

Modelling investigation of the thermal treatment of ash-contaminated particulate filters

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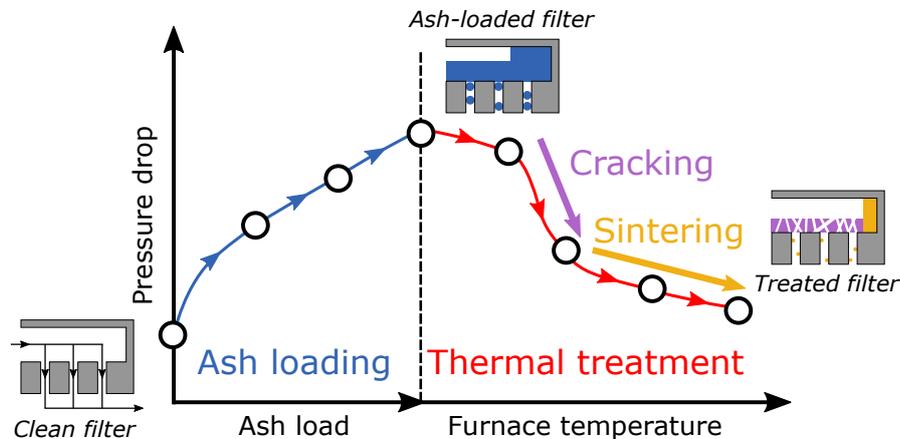
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Abstract

This paper investigates the impact of thermal treatment on the pressure drop of particulate filters containing ash deposits. A model has been developed and applied to describe the deposition of soot and ash particles, and estimate the spatial distribution of the deposits in such filters. Phenomenological models have been developed to describe the potential sintering and cracking of the ash deposits caused by thermal treatment of the filter. The model results are in good agreement with experimental measurements of the reduction in the pressure drop in thermally treated filters. It was found that crack formation in the ash layer can lead to significant reduction of the pressure drop at relatively low temperatures. Sintering of ash deposits in the wall and the ash plug also contributes towards a decrease in filter pressure drop at higher temperatures. This work is the first attempt to model the impact of the thermal treatment of ash in particulate filters in order to support the development of future ash management strategies. The cracking of the ash layer during the thermal treatment has been identified to be the most critical effect for pressure drop reduction.

Which mechanism contributes the most towards reducing ash-induced filter pressure drop?



Highlights

- Impact of thermal heating on ash deposits in particulate filters investigated
- Phenomenological model developed to describe sintering and cracking of ash
- Crack formation in ash responsible for substantial reduction in pressure drop

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1 Introduction

Internal combustion engines (ICEs) are a major source for particulate matter (PM) emissions [13]. Particulate matter is known to have an adverse impact on public health [32] and the environment [47]. Governing bodies around the world have been imposing increasingly stringent emission standards and more rigorous testing protocols to reduce pollution [1, 2, 10, 43] from ICEs. Diesel particulate filters (DPFs) have been widely adopted for diesel-fuelled engines in order to comply with these emission standards [24]. Recently, the introduction of regulations governing particle number (PN) emissions has also driven the introduction of gasoline particulate filters (GPFs) on gasoline direct injection (GDI) vehicles [5, 16, 37].

The typical operating cycle of a particulate filter involves the collection of solid particles, followed by an active regeneration process that reduces the pressure drop of the filter by burning off the trapped particles. The typical behaviour of a particulate filter during the filtration process is shown in Fig. 1. Particles are initially trapped in the porous walls of the filter by deep-bed filtration, leading to a rapid increase in the pressure drop and an increase in the filtration efficiency of the filter. As the pore volume within the filter walls reduces, particles can no longer penetrate the walls and a filter cake is formed on the surface of the walls. The pressure drop increases at a lower rate during the cake formation stage. Regeneration is triggered when the filter pressure drop reaches a preset threshold.

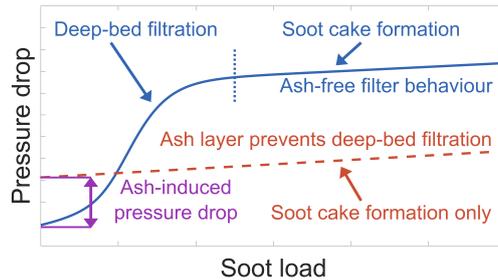


Figure 1: Typical behaviour of ash-free and ash-loaded filters during soot filtration. Deep-bed filtration via soot deposition inside the porous walls of the filter leads to a rapid increase in the pressure drop at low soot loading in the ash-free filter. Ash deposits in the ash-loaded filter lead to an increase in the pressure drop compared to the unloaded filter, but prevent deep-bed filtration of the soot in resulting a lower pressure drop compared to the ash-free filter at high soot loading.

In addition to soot, the particulate emissions from ICEs contain metallic ash originating from lubrication oil, fuel additives and abrasion [66]. Unlike soot, the metallic ash is incombustible, such that it accumulates over the lifetime of the filter [26]. While metallic ash makes up only a small amount of diesel PM [18, 19], PM emitted by GDI engines can have much higher ash content [8], with studies showing numbers in the range of 20%–50% by mass [59]. Whilst the accumulation of the ash contributes to the pressure drop [15], the presence of an ash layer on the filter walls can lead to improved filtration efficiency. The ash layer can eliminate the deep-bed stage of the filtration and, in so doing, can reduce the pressure drop over the filter at high soot loadings [68]. See Fig. 1. This

is especially beneficial for GPFs, where soot cake formation is harder to sustain [34]. On the other hand, ash deposits reduce the available storage capacity, necessitating an increase in the frequency of regeneration events and incurring an associated fuel penalty [39]. Ash deposits can also lead to an increase in the maximum filter temperature during regeneration [3] which can damage the filter and limits its service lifetime [6].

Many technologies have been considered to minimise the negative impact of ash accumulation and extend the service lifetime of particulate filters. The use of lubricants with minimal inorganic additives (“ashless oil”) can mitigate the production of lubricant-derived ash [62]. However, the contribution of corrosion and wear can still be significant, where it has been found that up to 50% of ash deposits in GPFs could come from corrosion of the moving parts in an engine and from eroded upstream catalysts [25]. Some progress has been made using alternative filter geometries. Asymmetric Channel Technology (ACT) has been shown to improve the ash storage capacity of particulate filters by having larger inlet channels than outlet channels [67]. Alternatives to square channel cross-section designs have also been shown to reduce ash-induced pressure drop *e.g.* hexagonal and octo-square geometries [65]. The engine system operation may also be optimised to minimise the impact of ash deposits by modifying the distribution and properties of the ash within a filter [66].

The deposition of ash in particulate filters has been studied by Sappok et al. [50, 51, 52, 53] using an accelerated ash loading system. Sappok et al. [52] observed that the pressure drop of ash-contaminated filters decreased after exposure to high temperatures. Analysis of the filters showed a reduction in the bulk volume of ash deposits due to sintering and crack formation in the ash layer. This indicates the potential of thermal treatment as a viable ash management strategy [66]. However, the relative contributions of the cracking and sintering are unexplored.

The **purpose of the paper** is to study the impact of thermal treatment on ash deposits in particulate filters using a computational model. Modelling allows the exploration of scenarios that are difficult to investigate using experiments. The development of the model provides helps quantify the effects of the thermal treatment process and provides insights that can aid the optimisation of ash management strategies. In this work we aim to investigate the relative importance of sintering and cracking on the reduction of the ash-induced pressure drop in a particulate filter.

The rest of the paper is structured as follows: The model is described in Section 2. The model is calibrated and assessed against the experimental data reported by Sappok et al. [50, 51, 52] in Section 3.2. The model for thermally treated ash deposits is applied in Section 3.3. Suggestions for future development are discussed in Section 3.4.

2 Model description

The model used in this work considers a pair of inlet and outlet channels in a Wall Flow Monolith (WFM) – a common design for particulate filters. Aspects of the model have been described previously, where it was used to consider soot filtration [27, 28]. The spatial representation of the model is shown in Fig. 2(a). The channels are discretised in

the axial and through-wall directions, resolving gradients along the channels and through the walls, but neglecting any variation across the soot cake. Particles are introduced into the filter via exhaust flowing into the inlet channel. The particles are then either filtered by the porous wall or exit the filter with the gas flow through the outlet channel.

