

**Contribution of
Aitken mode to cloud
droplet populations**

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On the contribution of Aitken mode particles to cloud droplet populations at continental background areas – a parametric sensitivity study

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Abstract

Aitken mode particles are potentially an important source of cloud droplets in continental background areas. In order to find out which physico-chemical properties of Aitken mode particles are most important regarding their cloud-nucleating ability, we applied a global sensitivity method to an adiabatic air parcel model simulating the number of cloud droplets formed on Aitken mode particles, CD_2 . The technique propagates uncertainties in the parameters describing the properties of Aitken mode to CD_2 . The results show that if the Aitken mode particles do not contain molecules that are able to reduce the particle surface tension more than 30% and/or decrease the mass accommodation coefficient of water, α , below 10^{-2} , the chemical composition and modal properties may have roughly an equal importance at low updraft velocities characterized by maximum supersaturations $<0.1\%$. For larger updraft velocities, however, the particle size distribution is clearly more important than the chemical composition. In general, CD_2 exhibits largest sensitivity to the particle number concentration, followed by the particle size. Also the shape of the particle mode, characterized by the geometric standard deviation (GSD), can be as important as the mode mean size at low updraft velocities. Finally, the performed sensitivity analysis revealed also that the chemistry may dominate the total sensitivity of CD_2 to the considered parameters if: 1) the value of α varies at least one order of magnitude more than what is expected for pure water surfaces ($10^{-2}-1$), or 2) the particle surface tension varies more than roughly 30% under conditions close to reaching supersaturation.

1 Introduction

One of the main sources of uncertainty in current predictions concerning the climate change arises from large difficulties in predicting reliably the microphysical structure of clouds, in particular the number concentration and size of cloud droplets (Menon, 2004; Chen and Penner, 2005). Since atmospheric aerosol particles act as nuclei onto which

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cloud droplets are formed, these uncertainties are closely tied to our incomplete knowledge regarding the sources and physico-chemical properties of atmospheric aerosols (Lohmann and Feichter, 2005; McFiggans et al., 2006).

Aerosol particles need to contain sufficient amounts of water-soluble material in order to form cloud droplets in the atmosphere. The minimum particle diameter required for acting as cloud condensation nuclei in the atmosphere is determined by complex interactions between cloud dynamics and aerosol particle population, but varies typically between 50 and 100 nm in the lower troposphere (Seinfeld and Pandis, 1998). This size range is also characteristic for the Aitken mode particles, the physico-chemical properties of which depend strongly on the aerosol origin. Given that the Aitken mode particles often make dominant contribution to the total particle number concentration in the size range >50 nm in continental areas (Tunved et al., 2003), it is therefore highly desirable to understand the connection between the Aitken mode particles and the cloud microphysics in these areas.

The climatic effects of Aitken mode particles, formed either in the atmosphere or emitted from surface-based sources, can be quantified with regional and/or global models (e.g. Adams and Seinfeld, 2002; Spracklen et al., 2005; Stier et al., 2005). Due to the various spatial scales involved, microphysical processes have to be described in a computationally efficient way while simultaneously maintaining a sufficient level of accuracy in such models. To make an optimal compromise between computational costs and realism, the key parameters governing the climatic effects of Aitken mode particles should be identified, and most of the effort should be devoted to capturing accurately the time development of these parameters. To this end, our aim is to give an answer to the following question: “which physico-chemical properties of Aitken mode particles are most important regarding their contribution to cloud droplet number concentrations?”. Providing an answer to this question would help us to prioritize the research needs also in the field of experimental aerosol research.

We approach the problem by investigating the sensitivity of the number concentration of cloud droplets formed on Aitken mode particles to the physico-chemical properties

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of these particles. The approach relies on performing model calculations with an adiabatic air parcel model and analyzing the model output with the probabilistic collocation method (PCM), a tool for sensitivity analysis (Tatang et al., 1997). Here we would like to note that studies adopting somewhat similar approaches have been conducted previously (e.g. Feingold, 2003; Rissmann et al., 2004; Ervens et al., 2005 and references therein; Chuang, 2006). However, the approach of the current paper is novel in two ways. First, the focus is solely on the impact of Aitken mode particles to cloud microphysics and not that of the whole particle population. This choice is largely motivated by the current research interest in new particle formation and its climatic implications (Kurten et al., 2003; Kulmala et al., 2004; Kerminen et al., 2005; Spracklen et al., 2006). Second, we employ a so-called “global” method for sensitivity analysis, in contrast to “local” methods used in the above-cited studies. There are dedicated papers discussing the differences between these two approaches (e.g. Saltelli, 1999a, Saltelli et al., 1999b), and thus it suffices to point out two major advantages of the “global” method over the “local” one: 1) the model sensitivity to uncertain input parameters is quantified over the whole parameter space or over a parameter space region, and 2) the net effects of simultaneously varying input parameters are accounted for. Furthermore, we focus on conditions typical to continental background air masses. New particle formation takes place regularly under such conditions and the newly-formed particles, after their growth to Aitken mode sizes, are able to contribute to the cloud droplet concentrations (Komppula et al., 2005 and references therein; Kerminen et al., 2005). Consequently, the results of the study are directly relevant to understanding the climatic effects of new particle formation taking place over large parts of the globe.

2 Approach

We approach the problem by a combination of model simulations performed with an adiabatic air parcel model (AAPM, Anttila and Kerminen, 2002) and sensitivity analysis using the probabilistic collocation method (PCM, Tatang et al., 1997). The AAPM

is used to predict the number concentration of cloud droplets formed on Aitken mode particles, CD_2 , during an air updraft. Here input parameters describing the physico-chemical properties of Aitken mode particles are treated as independent random variables.

5 The PCM is a technique that quantifies the sensitivity of the model output to uncertainties in input parameters. In order to save computing time compared with a full Monte Carlo analysis, the model output is approximated by polynomials termed as polynomial chaos expansions (PCEs), the terms of which are functions of the uncertain parameters. Free coefficients in the PCEs are determined so that the PCEs give
10 an optimal approximation for the true model output in the high probability regions of the parameter space. The required statistical properties of the model output can be readily extracted from the PCEs, allowing for a global characterization of the model sensitivity to uncertainties in model input parameters.

