

A zero-dimensional view of atmospheric degradation of levoglucosan (LEVCHEM_v1) using numerical chamber simulations

Loredana G. Suciu¹, Robert J. Griffin², Caroline A. Masiello^{1,3}

¹Department of Earth, Environmental and Planetary Sciences, Rice University, Houston, 77005, USA

²Departments of Civil and Environmental Engineering and Chemical and Biomolecular Engineering, Rice University, Houston, 77005, USA

³Departments of Chemistry and Biosciences, Rice University, Houston, 77005, USA

Correspondence to: Loredana G. Suciu (lgs4@rice.edu)

1 Levoglucosan and the products of its chemical degradation

The chemical structure of levoglucosan (LEV) and its gas-phase and aerosol-phase degradation products (LEVP#) are shown in Figure S1. Products LEVP1, LEVP2, LEVP3 and LEVP5 are generated in both phases (gas and aerosol), product LEVP4 is generated only in the gas phase, and products LEVP6 and LEVP7 are generated only in the aerosol phase. However, because the last three products can partition between phases via either evaporation or condensation, they are also considered degradation products in both the aerosol and gas phase. Henceforth, in their respective chemical mechanism, they are labeled as LEVP#_G or LEVP#_A, in which letters G and A refer to gas-phase and aerosol-phase, respectively. The chemical structures of LEV and LEVP1-LEVP4 are adapted from Bai et al. (2013) and those of LEVP6-LEVP7 are modeled with Chem3D 16.0, following the structures from Kessler et al. (2010). The chemical structure of LEVP5, also created with Chem3D 16.0, is introduced here for the first time based on the similarity of the reaction between an organic compound and N₂O₅ described by Gross et al. (2009).

Intermediary or radical products are shown in Figure S2. The chemical structures of LEVR, LEVRO, LEVRO2 and LEVROOH are adapted from Bai et al. (2013), while that of LEVR1 is re-created using Chem3D 16.0, based on Kessler et al. (2010). Radicals are also common to both phases as they are generated by the phase-specific chemical mechanisms, but they are not considered in the G/P mechanism based on the assumption that they react very quickly once produced.

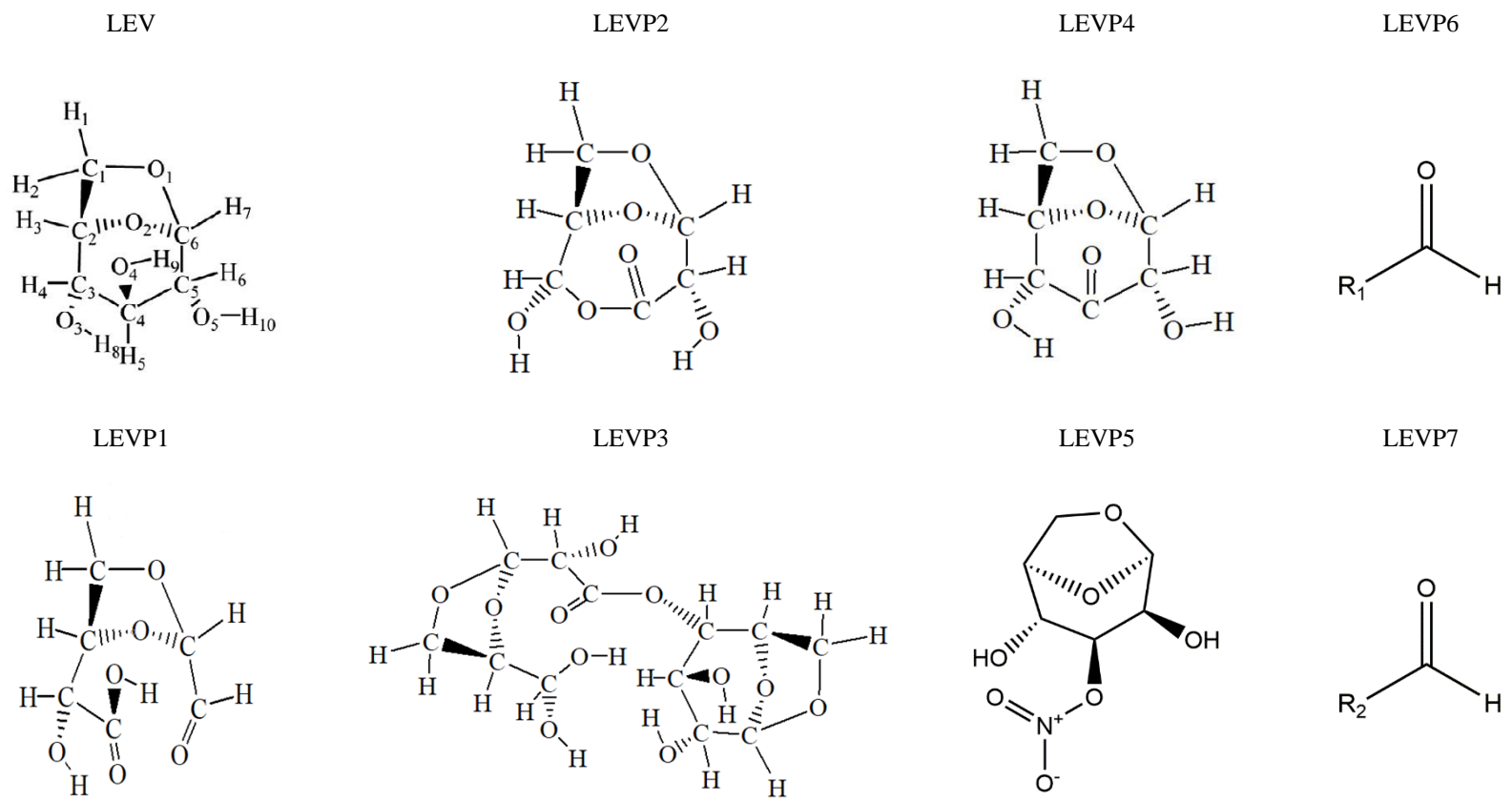
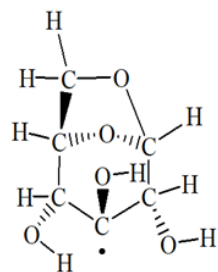
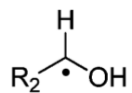