In this work, the model is extended to consider ash deposition. The spatial representation of the extended model is shown in Fig. 2(b). The presence of the ash layer is explicitly considered in the model. The ash layer is formed from incombustible residues that are left behind when the soot cake is burned off during regeneration of the filter. The ash layer is expected to be in contact with the porous wall, with a new soot cake eventually forming on top of the ash layer. The ash plug is assumed to be impervious to gas flow, and is outside the spatial domain considered by the model, but is implicitly considered in the model in the sense that it reduces the length of the inlet channel.

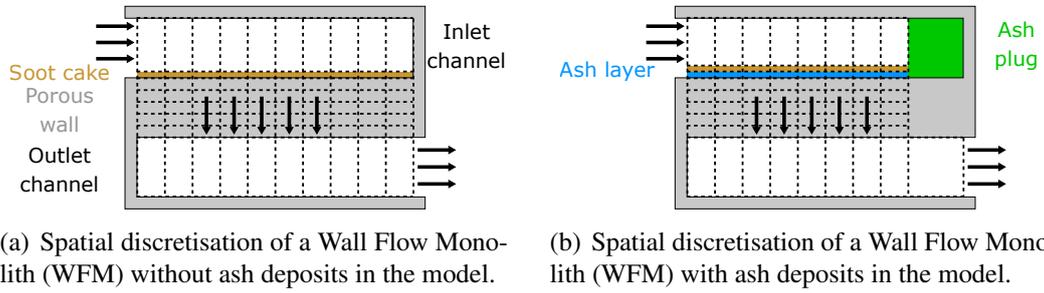


Figure 2: *Spatial representations of a Wall Flow Monolith (WFM). When ash deposits exist in the WFM, additional volume elements are introduced to describe the ash layer. The implicit reduction of the inlet channel volume due to the ash layer and ash plug formation is considered by the model.*

2.1 Governing equations

The engine exhaust first flows into the inlet channels of the WFM. Since the inlet channels are closed-ended, the exhaust is forced to flow through the soot and ash layers (should they exist) and the porous wall. The exhaust then exits the WFM via the outlet channel. The model permits gas flow in two directions: in the axial direction along the inlet and outlet channels, and through the porous media including the soot cake, the ash layer and the filter wall in the through-wall direction. These two flows are perpendicular [12].

The gas flow in the inlet and outlet channels is governed by the Bernoulli equation

$$P_{i-1} + \frac{1}{2}\rho_{i-1}u_{i-1}^2 = P_i + \frac{1}{2}\rho_i u_i^2 + \frac{F\mu L_i}{D^2}u_i, \quad i \in \begin{cases} S_{ci} & \text{inlet channel} \\ S_{co} & \text{outlet channel} \end{cases} \quad (1)$$

where P is the pressure, ρ is the mass density of the gas, u is the gas velocity along the channel, F is a friction factor ($F = 28.454$ for a square channel [56]), μ is the dynamic viscosity of the gas, L is the distance between neighbouring volume elements and D is the side length of the cross-section of a square channel. Subscript i indexes the volume

elements of the channels and increments in the axial direction. Eq. (1) applies to both inlet and outlet channels, where the index sets S_{ci} and S_{co} reference the volume elements in the inlet and outlet channels respectively. Eq. (1) is solved to obtain P and u . The parameters F , L and D are provided as inputs based on the geometry of the particulate filter. The ideal gas law is used to calculate ρ from the thermodynamic state of the gas phase. The dynamic viscosity of the gas mixture is assumed to be that of air.

The pressure and the gas flow from the inlet channel, through the porous media and into the outlet channel is governed by Darcy's law

$$P_j - P_{j+1} = \frac{\mu v_j L_j}{\kappa_j}, \quad j \in S_{ci} \cup S_{pm} \cup S_{co} \quad (2)$$

where v and κ are the gas flow velocity in the through-wall direction and the permeability across the porous medium. Subscript j indexes the volume elements in the through-wall direction. The index sets S_{ci} , S_{pm} and S_{co} reference the volume elements in the inlet channel, the porous media, including the soot cake layer, the ash layer and the filter wall, and the outlet channel respectively. Eq. (2) is solved to obtain P and v . The treatments of L and μ are identical to eq. (1). The permeability of porous medium κ is provided by the filtration model (see Section 2.2).

The temperature of the system is assumed to be constant and chemical reactions are neglected in the simulations in this work, such that the thermal energy balance and species equations are omitted.

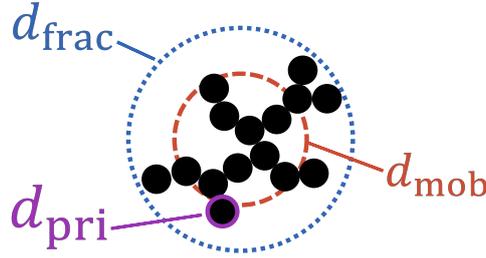


Figure 3: Illustration of the primary particle diameter d_{pri} , the mobility diameter d_{mob} and the fractal diameter d_{frac} of a typical aggregate considered by the particle model.

Previous applications of the model [27, 28] assumed all particles to be spherical and purely carbonaceous. These assumptions are relaxed in this work. Fig. 3 illustrates a typical aggregate considered by the particle model used in this work. Particles are described by elements of a type space E

$$E = \{m_{agg}, n_{pri}, m_{ash}\}, \quad (3)$$

where m_{agg} is the total aggregate mass, n_{pri} is the number of primary particles in the aggregate and m_{ash} is the mass of ash inside the aggregate. The aggregate mass other than ash is assumed to be carbonaceous material.

The size distribution of the particles is described by the particle number density N . The population balance model is solved using a sectional method [42]. The sectional method has been successfully applied in modelling the evolution of soot particle in flames [9, 69]

and engines [7, 30]. The particle population is divided into sections (*i.e.* size classes) according to the value of m_{agg} for each particle. All primary particles belonging to aggregates within a given size class are assumed to have the same size [9, 30]. In this work, population balance equations of the form

$$\frac{dN_{z,j}}{dt} = Q_{z,j}\dot{N}_{z,j-1}, \quad j \in S_{\text{ci}} \cup S_{\text{pm}} \cup S_{\text{co}} \quad (4)$$

are solved for N_{agg} , N_{pri} and N_{ash} , corresponding to each element of the type space defined in eq. (3) and representing the total number density of aggregates, primary particles and equivalent ash mass in a size class. $Q_{z,j}$ is the filtration efficiency of particles of size z calculated from the filtration model (see Section 2.2). $\dot{N}_{z,j-1}$ is the rate of change of the number density of particles flowing from volume element $j-1$ to volume element j

$$\dot{N}_{z,j} = (1 - Q_{z,j})\dot{N}_{z,j-1}, \quad j \in S_{\text{ci}} \cup S_{\text{pm}} \cup S_{\text{co}}. \quad (5)$$

The mass of an aggregate m_{agg} for a given size class is known *a priori*. The other type space variables are calculated

$$n_{\text{pri},z,j} = \frac{N_{\text{pri},z,j}}{N_{\text{agg},z,j}}, \quad (6)$$

$$m_{\text{ash},z,j} = m_{\text{agg},z,j} \frac{N_{\text{ash},z,j}}{N_{\text{agg},z,j}}. \quad (7)$$

The primary particle diameter is calculated

$$d_{\text{pri},z,j}^3 = \frac{6}{\pi} \frac{m_{\text{agg},z,j}}{n_{\text{pri},z,j} \rho_{\text{agg},z,j}}, \quad (8)$$

where ρ_{agg} is the material density of the aggregate.

2.2 Filtration model

The filtration of the particles is described using the classic unit collector model [31]. The unit collector model is used to calculate the filtration efficiencies and permeabilities that are required in eqs. (2) and (4). The unit collector model has been widely adopted in particulate filter modelling studies [21]. It considers the filter wall as a bed of spherical unit collectors. The model approximates the gas flow through the porous wall as the gas flow around a spherical unit collector and is able to describe the filtration of particles via diffusion and interception [21, 31], as illustrated in Fig. 4.