2.1 Air parcel model

15 The applied AAPM has been described in detail by Anttila and Kerminen (2002). Briefly, the model solves equations governing the time development of a population of aerosol particles and cloud droplets in an air parcel that rises adiabatically with a constant velocity. The particle size distribution is assumed to consist of an Aitken and accumulation mode, and particles are divided into two separate grids with 100 size bins
20 each according to their mode. This allows for a straightforward determination of CD_2 at the cloud top. The thermodynamic driving force behind the growth or evaporation of a particle/droplet during an air updraft is the difference ($S - S_{eq}$), where S is the saturation ratio of water vapour and S_{eq} is the equilibrium saturation ratio of water over the particle/droplet surface. The former quantity, S , is determined by a balance between
25 the cooling of the air parcel and transfer of water vapour onto particles and droplets. The quantity S_{eq} , in turn, is calculated using the Köhler equation (Seinfeld and Pandis, 1998) and is a function of the particle/droplet size and its chemical composition. To be more specific, S_{eq} depends on the particle/droplet surface tension, solution non-

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idealities and the number of molecules dissolved into the aqueous phase. For a particle with a given mass, the last of these quantities depends on the density and molecular weight of the solute molecules as well as on their tendency to dissociate in the aqueous phase.

5 Following previous modeling studies utilizing an adiabatic air parcel model, we quantify the uncertainties arising from the particle chemical composition by assuming that particles contain only a single solute (Feingold, 2003; Ervens et al., 2005). It is also assumed that the solute dissolves entirely into the aqueous phase. Slightly-soluble compounds, which dissolve only partially, are not considered because the effects of
10 limited solubility are clearly exceeded by those of varying soluble mass fraction (Ervens et al., 2005, McFiggans et al., 2006). Furthermore, our model does not account for the surface/bulk partitioning of the solute which may influence the cloud-nucleating ability of particles (Sorjamaa et al., 2004; Kokkola et al., 2006; Sorjamaa and Laaksonen, 2006). This is because the particle/droplet surface tension is assumed to be
15 constant in the model (Sect. 2.3), whereas the surfactant partitioning can be calculated only if the surface tension is allowed to depend on the solute concentration (Sorjamaa and Laaksonen, 2006).

2.2 Application of PCM

Comprehensive descriptions of PCM can be found in the literature (Tatang et al., 1997; Isukapalli, 1999; Lucas and Prinn, 2005), and therefore only a brief outline is given
20 here.

In our application, the goal is to find a PCE that approximates $\ln(CD_2)$. The natural logarithm of CD_2 is approximated rather than CD_2 in order to avoid unphysical predictions that might arise when approximating CD_2 with polynomials. We treat all uncertain
25 input variables or, if the uncertainty range is more than one order than magnitude, their logarithms as uniformly distributed random variables ϕ_i , $i=1, \dots, N$, where N is the number of the uncertain parameters. Furthermore, in order to simplify calculations, we re-scale the variables ϕ_i so that they are all distributed uniformly in the range $[-1, 1]$

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and consequently a new set of random variables ψ_i is obtained:

$$\psi_i = \frac{2\phi_i - (b_i + a_i)}{(b_i - a_i)}, \quad (1)$$

where $[a_i, b_i]$ is the value range of ϕ_i .

The PCEs generated here consist of a sum of orthogonal polynomials which are functions of ψ_i and depend on the probability distributions of ψ_i . For random variables distributed uniformly in the range $[-1, 1]$, the corresponding orthogonal polynomials are Legendre polynomials (Table 2). The accuracy of a polynomial approximation increases generally with increasing order of the polynomials, but consequently also the computing time increases. Hence, in order to keep the computational burden reasonable while maximizing the accuracy, PCEs used here are of fourth order with respect to homogeneous terms and of third order with respect to cross terms. Ternary or higher order products of ψ_i are not considered. Consequently, the generated PCEs for $\ln(CD_2)$ have the following form:

$$\Gamma = \alpha_0 + \sum_{j=1}^4 \sum_{k=1}^N \alpha_{j,k} P_j(\psi_k) + \sum_{k=1}^{N-1} \sum_{j=k+1}^N \beta_{j,k} P_1(\psi_k) P_1(\psi_j) + \sum_{k=1}^N \sum_{j=1, j \neq k}^N \gamma_{j,k} P_1(\psi_k) P_2(\psi_j). \quad (2)$$

Here α_0 and $\alpha_{j,k}$ are coefficients related to the homogeneous part of the PCE, P_j is j -th order Legendre polynomial (see Table 2), and $\beta_{j,k}$ and $\gamma_{j,k}$ are coefficients related to the heterogeneous part of the PCE. It should be noted that PCEs can be also used to parameterize model output (e.g. Calbó et al., 1998; Mayer et al., 2000), but here we use PCM solely as a tool for sensitivity analysis.

The coefficients α_0 , $\alpha_{j,k}$, $\beta_{j,k}$ and $\gamma_{j,k}$ are determined as follows. First CD_2 is calculated using the AAPM in certain points of the parameter space. In our case, these so-called collocation points are formed from the roots of fifth order Legendre polynomials (Tatang et al., 1997). In the standard formulation of PCM, the number of model runs used to determine coefficients in (2) is equal to the number coefficients (Tatang et al., 1997). In order to increase the accuracy of the PCEs, we performed twice that

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many simulations (Isukapalli, 1999). By substituting the model-generated output to the left-hand side of (2) and the corresponding input parameter values to the right-hand side of (2), we obtain a set of linear equations for the coefficients α_0 , $\alpha_{j,k}$, $\beta_{j,k}$ and $\gamma_{j,k}$. The system is solved using the singular value decomposition technique which yields also an optimal agreement between the model and PCE in the collocation points in the least-squares sense (Press et al., 1992).