Figure S1: Chemical structures of levoglucosan and its degradation products

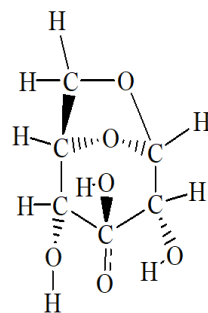
LEVRO



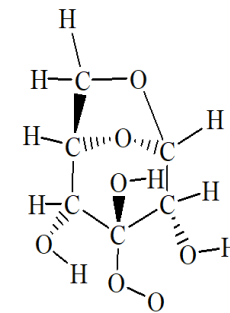
LEVR1



LEVRO



LEVRO2



LEVROOH

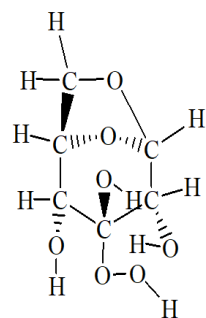


Figure S2: Chemical structures of intermediary (radical) products of levoglucosan degradation

2 Main conditions and parameters used in chamber simulations

The initial conditions of LEV concentration in both phases and main parameters used in the chamber simulations are given in Table S1. Note that the BOXMOXv1.7 model used to develop LEVCHEM_v1 does not explicitly model relative humidity, but it does model H₂O vapor concentration. Therefore, the values of RH used in chamber experiments were converted to H₂O vapor concentration and used in the simulations to roughly reproduce the conditions used in chamber experiments from two studies (Hennigan et al., 2010; Lai et al., 2014). All simulations were conducted at ambient atmospheric pressure (1013.35 hPa). Total aerosol number concentration (N_t) and surface area density (SAD) were taken or calculated based on Seinfeld and Pandis (2006). Enthalpy of vaporization ($\Delta H_{vap,i}$) at 298.15 K was taken from Xie et al. (2014) and applied to all species (both LEV and its products) that were subjected to the gas-particle (G/P) partitioning mechanism. Surface tension (σ) of LEV and related species was assumed to be similar to that of benzene and taken from Seinfeld and Pandis (2006). Saturation concentration of organics at 298 K ($C_i^*(298 K)$) was taken from May et al. (2013). The bulk density of the particle (ρ) was assumed to be the density of LEV. The diffusion coefficient for organic vapors in air (D) was taken from May et al. (2013) and assumed to be the same for all LEV-related species.