The diameter of the collector d_{coll0} representing the clean porous wall is calculated as [44]

$$d_{\text{coll0}} = \frac{3}{2} \frac{1 - \varepsilon_{\text{w0}}}{\varepsilon_{\text{w0}}} d_{\text{pore0}}, \quad (9)$$

where ε_{w0} and d_{pore0} are the porosity and pore diameter of the clean porous wall, and are inputs to the model. The filtration efficiencies of the porous wall and the ash layer are calculated using equations of the form

$$Q_{z,i} = 1 - \exp \left[-\frac{3\eta_{z,i}(1 - \varepsilon_i)w_i}{2\varepsilon d_{\text{coll},i}} \right], \quad (10)$$

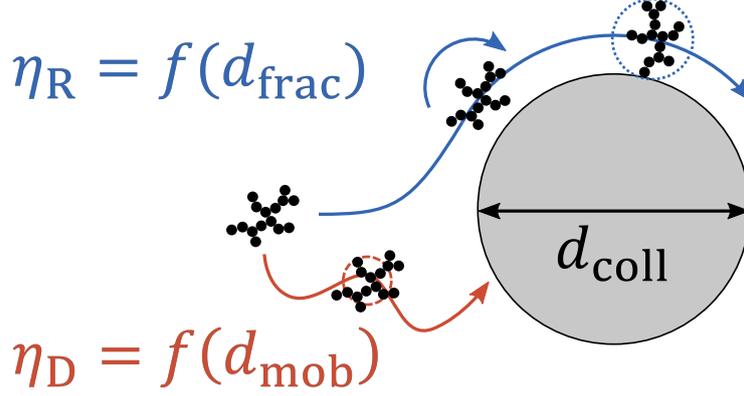


Figure 4: The unit collector model approximates the flow in the porous wall as gas flow through a bed of equivalent spherical collectors. Particles may be trapped by the collectors via diffusion (“D”) and interception (“R”).

where ε is the porosity of the filter medium, w is the thickness of the filter medium, d_{coll} is the diameter of the collector and η is the efficiency of a single collector. The single collector efficiency $\eta_{z,i}$ and hence the filtration efficiency $Q_{z,i}$ depend on both the size of the particles (indexed by z) and the state of the filter medium in volume element i . The values of ε , d_{coll} and w depend on the volume element index i which indicates whether the volume element represents the filter wall or the ash layer. In the case of the porous wall, the thickness of the wall is treated as a constant, whereas the porosity and collector diameter are calculated based on the particulate loading within the wall (see Section 2.2.1). In the case of the ash layer, the porosity and collector diameter are inputs to the model and are treated as constants during simulations. The thickness of the ash layer is a function of the mass of ash accumulated on the surface of the filter wall.

The single collector efficiency η in eq. (10) combines contributions from diffusion and interception (see Fig. 4). These two mechanisms are assumed to act independently, such that

$$\eta_{z,i} = \eta_{D,z,i} + \eta_{R,z,i} - \eta_{D,z,i}\eta_{R,z,i} \quad (11)$$

where subscripts D and R refer to diffusion and interception respectively.

The diffusional single collector efficiency η_D is a function of the filter porosity and the Peclet number of the particle [21, 31]

$$\eta_{D,z,i} = 3.5g(\varepsilon_i) \text{Pe}_{z,i}^{-\frac{2}{3}}, \quad (12)$$

where $g(\varepsilon)$ is a Kuwabara geometric function

$$[g(\varepsilon_i)]^3 = \frac{\varepsilon_i}{2 - \frac{9}{5}(1 - \varepsilon_i)^{\frac{1}{3}} - \varepsilon_i - \frac{1}{5}(1 - \varepsilon_i)^2}, \quad (13)$$

and Pe is the Peclet number which is calculated as per Konstandopoulos et al. [21]. The relevant length scale for the Peclet number is the *mobility diameter* of the particle d_{mob} . This is calculated as per Sorensen [60]

$$d_{\text{mob},z,i} = \begin{cases} d_{\text{pri},z,i} n_{\text{pri},z,i}^{0.46} & n_{\text{pri},z,i} < 100 \\ 0.65d_{\text{pri},z,i} n_{\text{pri},z,i}^{0.56} & n_{\text{pri},z,i} \geq 100 \end{cases}, \quad (14)$$

where d_{pri} is the diameter of the primary particles within the aggregate. The values of d_{pri} and n_{pri} are calculated as per eqs. (6) and (8).

The interceptional single collector efficiency η_R is calculated by [21, 31]

$$\eta_{R,z,i} = \frac{3}{2} [g(\varepsilon_i)]^3 \frac{R_{z,i}^2}{(1 + R_{z,i})^{1 - \frac{2}{3\varepsilon_i}}}, \quad (15)$$

where R is the interception parameter

$$R = \frac{d_{\text{frac},z,i}}{d_{\text{coll},i}}, \quad (16)$$

defined in terms of the collector diameter d_{coll} , see eq. (10), and the fractal diameter of the aggregate d_{frac} [9]

$$d_{\text{frac},z,i} = d_{\text{pri},z,i} (V_f n_{\text{pri},z,i})^{\frac{1}{D_f}}, \quad (17)$$

where the volume filling factor V_f and the fractal dimension D_f are set to 1.43 and 1.8 respectively [9]. The fractal diameter is the diameter of the minimal enclosing sphere of the aggregate (*i.e.* the *maximum diameter*). The choice of fractal diameter as the relevant length scale in the ratio defined by eq. 16 follows from the assumption that a particle will be collected by interception when the streamline carrying the particle passes within one particle radius of the collecting body [64].

2.2.1 Treatment of clogged filter wall

As particles accumulate in the porous wall, the pore volume available for gas flow is reduced, and the pressure drop of the filter increases accordingly. This implies a decrease in the permeability of the porous wall κ_w which is used in eq. (2). The unit collector model calculates the permeability of the wall as [21]

$$\kappa_{w,i} = \kappa_{w0,i} \frac{d_{\text{coll},i}^2}{d_{\text{coll0},i}^2} \frac{f(\varepsilon_{w,i})}{f(\varepsilon_{w0,i})}, \quad (18)$$

where ε_{w0} and ε_w are the porosity of the clean wall and the clogged wall. The permeability of the clean wall κ_{w0} is provided as an input to the model. The porosity of the clogged wall is a function of the collector diameter of the clogged wall d_{coll} [20, 21]

$$\varepsilon_{w,i} = 1 - (1 - \varepsilon_{w0,i}) \frac{d_{\text{coll},i}^3}{d_{\text{coll0},i}^3}, \quad (19)$$

which increases with the mass of particles accumulated in the wall [20, 21]

$$d_{\text{coll},i}^3 = d_{\text{coll0},i}^3 + \frac{6}{\pi} \frac{m_{p,w,i}}{\rho_w}, \quad (20)$$

where $m_{p,w}$ and ρ_w are the mass and packing density of the particles accumulated in the porous wall. The mass of particles $m_{p,w}$ is solved by the model. The packing density ρ_w

is a model parameter that is typically calibrated to match experimental observations of the pressure drop across the filter. It should be noted that ρ_w should not be interpreted as the physical bulk density of the ash deposits accumulated in the filter wall, but as the bulk density of particles accumulated in the bed of unit collectors. The accumulation of particles within the pores of the ash layer is neglected in this work. This assumption is based on the experimental observation that no deep-bed filtration occurs when the ash layer is established as shown in Fig. 1.

The function $f(\varepsilon)$ appearing in eq. (18) is the Kuwabara function with the following form [29]

$$f(\varepsilon_i) = \frac{1}{18} \frac{2 - \frac{9}{5}(1 - \varepsilon_i)^{\frac{1}{3}} - \varepsilon_i - \frac{1}{5}(1 - \varepsilon_i)^2}{1 - \varepsilon_i}. \quad (21)$$

Note that this has a factor of 4 difference from the equation reported by Konstandopoulos et al. [23]. Eq. (21) adopts the formulation from Lee et al. [29] which shows better consistency between the experimental pressure drop and primary particle measurements.

