocturnal stomatal behaviour, nocturnal ambient humidity is often elevated at locations with lush vegetation, as it is for summer months at Harvard Forest (monthly median RH 92–95 %; Table S3), whereby growth of aqueous films have been observed to occlude stomatal pores (Grantz et al., 2018); heterogeneous hydrolysis of NO₂ would be expected to proceed on foliar as well as other surfaces under these conditions. Furthermore, nocturnal uptake of NO₂ through a deposition 835 process involving HONO production has been implicated in the field on several occasions (Harrison and Kitto, 1994; Ren et al., 2020; Stutz et al., 2002), including with RH dependence (Stutz et al., 2004; VandenBoer et al., 2013).

Inferred values of γ_{NO_2} in Tables 1 & S2 fall within the range expected for uptake due to NO₂ heterogeneous hydrolysis, i.e., 10⁻⁶ to 10⁻⁵ (Section S3). For foliar surfaces under dark conditions, ‘total leaf area’ normalized uptake was not observed to exceed that of a planar surface of distilled water, supporting the possibility that reaction R1 is the predominant 840 mechanism driving uptake to both the non-foliar and foliar surfaces presented. Three features stand out from tabulated values of γ_{NO_2} in Tables 1 & S2. First, a dependence on surface moisture—dependence on RH or surface wetness is seen for uptake to wood board, concrete, and tree bark. The RH or wetness dependence for NO₂ uptake to wood board and bark is similar to the factor of two increase in γ_{g,NO_2} from Eq. (9) between RH 50% and 100%. Second, surface area available for heterogeneous reaction has direct influence on resulting surface-specific uptake, as expected for a collision-limited heterogeneous process 845 such as reaction R1. Surfaces with complex and undetermined microscopic surface areas (i.e., bark, coarse concrete, forest floors, and snow) exhibit higher V_d^{surf} and resulting γ_{NO_2} —a factor of 2 to 30 greater than to surfaces normalized by accurate predictions of available surface area (i.e., bulk water and foliar). This increased uptake to convoluted surfaces could be an indirect measure of the total microscopic surface area available for reaction, thus highlighting the utility of using field-derived uptake coefficients for parameterizing surface uptake in dry deposition models. Third, a feature stands out between NO₂ uptake 850 to coniferous versus deciduous leaves when stomatal aperture is at a minimum under dark conditions. When uptake is normalized by projected leaf area, inferred γ_{NO_2} to coniferous species is on average 2.6 times greater than to deciduous

species—a factor similar to the ratio of total-to-projected needle leaf area (Riederer et al., 1988). Since coniferous needles have stomata distributed across the entire leaf surface, whereas most deciduous leaves have stomata located on the lower (abaxial) leaf surface and a thicker hydrophobic wax cuticle on the upper (adaxial) leaf surface, the ~ 2.6 -fold greater uptake to coniferous species (when normalized by projected leaf area) may reflect the absence of thin water films on the adaxial surface of deciduous leaves in chamber studies from Table 1. These inferences are consistent with the work of Summer et al. (2004), where similar rates of NO_2 heterogeneous hydrolysis across a variety of hydrophilic and hydrophobic surfaces were understood in the context of available surface areas supporting thin water films. However, given the intra- and interspecies variability in nocturnal V_d^{surf} from leaf-level studies across a range of environmental conditions (i.e., Breuninger et al. (2013), Delaria et al. (2018, 2020), Rondón et al. (1993), and Wang et al. (2020)), further investigation into this simple generalization is warranted.

5 Conclusions

A stand-alone version of the gaseous dry deposition algorithm from GEOS-Chem, implemented to run in single-point-mode, enabled a detailed and more direct evaluation of various branches of the algorithm against eddy covariance inferred deposition velocities over two North American temperate forest ecosystems. Observations of deposition velocities for species that deposit under dynamical control (i.e., nominally small surface uptake resistance R_c) facilitated the identification of a large high bias in computed molecular diffusivities. Correction of this bias using Fuller’s method to calculate diffusivities in the absence of measured values resulted in improved simulation of $V_d(\text{HNO}_3)$. Consequently, this improved representation of simulated $V_d(\text{NO}_y)$ to an extensive dataset spanning many months wherein a potentially anomalous morning peak in observed $V_d(\text{NO}_y)$ was exposed, in agreement with previous work demonstrating challenges in interpretation of late morning NO_y fluxes over a similar forest ecosystem (Geddes and Murphy, 2014). Site-specific roughness length and reference height were found to be important constraints on the calculation of aerodynamic resistance for rapidly depositing species, whereas correction for the influence of the roughness sublayer was found to be of minor importance at the measurement heights involved and of negligible effect at the dry deposition reference height used in GEOS-Chem, in agreement with previous work (Simpson et al., 1998).