Table S1 Conditions used in chamber simulations for model evaluation

Reference study	LEV_G ₀ (ppmv)*	LEV_A ₀ (ppmv)	H ₂ O (ppmv)	RH (%)	T (K)	D _p (m)	N _t (m ⁻³)	SAD (m ⁻¹)	$\Delta H_{vap,i}$ (J mol ⁻¹)	σ (N/m)	C _i [*] (298 K) (kg m ³)	ρ (kg m ³)	D (m ² s ⁻¹)
Hennigan et al. (2010)	9.62x10 ⁻²	6.23x10 ⁻³	2.22x 10 ³	10	293.15	5x10 ⁻⁷	3.18x10 ⁷	2.50x10 ⁻⁵	84x10 ³	2.82x10 ⁻²	3x10 ⁻³	1.69x10 ³	5x10 ⁻⁶
Lai et al. (2014)	9.62x10 ⁻²	7.09x10 ⁻⁴	1.18x10 ⁴	40	298.15	2x10 ⁻⁷	9.96x10 ⁸	1.25x10 ⁻⁴	84x10 ³	2.82x10 ⁻²	3x10 ⁻³	1.69x10 ³	5x10 ⁻⁶
	9.62 x10 ⁻²	7.09x10 ⁻⁴	6.60x10 ³	40	288.15	2x10 ⁻⁷	9.96x10 ⁸	1.25x10 ⁻⁴	84x10 ³	2.82x10 ⁻²	3x10 ⁻³	1.69x10 ³	5x10 ⁻⁶

*The initial gas-phase LEV concentration was determined from its vapor pressure.

3 Evolution of LEV degradation products and their relative importance

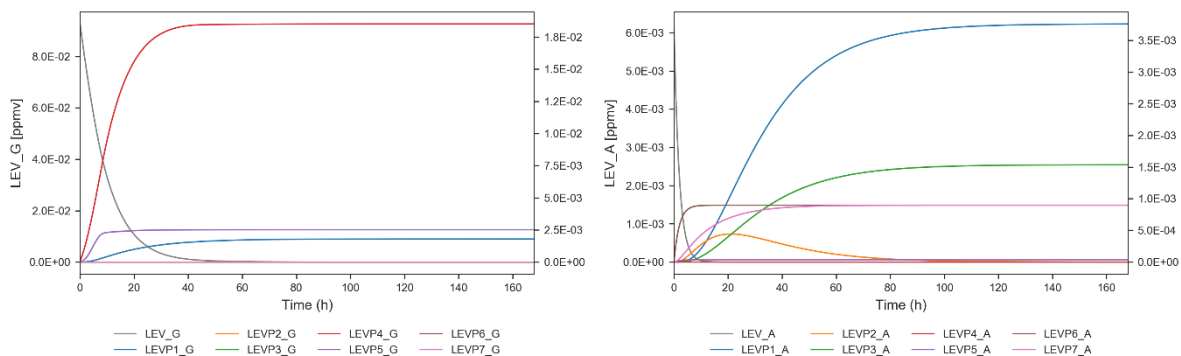


Figure S3: Products of LEV degradation in the gas phase (left) and the aerosol phase (right). ($F = 0.02$; $\alpha = 0.001$) (simulated conditions based on Hennigan et al. (2010)). Concentration of products (ppmv) is displayed on the secondary y-axis.