2.2.2 Treatments of particulate layers

As particles accumulate in the porous wall, the filtration mechanism changes and the particles start to form a soot cake at the interface between the wall/ash layer and the inlet channel. The transition from deep-bed filtration to soot cake formation is controlled by the partition coefficient

$$\Phi = \frac{d_{\text{coll,interface}}^2 - d_{\text{coll0}}^2}{(\psi d_{\text{coll,max}})^2 - d_{\text{coll0}}^2}, \quad (22)$$

where $d_{\text{coll,interface}}$ is the collector diameter in the volume element of the porous wall at the interface with the inlet channel, $d_{\text{coll,max}}$ is the maximum collector diameter possible, and ψ is the percolation constant. The percolation constant is a model parameter that is typically calibrated to match experimental observations of the change in the rate of increase in the pressure drop across the filter caused by the transition from deep-bed filtration to cake formation (See Fig. 1). The maximum collector diameter $d_{\text{coll,max}}$ is calculated from the porosity of the clean filter wall such that the unit collector has the same capacity for particles as the filter wall [20]

$$d_{\text{coll,max}}^3 = \frac{d_{\text{coll0}}^3}{1 - \varepsilon_0}. \quad (23)$$

The permeabilities of the soot cake and ash layer required by eq. (2) are calculated based on their microstructures [23]

$$\kappa = f(\varepsilon) d^2 C, \quad (24)$$

where ε is the porosity of the particulate layer, d is the diameter of the particles in the layer, C is the Stoke-Cunningham factor accounting for the slip-flow effect

$$C = 1 + \text{Kn} \left(1.257 + 0.4 \exp \left[-\frac{1.1}{\text{Kn}} \right] \right), \quad (25)$$

and Kn is the Knudsen number calculated based on the particle diameter d . It should be noted that eq. (24) requires a single value of d , even though the particles may be subject

to a polydisperse size distribution. In this work, the values of the particle diameter d for the soot cake and ash layer are supplied as single-valued inputs to the model. The value for soot is referenced from the experimental results of Sappok et al. [55] and that for ash is calibrated based on experimental pressure drop measurements of ash-loaded filters.

2.3 Regeneration model

The ash layer is formed on the surface of the inlet channels in the filter from incombustible residues (*i.e.* ash particles) in the soot cake that are left behind when the soot is burned off during regeneration. The ash layers in the particulate filters studied in this work were deposited over many loading-regeneration cycles [50]. The regeneration events are captured by post-processing simulations of the loading part of each cycle to provide initial conditions for the simulation of the next loading cycle. It is assumed that all the soot is burned off during each regeneration event, that all the ash particles trapped in the soot cake now form part of the ash layer (see Fig. 2), that the properties of the ash (in the ash layer and in the ash deposits in the porous walls of the filter) are unaltered by regeneration and that no axial transport of ash occurs. The consequence of neglecting axial transport of the ash is discussed in detail in Section 3.

2.4 Thermal treatment model

Two phenomena are expected to occur to ash deposits during the thermal treatment of particulate filters: sintering and crack formation. Both have been observed experimentally [52]. Sintering can reduce the bulk volume of ash deposits, which leads to decreased resistance to gas flow. On the other hand, sintering can alter the microstructure of the ash layer and therefore its permeability. The permeability of the ash layer can also be affected by crack formation, which allows more gas to flow through and hence results in an increase in the permeability.

The overall permeability of the ash layer κ_{ash} considering the impact of both sintering and cracking is

$$\kappa_{\text{ash}} = \kappa_{\text{s}} + \kappa_{\text{c}}, \quad (26)$$

where κ_{s} and κ_{c} are the contributions to the permeability of the ash layer arising from sintering and cracking respectively. In this work, the impact of the sintering and cracking of the ash deposits is incorporated in the model by post-processing the results of loading-regeneration simulations to modify the properties of the ash deposits to provide initial conditions for subsequent post-thermal-treatment simulations. Model equations for both phenomena are described below.

2.4.1 Sintering model

Sintering between ash particles occurs when the ash is exposed to high temperatures. The sintering leads to a reduction in the bulk volume of the ash deposits. A sintering progress

variable X is defined in terms of the relative reduction in bulk volume of ash deposits $V_{\text{ash,bulk}}$

$$X = 1 - \frac{V_{\text{ash,bulk}}}{V_{\text{ash,bulk}0}}, \quad (27)$$

where the subscript 0 refers to the volume of the ash deposit prior to thermal treatment. The value of the sintering progress variable is able to be inferred from experimental measurements (see Section 3.3) and is used to update the model input parameters to account for the effect of sintering.

Sintering of ash in the filter walls. The effect of sintering of ash in the walls of the filter is modelled by adjusting the packing density ρ_w that appears in eq. (20)

$$\frac{1}{\rho_w} = \frac{1}{\rho_{w0}} - X \left(\frac{1}{\rho_{w0}} - \frac{1}{\rho_{\text{ash}}} \right), \quad (28)$$

where ρ_{w0} is the packing density of unsintered ash deposits and ρ_{ash} is the material density of the ash ($\rho_{\text{ash}} = 3400 \text{ kg/m}^3$ in this work [50]).

Sintering of the ash layer. The effect of sintering in the ash layer is modelled by adjusting the porosity ε of the ash layer

$$\frac{1}{1 - \varepsilon} = \frac{1}{1 - \varepsilon_0} - X \left(\frac{1}{1 - \varepsilon_0} - 1 \right). \quad (29)$$

A fully sintered ash layer is expected to have $\varepsilon = 0$. In addition to the effect on bulk volume and porosity, sintering effects the permeability of the ash layer. The permeability of the sintered ash layer κ_s is evaluated using eq. (24), which requires the diameter of the particles and the porosity of the ash layer as inputs. Therefore it is necessary to model how sintering affects these two parameters.

The morphology of the ash layer may be deduced from the particle coordination number N_{cord} , which describes the (average) number of neighbouring particles for each particle in a bed of particles. A correlation between N_{cord} and the porosity of a particle layer ε is reported by van de Lagemaat et al. [63]

$$N_{\text{cord}} = \frac{3.08}{\varepsilon} - 1.13. \quad (30)$$

The correlation is shown in Fig. 5. Typical ash deposits have a porosity in the range 0.85–0.95 [66], implying a coordination number N_{cord} in the range 2.1–2.5. This is consistent with the finger-like structures predicted by particle deposition simulation studies [14].

In order to derive a relationship between the porosity and the diameter of the particles in the ash layer during sintering, the microstructure of the ash is approximated as having column-like structures, corresponding to $N_{\text{cord}} \approx 2$. The following relationship between the porosity ε_{ash} and the diameter d_{ash} of the particles in the ash layer can be derived

$$\varepsilon_{\text{ash}} = 1 - \frac{1 - \varepsilon_{\text{ash}0}}{d_{\text{ash}0}^2} d_{\text{ash}}^2, \quad (31)$$

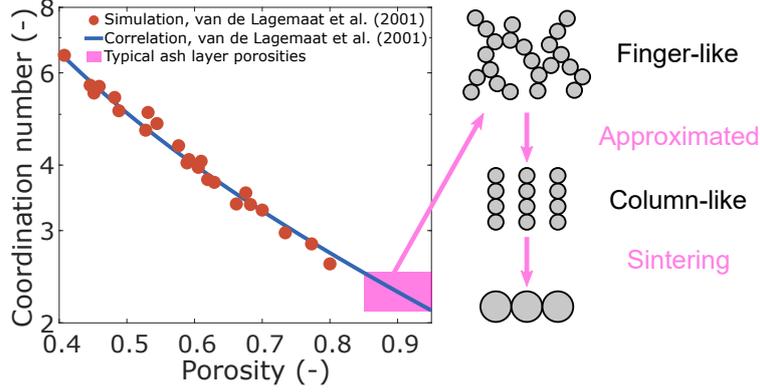


Figure 5: The combination of the observed range of ash-layer porosities and the correlation proposed by van de Lagemaat et al. [63] suggests that a typical ash layer would have a coordination number in the range 2.1–2.5, corresponding to a finger-like microstructure. The ash layer structure is approximated as particle columns for the purpose of developing a relationship between the particle size and porosity in the sintering model.

subject to the assumption that all primary particles in the ash layer have the same size and that sintering causes the particles in the columns to merge and grow until they form a mono-layer of particles in point contact. These assumptions are illustrated in Fig. 5. The symbols $\epsilon_{\text{ash}0}$ and $d_{\text{ash}0}$ denote the porosity and particle diameter of the ash layer prior to thermal treatment. The detailed derivation of the relationship in eq. (31) is given in Appendix A. Note that the relationship breaks down once the particles sinter beyond the mono-layer. The porosity of a mono-layer is $1 - \frac{\pi}{6} \approx 0.417$. If the porosity of the ash layer prior to thermal treatment $\epsilon_0 \geq 85\%$, then eq. (31) is subject to the limit $X \leq 0.713$. These limits are never exceeded in this work.