Γ is a random variable which approximates the original model output, and therefore several useful statistical properties describing the model behavior can be extracted from (2) once the coefficients have been determined. The subsequent results are based on calculating the following integrals:

$$E(P_n^j) = \frac{1}{2} \int_{-1}^1 P_n^j(x) dx, \quad (3)$$

where $E(P_n^j)$ is the expectation value of n-th order Legendre polynomial which is raised to the j-th power. The values of the required integrals are shown in Table 2. By using the independency of the variables ψ_j and the orthogonality of Legendre polynomials, the expected value, $E(\Gamma)$, and variance, $Var(\Gamma)$, readily follow:

$$E(\Gamma) = \alpha_0, \quad (4)$$

$$Var(\Gamma) = \sum_{j=1}^N \left(\frac{1}{3}\alpha_{1,j}^2 + \frac{1}{5}\alpha_{2,j}^2 + \frac{1}{7}\alpha_{3,j}^2 + \frac{1}{9}\alpha_{4,j}^2 \right) + \frac{1}{9} \sum_{k=1}^{N-1} \sum_{j=k+1}^N \beta_{j,k}^2 + \frac{1}{15} \sum_{k=1}^N \sum_{j=1, j \neq k}^N \gamma_{j,k}^2.$$

As seen, the expression for $Var(\Gamma)$ includes a summation over N , which provides a means to decompose the total variance into the contributions from each variable ψ_j :

$$Var(\psi_j) = \frac{1}{3}\alpha_{1,j}^2 + \frac{1}{5}\alpha_{2,j}^2 + \frac{1}{7}\alpha_{3,j}^2 + \frac{1}{9}\alpha_{4,j}^2 + \frac{1}{9} \sum_{j=1, j \neq i}^N \frac{\beta_{j,i}^2}{2} + \frac{1}{15} \sum_{j=1, j \neq i}^N \left(\frac{\gamma_{j,i}^2}{2} + \frac{\gamma_{i,j}^2}{2} \right). \quad (5)$$

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The factor $1/2$ appearing in the last two terms in the right-hand side of (2) is due to the fact that ψ_i and ψ_j have the same probability distribution, which implies that the cross-terms contributions distribute evenly between these variables. Expression (5) is of central importance to the further considerations, since it provides a measure to the contribution of each uncertain model parameter ϕ_j to the total variance in the model output.

2.3 Performed sensitivity studies

Here the focus is on uncertainties arising from the properties of Aitken mode particles, not on those related to background particles. Therefore we assumed that the background particles comprise a single mode with constant properties characteristic to continental remote areas. Thus, the background mode particles consist of ammonium bisulfate and have mode mean diameter of 200 nm, total number concentration of 250 cm^{-3} and geometric standard deviation of 1.45. The surface tension of background particles and droplets formed on them is assumed to be equal to that of pure water. In addition, the mass accommodation coefficient of water onto background particles and droplets formed on them is set equal to unity.

The updraft velocity of an air parcel, V , is a crucial model parameter which determines, together with the aerosol population, the number concentration of cloud droplets formed during an air ascent. In order to explore a range of supersaturations and to improve the accuracy of PCEs, the PCEs were generated separately for each applied value of V which were 0.2, 0.4, 0.6, 0.8 and 1.0 m/s. We also generated PCEs for lower updraft velocities, but the accuracy of the PCEs were notably worse compared to results obtained for $V=0.2$ m/s. The reason for this is discussed in detail in Sect. 3. However, we emphasize that the choice does not limit the validity of our conclusions, since the obtained results show a coherent behavior that can be extrapolated to smaller updraft velocities.

Uncertain parameters describing the physico-chemical properties of the Aitken mode particles are shown Table 1. Our focus is on particles that are able to act as CCN at

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supersaturations characteristic for clouds formed in continental background areas, and we have utilized available empirical data in choosing the value ranges over which these parameters vary. Attributing specific probabilities to different parameter values is not, however, possible at the present due to large gaps in the current knowledge regarding the statistical distribution of the properties of atmospheric Aitken mode particles. Rather, only a value range can be ascertained with confidence. Accordingly, it was assumed that all the uncertain parameters are distributed uniformly, excluding CN and α of which values span several orders of magnitude. In order to better account for the larger uncertainties, it was assumed that the logarithms of CN and α have uniform distributions.

The parameters listed in Table 1 can be divided into two groups: those describing the modal properties of Aitken mode particles (first three parameters) and those related to their chemical composition (last six parameters). The parameters belonging to first group are the number concentration of Aitken mode particles, CN , and the mean size and geometric standard deviation of the Aitken mode, D_m and σ_g , respectively. The value ranges of these parameters are chosen according on the particle size distribution measurements conducted in continental background areas (e.g. Tunved et al., 2003) and represent thus observationally constrained value ranges. Here we would like to point out that one factor causing uncertainty is the mixing state of Aitken mode particles (Rissman et al., 2006). The mixing state, however, is a qualitative concept and as such cannot be incorporated into the current framework. Therefore it is assumed that particles are internally mixed, but we acknowledge the fact that the effect of the particle mixing state is not captured by our approach.

The uncertain parameters related to the particle chemical composition include the solubility, density and number-averaged molecular weight of the matter comprising the Aitken mode particles, ε , MW_{avg} and ρ , respectively. These parameters include also the so-called “effective” Van’t Hoff factor, $\nu\Phi$, which is the product of the osmotic coefficient of the solute, Φ , and the number of ions resulting from dissociation of a solute molecule in the aqueous phase, ν . The remaining two parameters in this group are

the surface tension of the Aitken mode particles and droplets formed on them, σ_s , and the mass accommodation coefficient of water onto the surfaces of the Aitken mode particles and resulting droplets, α .

A proper choice of the value ranges of the last six parameters listed in Table 1 is problematic due to the organic aerosol component which is not completely characterized at the present. Therefore it was decided to make several sensitivity studies with the same approach but using different value ranges for the most poorly-constrained parameters. The first sensitivity study can be viewed conservative because the full variability in the chemical composition of atmospheric aerosols is not accounted for. We term this scenario as “BASE”. Since this scenario may underestimate the importance of the particle chemical composition, we performed two additional sets of sensitivity studies, called “MACRO” and “FILM”. In these scenarios, we adopted larger value ranges for the most uncertain parameters.