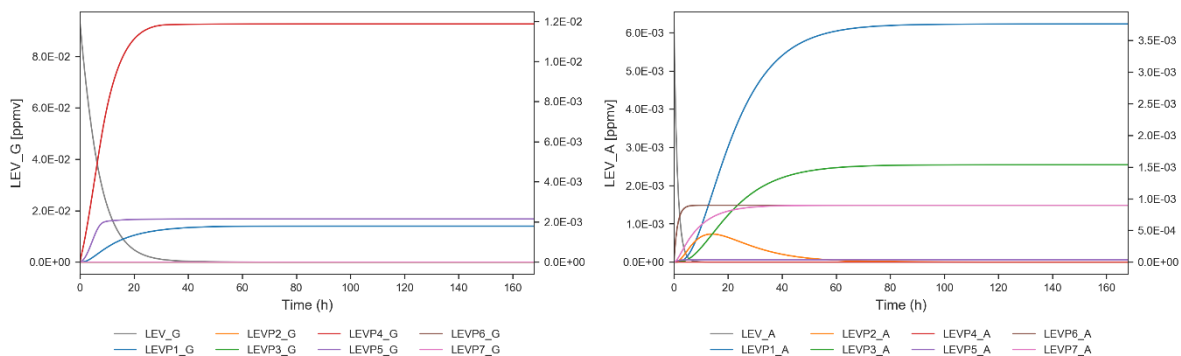


Figure S4: Products of LEV degradation in the gas phase (left) and the aerosol phase (right). ($F = 0.03$; $\alpha = 0.001$) (simulated conditions based on Hennigan et al. (2010)). Concentration of products (ppmv) is displayed on the secondary y-axis.

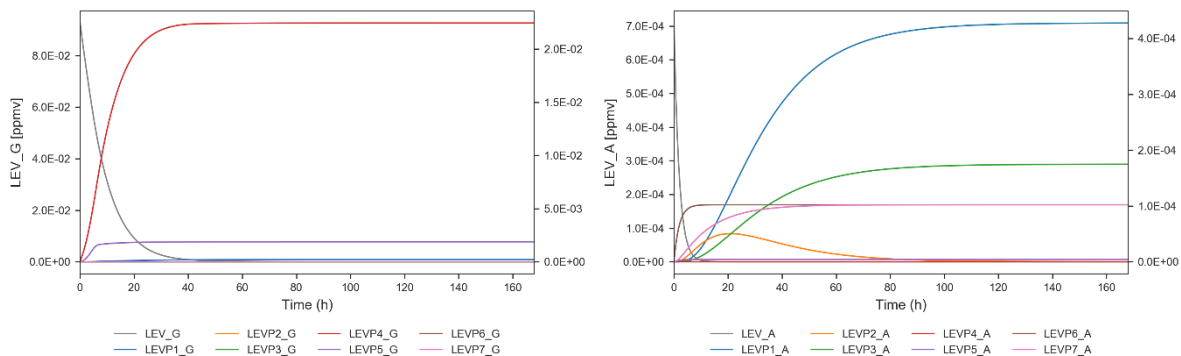


Figure S5: Products of LEV degradation in the gas phase (left) and the aerosol phase (right). ($F = 0.004$; $\alpha = 0.001$) (simulated conditions based on Lai et al. (2014)). Concentration of products (ppmv) is displayed on the secondary y-axis.

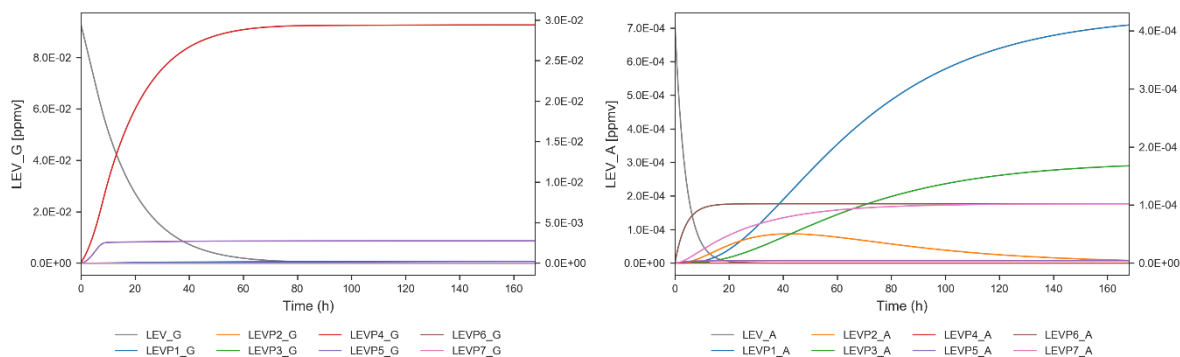


Figure S6: Products of LEV degradation in the gas phase (left) and the aerosol phase (right). ($F = 0.002$; $\alpha = 0.001$) (simulated conditions based on Lai et al. (2014)). Concentration of products (ppmv) is displayed on the secondary y-axis.

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