Sintering of the ash plug. In the case of an ash plug, the effect of sintering is modelled by adjusting the length of the ash plug

$$L_{\text{plug}} = L_{\text{plug}0} - X (L_{\text{plug}0} - \min[L_{\text{plug}}]), \quad (32)$$

where $\min[L_{\text{plug}}]$ is calculated under the assumption that a fully sintered ash plug would have the material density of the ash. As the modelled ash plug shrinks, a new portion of the filter wall and ash layer are exposed to the gas flow in the inlet channel. The state of this portion of the wall and ash layer are not tracked by the model (see Fig. 2(b)). In this work, it is assumed that they have the same state as the upstream neighbouring volume elements in the inlet channel. This approach is supported by the observation that the axial variation of ash deposits in the porous wall predicted by the model is negligible ($\leq 1\%$). This is substantiated in Fig. B.1 in the Appendix.

2.4.2 Cracking model

Crack formation in ash layers has been observed after thermal treatment [52]. The cracks significantly increase the permeability of the damaged ash layer. Mahadevan et al. [36]

adapted a damage permeability function from a study of concrete [45] to model the damage to a soot cake during regeneration

$$\kappa_d = \kappa_0 \exp \left[(AD)^B \right], \quad (33)$$

where κ_d is the permeability of the damaged media, κ_0 is the permeability of the intact media, A and B are model parameters and D is a damage variable. The actual form of D depends on context: it was chosen to be the dimensionless soot cake mass by Mahadevan et al. [36] and the relative change in elastic modulus by Picandet et al. [45].

Whilst eq. (33) has been shown to be consistent with experimental observations, two problems need to be solved before it can be applied to the thermal treatment of ash. Firstly, eq. (33) needs to be modified in order to consider the impact of sintering. If sintering occurs, κ_0 would not be constant. The change in permeability due to the cracks alone κ_c is [36]

$$\kappa_c = \kappa_d - \kappa_0 = \kappa_0 \left(\exp \left[(AD)^B \right] - 1 \right). \quad (34)$$

Secondly, eq. (33) can predict extremely high permeabilities for severely damaged media, which leads to numerical difficulties in the solution procedure. This was addressed by Pijaudier-Cabot et al. [46] who opted to use a Taylor expansion of eq. (33) to avoid the rapid increase in permeability for severely damaged media. In this work an alternative formulation of the cracking model in eq. (34) is proposed

$$\frac{\kappa_c}{\kappa_0} = \theta + \gamma \left(X - \frac{1}{\alpha} \exp[\alpha(X - \beta)] \right), \quad (35)$$

where α , β , γ and θ are model parameters. The damage variable is chosen to be the sintering progress variable X , see eq. (27). The apparent increase in number of parameters is negated by imposing two additional boundary conditions. First, the minimum value of the crack permeability κ_c is set to be 10^{-19} m². This value corresponds to granite which is practically impervious to gas flow [57]. Second, the maximum value of κ_c is chosen to be 3 orders of magnitude higher than the permeability undamaged media κ_0 . If the pressure drop due to the undamaged media is $O(10^3)$ Pa, then the pressure drop of the damaged media is capped at $O(1)$ Pa, sufficiently smaller than the expected experimental error.

The crack model proposed in this work in eq. (35) therefore has two degrees of freedom, the same as the cracking model in eq. (33). Fig. 6 compares the two models. The ratio between the permeability of the damaged media and that of the intact media, κ_d/κ_0 is presented. Excellent agreement between the models can be observed for $1 \leq \kappa_d/\kappa_0 \leq 10$, the range of practical interest. At large values of the damage variable, the model used in this work predicts a finite maximum permeability. This provides better numerical behaviour.

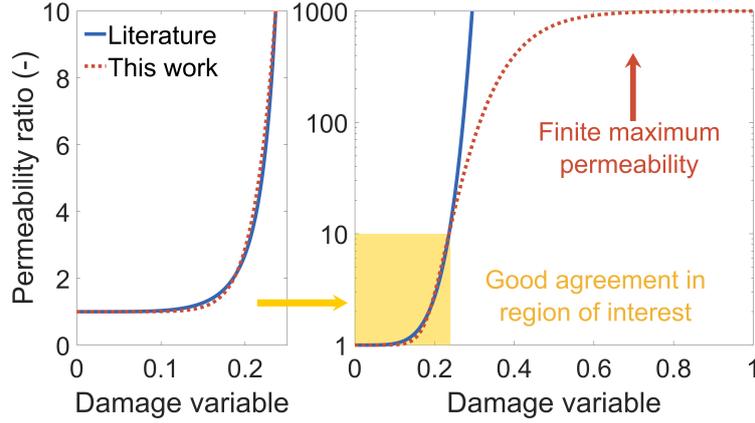


Figure 6: The cracking model parameterises the permeability of the ash layer as a function of the damage variable. The model used in this work ensures that the value of the permeability remains finite at large values of the damage variable, whilst retaining good agreement with the original model proposed by Picandet et al. [45] in the region of interest.

3 Numerical study and discussions

The model was applied to investigate the effect of ash loading and the subsequent thermal treatment of the ash on the pressure drop through a particulate filter by simulating a series of experiments by Sappok et al. [50, 51, 52, 53]. All simulations in this work were performed using the same spatial discretisation, where the number of control volumes in the axial and the through-wall directions are $n_{ax} = 10$ and $n_w = 8$ respectively. This choice of discretisation is consistent with that in other studies [27, 36].

Key details of the experiments are summarised in Section 3.1. The initial calibration of the model and simulation of the pressure drop whilst loading the filter is reported in Section 3.2. The impact of high temperature exposure on the ash-related pressure drop is investigated in Section 3.3. The implication of the results and suggestions for future studies are discussed in Section 3.4.

3.1 Experimental set-up

The experiments modelled in this work were performed using an accelerated ash loading system [50–53]. The experimental setup is shown in Fig. 7. Two particle sources were included in the accelerated loading system. A Cummins ISB 300 diesel engine that emitted soot aggregates with negligible ash content, and a custom-made burner equipped with an oil injector. The burner was used to emit soot aggregates with various levels and compositions of ash content depending on the type and rate of injection of lubricant oil into the burner. A downstream filter was loaded with particles from either the engine or the burner. This work focuses on experiments with platinum-coated DPFs [52].

The experiments were performed to measure the pressure drop across the filter in four steps. First, the pressure drop was measured at different space velocities when the filter

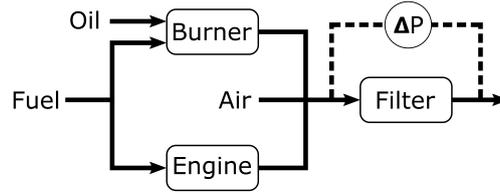


Figure 7: Schematic of the experimental setup used by Sappok et al. [50, 51, 52, 53]. The filter can be subject to in-flow from the engine, the burner or ambient air. The reported pressure drop was always measured with ambient air.

was clean (*i.e.* without any particle loading). Second, the filter was loaded with particles from the engine and the pressure drop measured as a function of soot loading. Third, the filter was loaded with particles from the burner, using CJ-4 engine oil to provide a source of ash. The ash was allowed to accumulate in the filter over multiple loading-regeneration cycles, and the pressure drop measured as a function of ash loading after each regeneration event. Two sets of measurements were performed. One that stopped when the filter reached a nominal ash loading of 12.5 g/L, the other when the filter reached 42 g/L. Fourth, once each filter reached its target loading, a cylindrical core sample was removed from the filter and subsequently heated at 650 °C for an hour to ensure that no soot remained within the sample. The samples were then heated in a furnace at a specified temperature for 30 minutes before measuring the pressure drop across each filter sample. This was repeated for furnace temperatures increasing from 650 to 1100 °C. Finally, note that the reported pressure drops were all measured using ambient air.