The “BASE” scenario is based on the following assumptions: particles do not contain 1) macromolecules having a large molecular weight (>250 g/mol), 2) compounds having more than two carboxylic groups, 3) surface-active compounds that decrease the value of σ_s more than roughly 30%, or 4) surface-active compounds that are able to form a thick film onto the particle/droplet surface and thereby reduce the value of α below the range that has been reported for pure water surfaces (0.01–1.0, see Laaksonen et al., 2005, and references therein). Examples of organic compounds that meet these criteria are various alcohols, polyols, ketones, aldehydes and acids containing one or two functional groups (Saxena and Hildemann, 1996). The maximum allowed reduction in σ_s is consistent with surface tension measurements of atmospherically relevant organics at relatively dilute solutions (Shulman et al., 1996; Facchini et al., 2000; Tuckermann and Cammenga, 2004; Hyvärinen et al., 2006; Salma et al., 2006; Sveningsson et al., 2006). It should be noted that surface tension is a dynamic parameter which depends on the particle size and relative humidity (Ervens et al., 2005; Asa-Awuku and Nenes, 2006; Dinar et al., 2006). We have assumed, however, that the value of σ_s is constant during a model run, i.e. it does not depend on the parti-

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cle/droplet size or its composition. This allows for assessing unambiguously the importance of σ_s in the considered value range. It should be further noted that the soluble mass fraction, ε , in atmospheric Aitken mode particles is poorly constrained as well, and indirect measurements on the chemical composition of sub-100 nm particles suggest that ε can be even lower than the minimum value chosen here, 0.2 (Sveningsson et al., 1997; Ehn et al., 2007). However, given that the model solute compound in the cited studies was ammonium sulfate, soluble fractions <0.2 translate to critical supersaturations $>0.5\%$ for Aitken-mode sized particles. The range is clearly higher than the maximum supersaturations reached at continental environments with updraft velocities <1.0 m/s (Seinfeld and Pandis, 1998), and therefore such particles cannot be regarded as “potential” CCNs under conditions relevant to this study. Regarding the particle dry density, ρ , the range chosen here spans the range expected for atmospheric aerosols. Taken together, the discussed choices limit the value ranges of MW_{avg} , ρ , $\nu\Phi$, σ_s and α to those shown in Table 1.

The second sensitivity study, the “MACRO” scenario, differs from the “BASE” scenario by larger value ranges applied for the parameters MW_{avg} , $\nu\Phi$, σ_s and α (Table 1). The larger value ranges reflect the findings that atmospheric aerosols, including Aitken mode particles, may contain polyfunctional compounds with large molecular weights and ability to act as effective surfactants (Facchini et al., 1999, Graber and Rudich, 2006 and references therein). The upper limit for MW_{avg} , 600 g/mol, is chosen according to experimental information on the number-averaged molecular weight of humic-like substances (HULIS) extracted from atmospheric aerosols (Dinar et al., 2006). Although several studies indicate that larger macromolecules with MWs over 1000 g/mol can be present in aerosols (Graber and Rudich, 2006 and references therein), we use a smaller limit for the following reasons. First, the studies reporting the large MWs are mainly smog-chamber experiments in which conditions may not be entirely representative to those in the atmosphere (Graber and Rudich, 2006). Second, the available evidence seems to suggest that the MW distributions have a maximum value below 600 g/mol in these experiments (Graber and Rudich, 2006). Since we assume only a

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single solute in the AAPM, the use of larger solute MWs may thus grossly overestimate the atmospherically realistic range of number-averaged MW, the fundamental quantity here (Dinar et al., 2006), and may therefore also overestimate the uncertainties arising from MW. For these reasons, the value of MW_{avg} is not varied by more than one order of magnitude. The maximum value of $\nu\Phi$, in turn, is based on the properties of a standard Fulvic acid compound that has been used in several studies as a model compound for aerosol-bound polyfunctional compounds (Mircea et al., 2002; Nenes et al., 2002). Finally, the lower limit for σ_s , 0.02 N/m, is chosen according to the estimate of Ervens et al. (2005) for the maximum reduction of the droplet surface tension due to the presence of organics at relative humidities close to 100%.

The third scenario, "FILM", differs from the "BASE" scenario only by the larger value range of α (Table 1). The minimum values adopted here are based on available experimental evidence suggesting that the mass accommodation coefficient of water on atmospheric aerosols can be as low as 10^{-5} (Chuang, 2003, and references therein). The possible value range spans thus five orders of magnitude. However, in the discussed scenario, "FILM", the minimum value of α was decreased only down to 10^{-3} , and the reason for not using smaller values is discussed in Sect. 3.2.3.

2.4 Validation of the method

The validity of our approach was evaluated as follows. For each PCE generated, we performed 750 additional AAPM simulations, in which the uncertain input parameters were varied randomly according to their probability distributions. The results were compared with the corresponding predictions of the PCEs. Based on the AAPM calculations, probability density functions (PDFs) for CD_2 were then constructed, and these PDFs were compared with PDFs obtained by sampling from the corresponding PCEs. Finally, the PCE and model-based expected values and variances of $\ln(CD_2)$ were compared.

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3 Results

3.1 Performance of the PCEs

Figure 1 shows a comparison of the true model output and corresponding PCE-based predictions for $\ln(CD_2)$. Further, the corresponding coefficients of determinations, R^2 , are displayed in Table 3. The comparison has been made for $\ln(CD_2)$ and not for CD_2 , since $\ln(CD_2)$ was the approximated model output. Results are shown for all the three scenarios and for three updraft velocities ($V=0.2, 0.4$ and 1.0 m/s). The average maximum supersaturations in the AAPM calculations for these three updraft velocities were around 0.12, 0.16, and 0.28, respectively, regardless of the scenario. These values compare favorably with estimated supersaturations reached in continental clouds (Pruppacher and Klett, 1997; Cantrell et al., 1999).

As can be seen from Fig. 1, the PCEs approximate the true model output generally well in all scenarios, showing that the algorithm for determining PCEs was properly implemented. The corresponding R^2 values ranged between 0.78 and 0.94 (Table 3). Figure 1 and Table 3 show also that the degree of agreement is fairly independent of the scenario. Furthermore, the largest errors take place for the smallest updraft velocity, 0.2 m/s, and the degree of agreement generally increases with increasing V . This result can be explained in the following way. At low updraft velocities, no cloud droplets are predicted to be formed on Aitken mode particles in large parts of the parameter space due to low supersaturations reached during an air ascent. On the other hand, small changes in the input parameter values may produce notable changes in CD_2 , such that CD_2 increases or decreases steeply with the changing value of the parameter. Thus CD_2 exhibits a “threshold behaviour” that is difficult to be captured using a polynomial approximation, and this is also the reason why updraft velocities lower than 0.2 m/s are not explicitly considered. In contrast, activation of Aitken mode particles to cloud droplets is more favorable at higher updraft velocities where CD_2 also tends to be less sensitive to the input parameter values.