A large low bias ($\sim 80\%$) in simulated nocturnal $V_d(\text{NO}_2)$ against eddy covariance inferred values spanning many months over Harvard Forest was identified in accordance with previous work (Horii, 2002; Horii et al., 2004). We addressed this low bias by representing NO_2 heterogeneous hydrolysis—a well-known surface reaction (Finlayson-Pitts et al., 2003) shown to be of importance in the field (VandenBoer et al., 2013), however, yet to be represented in dry deposition parameterizations to our knowledge—in the calculation of non-stomatal surface resistance. A literature review of surface-specific NO_2 deposition velocities to both non-foliar and nocturnal foliar surfaces highlights the importance of considering microscopic surface area for heterogeneous reaction, and enabled estimates of bottom-up $V_d(\text{NO}_2)$ for Harvard Forest which agree well with top-down estimates from eddy covariance inferred values when uptake to bark was reduced by a factor of two—indicating a need for further laboratory study of uptake to bark for representative samples. Consideration of soil NO

emission on eddy covariance inferred $V_d(NO_2)$ at Harvard Forest was found to be important, as was representative canopy surface area when applying NO_2 uptake coefficients from field and laboratory observations.

We persist with the assumption that nocturnal uptake of NO_2 follows non-stomatal pathways, as is currently the case in dry deposition schemes widely used in atmospheric CTMs. Meanwhile, the nocturnal behaviour of stomata remains an active area of research. Confounding processes such as the hydraulic activation of stomata (HAS) and condensation at elevated RH complicate the inference of stomatal conductance to trace gases from observations of water vapor flux, especially in well mixed chambers under dark conditions. It would be helpful for future enclosure studies of NO_2 uptake to consider the effects of heterogeneous hydrolysis of NO_2 on foliar and non-foliar surfaces, as well as potential biases in estimates of stomatal conductance resulting from possible HAS.

Although we find that the Z03 dry deposition scheme adequately captures the magnitude of nocturnal $V_d(NO_2)$ over Harvard Forest, formulating uptake relative to $V_d(O_3)$ neglects contributions from reaction R1, and therefore may be a misrepresentation for NO_2 with implications for surface HONO production in models. We recommend consideration of a non-stomatal dry deposition scheme for NO_2 in atmospheric models that considers the effect of NO_2 hydrolysis on surface resistances. This represents a significant depositional sink for NO_2 under conditions when both the lifetime and near-surface concentration of NO_2 may be elevated, i.e., nocturnal and urban wintertime conditions. We present two approaches that result in general agreement for a mature temperate forest ecosystem. The simplest approach being to represent non-stomatal resistance to NO_2 uptake as r_{hyd} following Eq. (8), with a surface area scale factor $\alpha = 2$ for high surface area land types such as urban and forest, and $\alpha = 1$ for remaining land types. We caution that the value $\alpha = 2$ is not a fit to observations at Harvard Forest, rather, a likely upper estimate from a sensitivity study. Long-term field studies quantifying atmosphere–surface exchange across a variety of land types and seasons would facilitate further development of this unique dry deposition pathway.

Although a main focus of this work was on the effect that NO_2 hydrolysis has on $V_d(NO_2)$, there is much interest in an accurate HONO simulation given that the near-surface nocturnal build-up of HONO results in an early morning burst of OH and NO radicals as HONO photolyzes (Finlayson-Pitts, 2009; Ren et al., 2020), initiating photochemistry prior to other HO_x precursors (Platt et al., 1980). Future work will present the implementation of updates to $V_d(NO_2)$ developed herein into the GEOS-Chem CTM, including analysis of impacts on simulated concentration fields (i.e., NO_2 , HONO, OH, and O_3) and deposition budgets.

Code and data availability. The source code for the dry deposition module from GEOS-Chem version 10-01 is available in a Zenodo repository (<https://zenodo.org/records/13892258>) (Boys, 2024). Harvard Forest data used herein, including trace gas concentrations and fluxes (Munger and Wofsy, 2004, 2023), meteorological observations (Fitzjarrald and Sakai, 2023; Munger and Wofsy, 2024), and LAI (Matthes et al., 2024) are publicly available from the Harvard Forest Data Archive (<https://harvardforest.fas.harvard.edu/data-archive>).

Supplement. The supplement for this article contains additional model and measurement descriptions, review material, and supplemental figures and tables referenced herein.

Author contributions. The manuscript was written using contributions from all authors. BLB designed the study under the supervision and support of RVM. BLB performed the model simulations and data analysis with feedback from RVM. BLB wrote the original draft with feedback from RVM. TCV provided an extensive review, edit, and direction of the original draft which was incorporated into subsequent drafts. All authors have reviewed, edited, and approved the final version of the manuscript.

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