3.2 Model calibration

In order to study the impact of thermal treatment on the pressure drop of ash-contaminated filters, the spatial distribution of ash within the porous wall, along the inlet channel surface and within the ash plug must be known. Likewise, the porosity and the size of the particles in the ash layer must be known in order to evaluate the permeability using eq. (24). Whilst the porosity and thickness profile of the ash in selected filter channels are reported [49], the spatial distribution of the ash in the filter wall and the size of the particles in the ash layer are unknown.

The unknown model parameters (including the size of the particles in the ash layer) are estimated by progressively calibrating the model against experimental data, initially for the pressure drop across a clean filter, then for the pressure drop across a soot-loaded filter and eventually for the pressure drop across an ash-loaded filter. Once this is complete, the spatial distribution of the ash in the filter wall can be obtained as an output from the calibrated model.

The rest of the section is organised as follows. First, the contribution of the clean filter towards overall pressure drop is investigated in Section 3.2.1. Second, the unit collector model is calibrated using data from soot filtration experiments in Section 3.2.2. Finally, ash accumulation experiments are simulated in Section 3.2.3, where the model is used to estimate the particle diameter and spatial distribution of the ash deposits.

3.2.1 Pressure drop across a clean filter

The pressure drop across the clean filter arises due to frictional losses in the flow along the channels and through the porous wall. These losses are included in the model via eqs. (1) and (2). Losses due to contraction and expansion are neglected since they are only significant at high flow rates [22]. Since the frictional loss along the channels is determined by the geometry of the filter, only the permeability of the clean filter wall κ_{w0} needs to be calibrated. The calibration is executed as per Tronconi et al. [61].

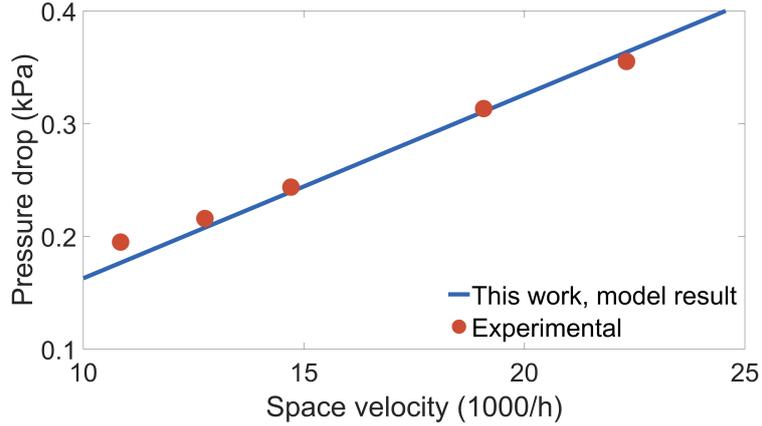


Figure 8: Pressure drop across a clean filter. The model shows good agreement with the experimental data reported by Sappok et al. [50].

A comparison of the pressure drop predicted by the model versus experimental data [50] for a clean filter is shown in Fig. 8. Good agreement is observed. The pressure drop increases with the space velocity. The experimental pressure drop has a linear relationship with the space velocity, indicating that the inertial pressure drop across the filter is negligible. The value of κ_{w0} after calibration was $1.3 \times 10^{-13} \text{ m}^2$. This value was held constant for all subsequent simulations in this work.

3.2.2 Pressure drop during soot filtration loading

The unit collector model was calibrated in order to simulate how the filtration efficiencies and permeabilities in eqs. (2) and (4) change during soot (and ash) loading.

The porosity ε_{w0} and the pore diameter $d_{\text{pore}0}$ of the clean filter wall are required as model inputs. No filtration efficiency measurements were reported for the filters considered in this study, so the values of these parameters could not be estimated as per Lao et al. [27]. Rather, the Rumpf-Gupte correlation [48]

$$\kappa_{w0} = \frac{1}{5.6} \varepsilon_{w0}^{5.5} d_{\text{pore}0}^2, \quad (36)$$

was used to estimate ε_{w0} and $d_{\text{pore}0}$ from the calibrated value of κ_{w0} . It was assumed that $d_{\text{pore}0}$ has a value of $10 \mu\text{m}$ in this work. This is on the lower end of the typical range of pore diameters ($10\text{--}30 \mu\text{m}$ [49]), but is consistent with the fact that the filters considered

in this study are coated with a platinum-based catalyst, and are therefore expected to have smaller pore diameters than uncoated filters. The corresponding value of ϵ_{w0} was 41%, which is considered to be reasonable for a coated filter. In practice these two parameters could also be measured experimentally (*e.g.* using X-ray tomography [58]). The remaining unit collector parameters were calibrated as per Kladopoulou et al. [20] and Zhang et al. [70]

- Particle packing density in porous wall $\rho_w = 8.0 \text{ kg/m}^3$.
- Soot cake porosity $\epsilon_{\text{cake}} = 0.958$.
- Percolation constant $\psi = 0.96$.

The particle packing density is within the range of values reported in the literature (6.24–14.1) [21, 35]. The soot cake porosity is within the range of 0.93–0.97 suggested by Liu et al. [33]. The percolation constant is similar to that reported by Gong et al. [11] (0.95).

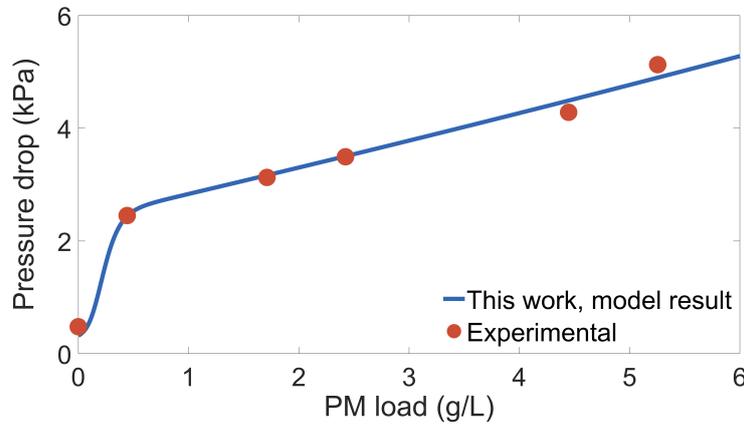


Figure 9: Pressure drop versus soot load. The calibrated model shows good agreement with the experimental data reported by Sappok et al. [50]. The change in gradient at 0.5 g/L is due to the transition between deep-bed filtration and soot cake formation as illustrated in Fig. 1.

Fig. 9 shows the pressure drop predicted by the calibrated model during soot loading. Good agreement is observed with the experimental data reported by Sappok et al. [50]. The following assumptions were made in the simulations.

- The in-situ density of soot particles was 1800 kg/m^3 [41].
- The space velocity of the soot filtration experiment was 20000 1/h, which is the same as the space velocity used for pressure drop measurement (see Section 3.1).
- The morphology of the inlet soot particles was assumed to be the same as those produced by ISB 300 engine at 25% load [55].
- The soot filtration experiment operated at $300 \text{ }^\circ\text{C}$ [55].
- No passive regeneration occurred at this condition.

3.2.3 Pressure drop due to ash

The model was applied to simulate the ash-loading of the filter. The experimental investigation of the system used an accelerated loading system that consisted of multiple loading-regeneration cycles in which the filter was loaded with burner-generated particles at 250 °C for 60 minutes, and then regenerated at 620 °C for 15 minutes every cycle [51]. Data from two experiments were reported. One that loaded a filter up to a nominal ash load of 12.5 g/L, the other that loaded a filter up to a nominal ash load of 42 g/L. Time-resolved simulations were performed for the loading part of each experiment. The simulations were stopped after each loading cycle, the results post-processed to account for regeneration, and then restarted to simulate the next loading cycle as described in Section 2.3.