For the purposes of this study, it is more important that the statistical features of the

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model output are produced with a sufficient accuracy than that a good approximation for the true model output is obtained in every point of the parameter space. Therefore we compared also the PCE-based PDFs of CD_2 with corresponding ones generated from the true model output. Figure 2 shows the comparison for two updraft velocities, $V=0.2$ and 1.0 m/s which include cases with the least and largest degree of agreement (Table 3). As can be seen, a quantitative agreement is reached in most cases: the PCEs produce the basic characteristics, such as the shape and peak, of the PDFs describing the true model output. For cases with $V=0.2$ m/s, however, the PCM-based PDFs are biased towards smallest (<10 cm⁻³) and largest (>1000 cm⁻³) values of CD_2 as compared to the PDFs of the true model output. In addition, the peak of the PDF is notably shifted to smaller concentrations in the “BASE” and “FILM” scenarios. These discrepancies are caused by the “threshold” behavior discussed above. However, the agreement improves rapidly when V increases and as Fig. 2 illustrates, a very good agreement is reached for higher updraft velocities.

The PCE and model-based expected values and variances of $\ln(CD_2)$, $E[\ln(CD_2)]$ and $Var[\ln(CD_2)]$, respectively, were also compared, and the relative errors in predicting $E[\ln(CD_2)]$ and $Var[\ln(CD_2)]$ are shown in Table 3. As seen, $E[\ln(CD_2)]$ is reproduced accurately in most of cases, and the maximum error is 12%. Errors in $Var[\ln(CD_2)]$ are generally slightly larger, the maximum error being 15%.

Results shown in Figs. 1 and 2 and in Table 3 provide a comprehensive characterization of the accuracy of the PCEs generated. To summarize, in spite of the biases exhibited by the PCEs at low updraft velocities, the agreement is sufficient to warrant the conclusions based on the sensitivity analysis which is discussed next.

3.2 Sensitivity study

After the PCEs were generated, the contribution of each uncertain input parameter (listed in Table 1) to the total variance of the model output was calculated using (5). As discussed in Sect. 2.3, these input parameters can be divided into two groups: those related to the modal properties (first three parameters in Table 1) and those

related to the chemical composition of the Aitken mode particles (last six parameters in Table 1). Here we call these parameters as physics- and chemistry-related parameters, respectively. With an aim to find out which of them cause most of the uncertainty, Fig. 3 shows the relative importance of the particle size distribution versus the particle chemical composition for all three scenarios.

Common to all the results is that the importance of the particle chemical composition decreases as V increases. This feature is consistent with the results of Ervens et al. (2005) who predicted that the cloud droplet number concentration becomes less sensitive to the particle chemical composition as the updraft velocity increases. Also, the net contribution of the physics-related parameters to the total variance becomes larger than that of the chemistry-related parameters at updraft velocities of ~ 0.3 and ~ 0.9 m/s in the “MACRO” and “FILM” scenarios, respectively. In the “BASE” scenario, the physics-related parameters dominate the total variance of the model output regardless of the updraft velocity. The larger roles of chemistry in the “MACRO” and “FILM” scenarios are mainly due to larger value ranges adopted for the surface tension and mass accommodation coefficient, respectively (Table 1), as will be shown below. Overall, these results suggest that the chemical composition can be as important as the physical properties what it comes to the cloud-nucleating ability of Aitken mode particles in the continental background areas.

Next we elucidate which individual parameters are behind the features displayed in Fig. 3. Figures 4, 5 and 6 show the contribution of the uncertain input parameters (listed in Table 1) to the total variance of the model output for the “BASE”, “MACRO” and “FILM” scenarios, respectively. Furthermore, the average contribution of each parameter is shown in Table 4 for all the scenarios. The averaging is performed over the considered updraft velocities with the same weight given for each value of V .

3.2.1 “BASE” scenario

The “BASE” scenario is considered first. Figure 4 shows that the physics-related parameters CN , D_m and σ_g , dominate the total variance of the model output, CN being the

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most important parameter in this respect. It is also seen that the relative importance of CN increases with increasing V . This is due to larger maximum supersaturations reached at higher updraft velocities which allows for a larger fraction of Aitken mode particles to form cloud droplets. Consequently, the value of CD_2 reflects that of CN at higher updraft velocities. This is also the main reason for the increasing importance of the physics-related parameters with increasing V (Fig. 3).

Figure 4 shows also that the relative importance of σ_g decreases with increasing V . This is because typically only a small fraction of the Aitken mode particles, i.e. particles belonging to the “tail” which extends to larger sizes, are able to form cloud droplet at low updraft velocities. Under these conditions, increasing σ_g increases the number of particles in the “tail” and hence also CD_2 given that other factors remain constant. At higher updraft velocities, however, CD_2 becomes less sensitive to σ_g since larger fractions of the Aitken mode particles generally activate. To illustrate this point, let us consider an extreme case in which exactly half of the Aitken mode particles form cloud droplets and the mode is internally mixed. In this situation, CD_2 does not depend on σ_g at all, provided that other factors do not change with changing σ_g . When more than half of the Aitken mode particles nucleate to cloud droplets, the sensitivity of CD_2 to σ_g starts to increase again. However, this does not take place in most of the calculations which together with the other factors discussed here explains the feature.

The relative importance of the third physics-related parameter, D_m , decreases slightly with increasing V . The decrease is because the minimum particle diameter decreases with increasing maximum supersaturations and hence the particle size plays smaller role at higher updraft velocities. Another interesting result is that σ_g and D_m have both approximately equal importance at $V < 0.4$ m/s. This suggests that the shape of the Aitken mode has also to be accounted for when predicting the contribution of sub-100 nm particles to cloud droplet concentrations at regimes with low updraft velocities.

The parameters ε , MW_{avg} and σ_s are the three most important chemistry-related parameters, as can be seen from Fig. 4 and Table 4, with roughly equal contribution from each parameter. Their contributions are, however, smaller than those of CN and

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D_m . More generally, the net contribution of the chemistry-related parameters is smaller than that of physics-related parameters (Fig. 3), which implies that if the chemical composition of the atmospheric Aitken mode particles varies in the range characteristic to the “BASE” scenario, the modal properties have a larger overall effect on the cloud-nucleating ability of the Aitken mode particles than the chemical composition. Regarding the other uncertain parameters, it is worth noting that α contributes only marginally to the total model variance, except in the case with the lowest updraft velocity. Such a drastic change, or “jump”, in the relative importance of α is not intuitive in view of the fact that the relative contributions of other uncertain parameters display a coherent behavior. Moreover, it is not seen in the “MACRO” scenario (see below) even though α has the same value range in these two scenarios (Table 1). We speculate that the “jump” is caused by the inability of the PCE to capture the effect of α at low updraft velocities and should thus be considered as an artifact.