Fig. 10 shows the pressure drop at the end of each loading-regeneration cycle versus the ash load accumulated in the filter. The model (black line) uses a calibrated value of 220 nm for the size of the particles in the ash layer. This value affects both the filtration efficiency and permeability of the ash layer (see Section 2.2). The filtration efficiency controls the proportion of ash deposited in the wall, and hence the point at which the transition from deep-bed to cake filtration is observed. The permeability controls the subsequent rate of increase in the pressure drop as the ash layer grows during cake formation.

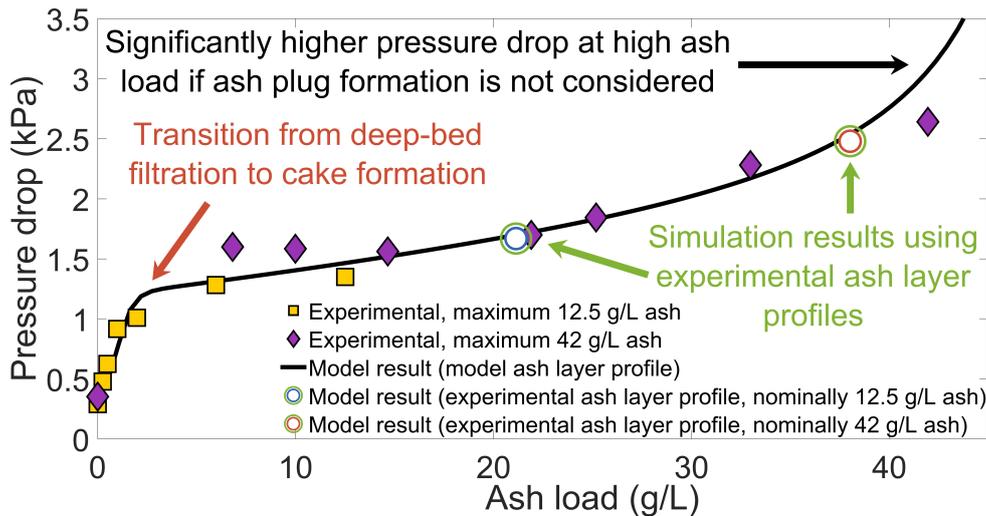


Figure 10: Pressure drop after regeneration versus the ash load accumulated in the filter. The pressure drop of the ash loaded filters predicted by the model is compared against experimental measurements reported by Sappok et al. [50]. The model describes lightly loaded filters adequately, but the pressure drop is overestimated at high ash loading if ash migration along the channel is neglected.

The calibrated model shows good agreement with the experimental data up to 40 g/L ash load. At ash loads above this, the model overestimates the pressure drop. The discrepancy is possibly related to the choice to neglect the axial transport of ash in the model. One consequence of this is that the model cannot describe the formation of an ash plug, and instead predicts an ash layer that is almost uniform in thickness. This is reasonable at low ash loads, but disagrees with experimental observations at high ash loads, which show

that the ash plug can occupy 32%–48% of the inlet channel [49].

Sappok [49] measured the ash layer profile in three inlet channels of the filters used in each experiment shown in Fig. 10. In order to investigate the impact of the ash layer, additional simulations were performed in which the experimental ash layer profiles were imposed on the model. The imposed profiles are shown in Fig. 11 and were taken as the average of the profiles measured in each filter. In addition, the ash load in each simulation was imposed based on the ash load implied by the volume of the imposed profile, corresponding to 21.1 g/L and 38.0 g/L, as opposed to the nominal ash loads of 12.5 g/L and 42 g/L respectively. It is hypothesised that the difference between the ash loads implied by the imposed profiles and the nominal ash loads arises due to variations between the channels in each filter. As a consequence of this, the pressure drops calculated by the simulations with the imposed ash layer profiles are discussed in the context of the pressure drop across the simulated channel-pairs, as opposed to the pressure drop across the entire filter.

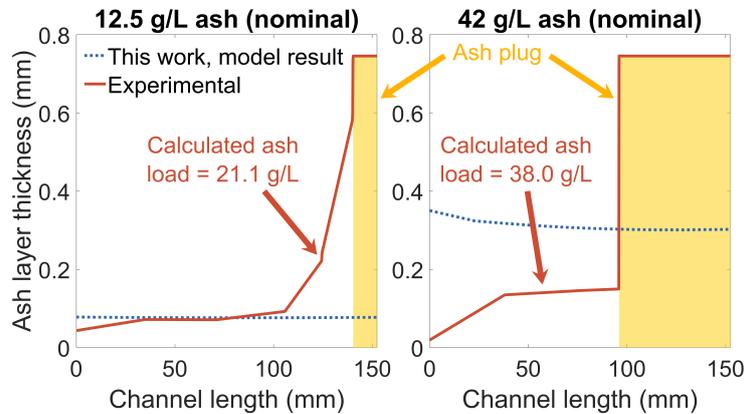


Figure 11: Ash layer profiles from experimental measurements and simulation results.

The hollow circles on Fig. 10 show the channel-pair pressure drops calculated after imposing the experimental ash layer profiles on the model. The predicted pressure drop for the lightly ash-loaded (nominal 12.5 g/L) filter is almost unchanged. This suggests that the pressure drop is insensitive to the shape of the ash layer profile at this ash load. On the other hand, the imposed ash layer profile had a significant effect on the simulation of the heavily ash-loaded (nominal 42 g/L) filter, resulting in a pressure drop of approximately 3.5 kPa (not shown).

In order to investigate the impact of thermal treatment on the pressure drop (in the next section), the simulation of the heavily ash-loaded (nominal 42 g/L) filter shown on Fig. 10 was re-calibrated to use a value of 350 nm for the size of the particles in the ash layer. It is noted that Sappok [49] found that the ash layer in the heavily loaded filter had a lower porosity (0.911) than that in the lightly loaded filter (0.946), which may be indicative of an increase in particle size due to sintering during the repeated regeneration events. By assumption, sintering during regeneration is excluded from the model, and the impact of its omission remains an open question. Finally, it should be emphasised that *all* other simulation results shown in Fig. 10 were generated using the original value of 220 nm for the size of the particles in the ash layer.

Fig. 12 shows the amount of ash accumulated in the filter wall and in the ash layer at the end of each loading-regeneration cycle. The data correspond to the model result (black line) shown in Fig. 10 and apply to both the 12.5 g/L and 42 g/L filters. The ash load in the filter wall reaches a maximum value of 0.12 g/L after four cycles. No further particle deposition within the filter wall occurs after this point, indicating that the filtration efficiency of the ash layer is now 100%. The total ash load in the filter at this point is estimated to be 2.2 g/L. Given that the filter wall is saturated at low ash loads, the ash plug (if it is present) would have negligible impact on the ash deposition in the wall.

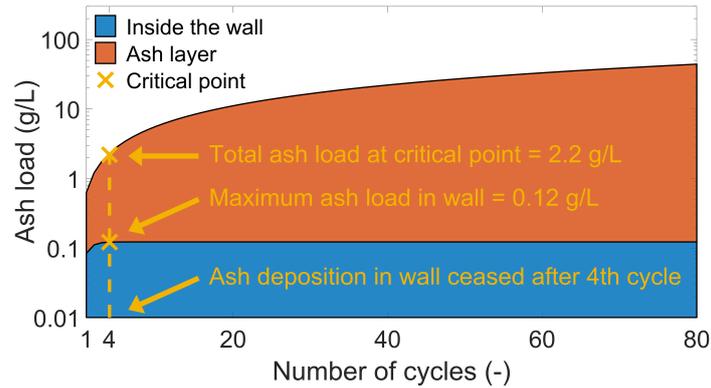


Figure 12: Ash load in the filter wall and in the ash layer at the end of each loading-regeneration cycle. The ash load in the filter wall reaches a maximum value of 0.12 g/L after four cycles. The total ash load at this point is 2.2 g/L.