3.2.2 “MACRO” scenario

The “MACRO” scenario differs from the “BASE” scenario by larger value ranges adopted for MW_{avg} , $\nu\Phi$ and σ_s (Table 1). The difference is also reflected in the results: the chemical composition of the Aitken mode particles is more important in the “MACRO” scenario than in the “BASE” scenario (Fig. 3). As seen from Fig. 5, this is mainly due to σ_s which is the most important parameter regarding the total variance of the model output at $V < 0.3$ m/s. The contribution of σ_s is exceeded by that of CN at larger updraft velocities, and the reason for the growing importance of CN is the same as in the “BASE” scenario discussed above. The results for “MACRO” and “BASE” have also other common features: the importance of σ_g decreases strongly with increasing V . It is further seen that the parameter CD_2 is rather insensitive to α and ρ , but exhibits notable sensitivity to D_m regardless of the updraft velocity. The parameters ε , MW_{avg} and $\nu\Phi$ have similar contributions to the total uncertainty, but the relative importance of these three parameters shows some variation with the updraft velocity.

To summarize, the largest change compared to the “BASE” scenario is the increased

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role of the surface tension, which leads to the following conclusion: if particle-phase organics do not decrease the surface tension of Aitken mode particles by more than approximately 30%, the uncertainties caused by the surface tension are comparable to those caused by uncertainties in the solubility and molecular weight of the organics.

5 However, a sufficient presence of extremely surface-active organics may cause substantial uncertainties in predictions concerning the cloud-nucleating ability of Aitken mode particles.

3.2.3 “FILM” scenario

The contributions of the uncertain model parameters to the total variance of the model output are shown in Fig. 6 for the “FILM” scenario (see also Table 4 for the average values). The most notable feature of the results is that α makes the largest contribution to the total variance of the model output at updraft velocities of approximately <0.9 m/s. It is also seen that only α and CN contribute more than 10% to the total variance, CN becoming more important at larger updraft velocities. The relative contributions of the other model parameters are similar to those in the “BASE” scenario and are not described explicitly here. We note, however, that the contribution of σ_s varies between 3 and 8% in a manner that is difficult to interpret. In any case, it is clear that the importance of σ_s is much smaller than those of α and CN .

The most important conclusion following from our results is that when the value of the parameter α is in the range expected for pure water surfaces, i.e. between 10^{-2} and unity, it forms a relatively small source of uncertainty in the cloud droplet formation predictions. In contrast, if α varies more than three orders of magnitude at conditions close to reaching supersaturation, the variability translates to large uncertainties regarding the ability of atmospheric sub-100 nm particles to form cloud droplets. In comparison, Chuang (2006) predicted that the cloud droplet formation exhibits large sensitivity to α when the value of α is below a critical value that ranges between 0.1 and 10^{-3} depending on the droplet size.

In view of the fact that the minimum value of α can potentially be even lower than

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the value applied here (see Sect. 2.3.), it can be asked how results would change by decreasing the minimum value of α further by one or two orders of magnitude. To this end, we performed also calculations using 10^{-4} as a minimum value for α , but the agreement between the PCEs and the true model output was notably worse compared with the results for the three scenarios considered here. This was probably due to complicated cloud formation dynamics caused by extremely small values of α . In view of this, it is expected that the importance of α increases even further with increasing level of uncertainty.

4 Discussion and conclusions

The present study attempts to identify and rank the physico-chemical properties of Aitken mode particles that determine the particle cloud-nucleating ability in continental background areas. The approach is based on performing model calculations with an adiabatic air parcel model and analyzing the model output with the probabilistic collocation method (PCM). The PCM is a tool for “global” sensitivity analysis and it allows for quantification of the uncertainties in the model output. Here the model output of interest is the number concentration of cloud droplets formed on Aitken mode particles, CD_2 , and the uncertain model parameters were those describing the modal and chemical properties of the Aitken mode particles.

The relative roles played by the particle size distribution and chemical composition in determining the cloud-nucleating ability of atmospheric particles is a subject of intense research at the present (McFiggans et al., 2006; Dusek et al., 2006; Ervens et al., 2007), and the results of our study have also implications on this issue. First, given that the Aitken mode particles do not contain molecules that are able to reduce the particle surface tension more than 30% and/or decrease the mass accommodation coefficient of water, α , below 10^{-2} , the chemical composition and modal properties may have roughly an equal importance at low updraft velocities characterized by maximum supersaturations $<0.1\%$ (Fig. 4 and Table 3). For larger updraft velocities, however,

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the particle size distribution is more important than the chemical composition. Furthermore, the largest uncertainties generally arise from the particle number concentration, followed by the particle size. Second interesting result is that the shape of the particle mode, characterized by the geometric standard deviation (GSD), can be as important as the mean size of the mode at low updraft velocities. This suggests that using a prescribed value for the GSD (see Vignati et al., 2004 for example) might cause errors to the predicted effect of sub-100 nm sized particles on the cloud droplet number concentrations.

The performed sensitivity analysis revealed also that the chemistry may dominate the total uncertainty in CD_2 if: 1) the value of α varies at least one order of magnitude more than what is expected for pure water surfaces (10^{-2} –1), or 2) the particle surface tension varies more than roughly 30% under conditions close to reaching supersaturation. These results provide motivation for experimental studies aiming to find out if α may reduce below 10^{-2} or if the surface tension may reduce below ~ 0.05 N/m for aerosols comprising of atmospherically relevant mixtures at relevant dilution levels.

The largest sources of uncertainty in the conclusions presented above arise from poorly characterized chemical composition of sub-100 nm atmospheric particles. When more information on the particle chemical composition emerges, however, it can be utilized to constrain the probability distributions of uncertain model parameters. Consequently, the accuracy of the sensitivity analysis will improve (Tatang et al., 1997). We also expect that the results of the study are generally applicable to atmospheric conditions where supersaturations reached during cloud formation are of similar magnitude than in simulations considered here. On the other hand, the results are probably not applicable to polluted conditions where supersaturations are considerable lower. Because of the low supersaturations, however, the contribution of Aitken mode is expected to be insignificant in such areas.

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Table 1. Investigated parameters, their abbreviations and ranges over which their values were varied. All the parameters refer to Aitken mode particles and not to the whole particle population.

Parameter	Abbreviation	“BASE”	“MACRO”	“FILM”
Geometric standard deviation	σ_g	1.3–1.9	1.3–1.9	1.3–1.9
Total particle concentration (cm^{-3})	CN	10–10 000	10–10 000	10–10 000
Particle mean diameter (nm)	D_m	50–100	50–100	50–100
Average molecular weight (g mol^{-1})	MW_{avg}	60–250	60–600	60–250
Water-soluble mass fraction	ε	0.25–1.0	0.25–1.0	0.25–1.0
Particle dry density (g cm^{-3})	ρ	1.0–2.0	1.0–2.0	1.0–2.0
“Effective” Van’t Hoff factor	$\nu\Phi$	1–3	1–5	1–3
Particle surface tension (N m^{-1})	σ_s	0.05–0.072	0.02–0.072	0.05–0.072
Mass accommodation coefficient	α	10^{-2} –1	10^{-2} –1	10^{-3} –1

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Table 2. Legendre polynomials (P_n) contained by Eq. (2) and the integrals $E(P_n)$ and $E(P_n^2)$ (Eq. 3).

Order	$P_n(x)$	$E(P_n)$	$E(P_n^2)$
0	1	1	1
1	x	0	1/3
2	$1/2 \times (3x^2 - 1)$	0	1/5
3	$1/2 \times (5x^3 - 3x)$	0	1/7
4	$1/8 \times (35x^4 - 30x^2 + 3)$	0	1/9

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Table 3. The coefficients of determination, R^2 , relative errors in the expected values and total variances of $\ln(CD_2)$ ($E[\ln(CD_2)]$ and $Var[\ln(CD_2)]$, respectively) for updraft velocities $V=0.2$, 0.4 and 1.0.

	"BASE"			"MACRO"			"FILM"		
	0.2	0.4	1.0	0.2	0.4	1.0	0.2	0.4	1.0
V (m/s)	0.2	0.4	1.0	0.2	0.4	1.0	0.2	0.4	1.0
Average SS_{\max} (%)	0.12	0.16	0.28	0.12	0.15	0.27	0.12	0.17	0.29
R^2	0.78	0.83	0.94	0.8	0.77	0.83	0.79	0.81	0.82
Error in $E[\ln(CD_2)]$ (%)	6	0.8	0.6	1	0.6	0.3	12	8	2
Error in $Var[\ln(CD_2)]$ (%)	5	11	1	10	3	3	14	0.3	15

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Table 4. Average contributions of the uncertain input parameters to the total variance of the model output (in percentages). Averaging is performed over all considered updraft velocities for a scenario with equal weight given for each case. Three largest sources of the variance are indicated with bold.

Abbreviation	“BASE”	“MACRO”	“FILM”
σ_g	8	6	6
CN	51	43	23
D_m	13	9	8
ε	7	5	3
MW_{avg}	6	6	4
ρ	2	2	2
$v\Phi$	4	6	3
σ_s	5	22	4
α	3	1	46

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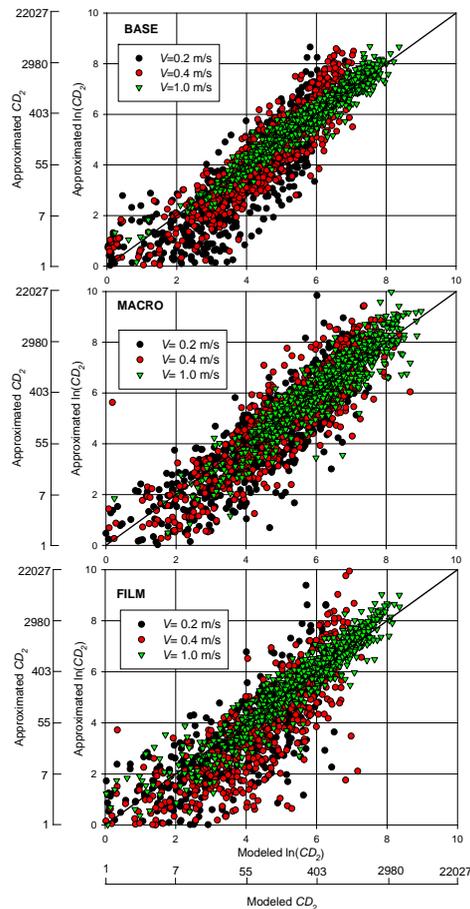


Fig. 1. A comparison of $\ln(CD_2)$ predicted by the adiabatic air parcel model and by the corresponding PCEs. The scenario is shown in each plot, and the updraft velocity (V) is shown in the legend. Furthermore, 1:1 line is added to each plot to guide the eye.

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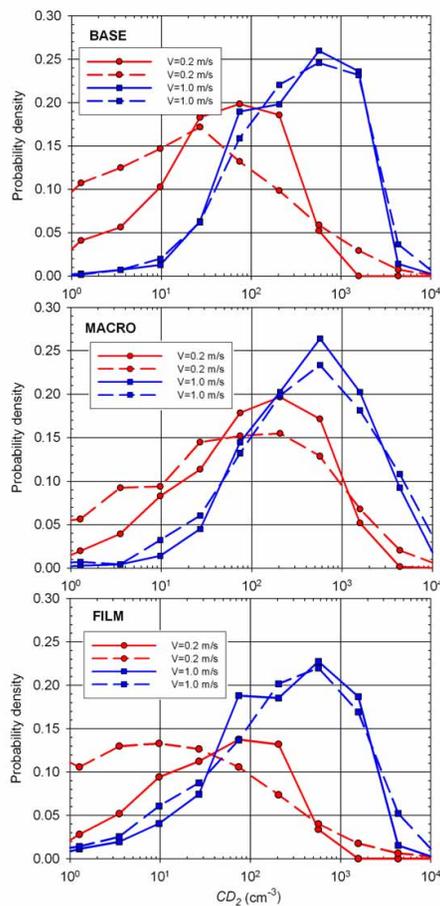
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Fig. 2. The probability density functions representing the original model output (solid lines) and samples from the PCEs (dashed lines). The scenario is shown in each plot, and the updraft velocity (V) is shown in the legend.

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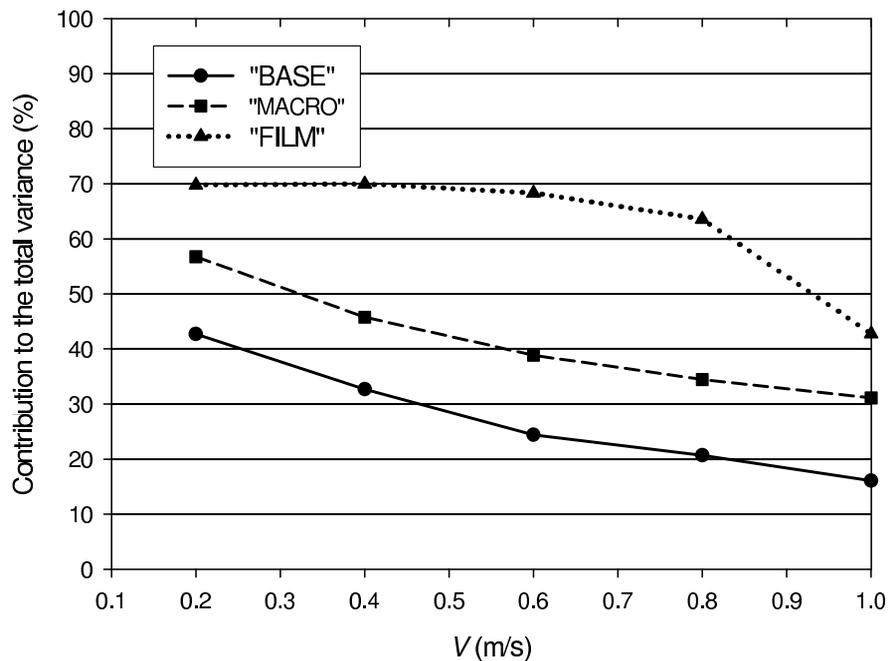


Fig. 3. The net contribution of chemistry-related parameters to the total variance of the model output as a function of the updraft velocity, V . The scenario is indicated in the legend.

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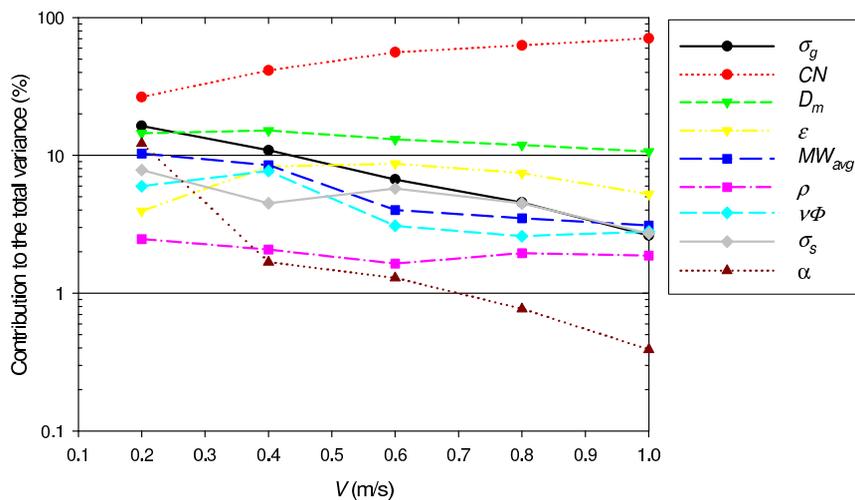


Fig. 4. The contributions of the uncertain model parameters to the total variance of the model output as a function of the updraft velocity, V , for the “BASE” scenario. The model parameters are indicated in the legend.

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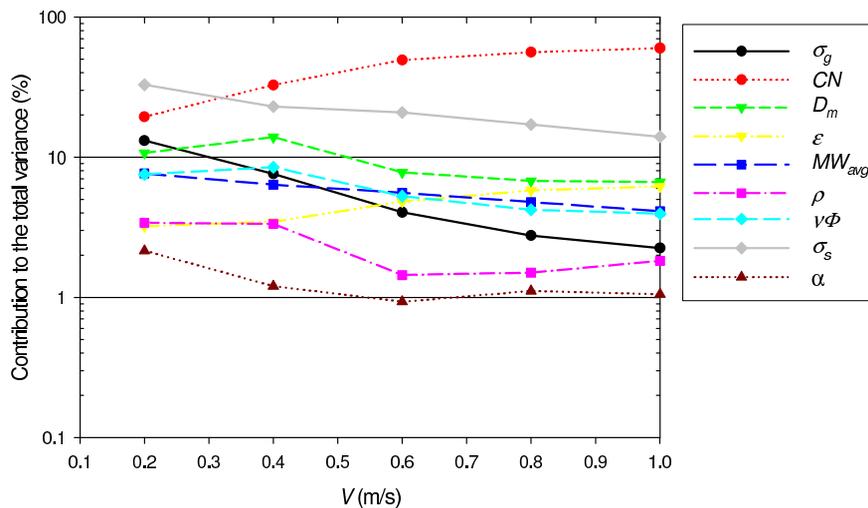


Fig. 5. Same as Fig. 4, but for the “MACRO” scenario.

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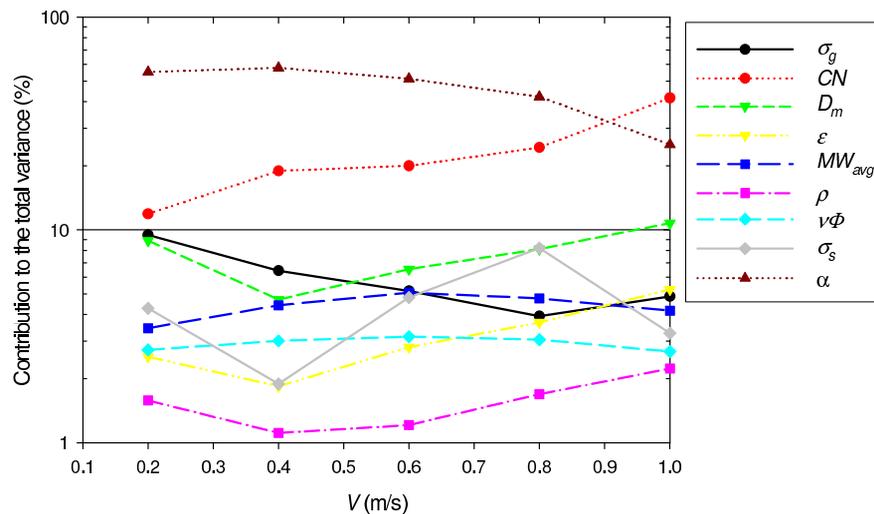


Fig. 6. Same as Fig. 6, but for the “FILM” scenario.

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