3.3 Impact of thermal treatment

Sappok et al. [52] observed that the pressure drop across the ash-loaded filters decreased after the filters were heated in a furnace, with higher furnace temperatures resulting in larger decreases in the pressure drop. The model developed in this work was applied to investigate the mechanisms responsible for this decrease in the pressure drop.

Fig. 13 shows the experimental measurements of the pressure drop across the thermally treated filters versus the furnace temperature [52]. In addition, the figure shows the pressure drop over individual channel-pairs from each filter. The points at 650 °C were simulated using the imposed experimental ash layer profiles (these data were also shown in Fig. 10). The points at 700–900 °C were estimated based on the corresponding experimental measurements across the entire filter and the simulation results at 650 °C. It should be noted that the pressure drop over the channel-pair in the lightly loaded (nominal 12.5 g/L) filter is larger than the pressure drop measured across the entire filter. This is consistent with the difference between the ash load implied by the imposed ash layer profile (21.1 g/L) versus the nominal ash load (12.5 g/L), as discussed in Section 3.2.3. The discrepancy is less severe for the heavily loaded (nominal 42 g/L) filter.

Both filters show similar pressure drops for furnace temperatures of 900 °C and above. This suggests that the ash loading in the wall (which is the same for both filters, see Fig. 12) dominates the pressure drop and that the ash layer (which is different between

the two filters) has negligible impact at these conditions. One consequence of this is that the channel-pairs also show the same pressure drop as the filters under these conditions.

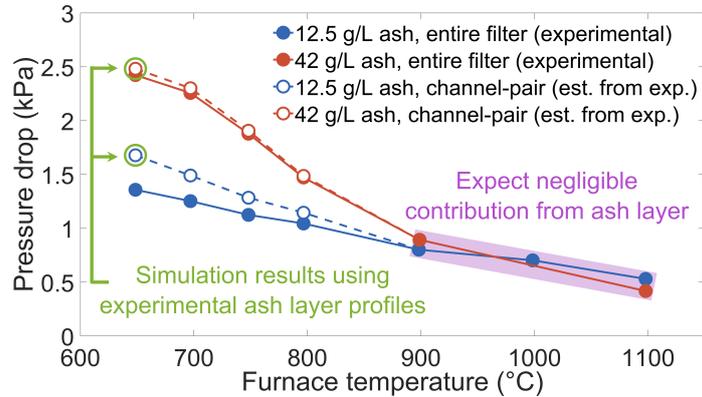


Figure 13: Pressure drop of thermally treated particulate filters versus furnace temperature. Solid circles: Experimental data [52]. Hollow circles: Estimates of the pressure drop across individual channel-pairs. The points at 650 °C were simulated using the imposed experimental ash layer profiles (these data were also shown in Fig. 10). The points at 700–900 °C were estimated based on the corresponding experimental measurements across the entire filter and the simulation results at 650 °C.

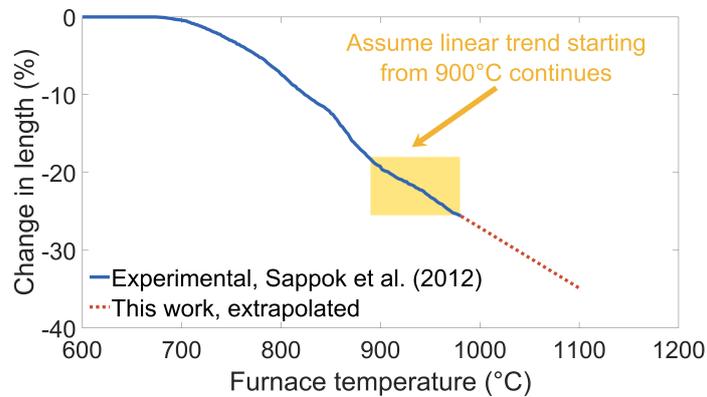
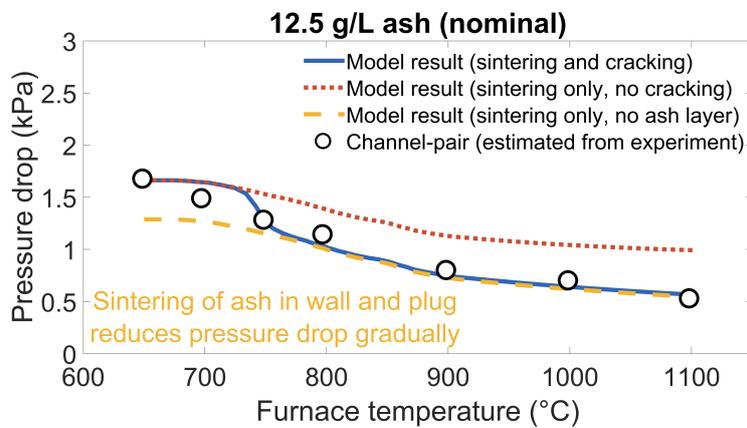


Figure 14: Experimental data showing dilatometer measurements of the change in the length of ash deposits retrieved from field DPFs versus furnace temperature [52]. These data are used to define the reduction in the bulk volume of the ash deposits induced by the thermal treatment. The volume reduction at high furnace temperatures is extrapolated from the experimental data under the assumption that the apparent linear trend continues.

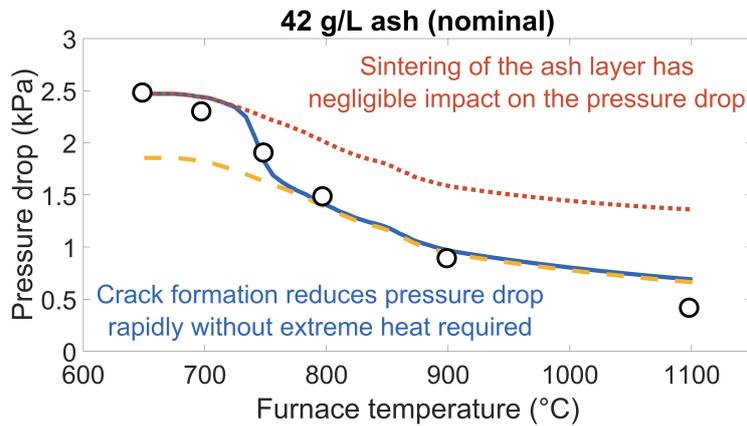
The thermal treatment model (see Section 2.4) uses the change in the bulk volume of the ash to define the sintering process variable X . In this work, the relationship between the bulk volume of the ash and the furnace temperature is known experimentally from dilatometer testing of ash samples removed from the filters [52]. Measurements were performed using ash from the filters studied with the accelerated loading system and from

filters installed on vehicles operating in the field. Unfortunately, only the samples from the vehicles in the field produced meaningful dilatometer results, and it is these data that are used here. The data are shown in Fig. 14. The data only extend up to 1000 °C, which is less than the maximum furnace temperature of 1100 °C. The data are therefore extrapolated, assuming that the linear trend observed above 900 °C continues up to 1100 °C.

Fig. 15 shows the results of simulations of thermally treated channel-pairs in the filters with nominal ash loads of 12.5 g/L and 42 g/L. The model shows good agreement with the pressure drop estimated from experimental data when both the sintering and cracking sub-models are enabled. The cracking model parameters were $\alpha = 55.33$ and $\beta = 0.00266$ for both cases shown in Figs. 15(a) and 15(b), and control the temperature at which cracking starts to occur in the ash layer. A corresponding sharp decrease in the pressure drop is observed between 700 and 800 °C. The pressure drops continues to decrease for furnace temperatures above 800 °C, albeit at a lower rate.



(a) Channel-pair from the filter with 12.5 g/L (nominal) ash load.



(b) Channel-pair from the filter with 42 g/L (nominal) ash load.

Figure 15: Pressure drop of thermally treated filter channel-pairs versus furnace temperature. Good agreement between the model and the pressure drop estimated from experimental data is observed when both sintering and cracking are considered by the model. The cracking of the ash layer causes a rapid decrease in the pressure drop between 700 and 800 °C.

Additional simulations were performed to investigate the impact of the ash layer. The model predictions with and without the ash layer sub-model enabled (blue solid lines versus yellow dashed lines in Fig. 15) are coincident for furnace temperatures above 800 °C. This indicates that the cracked ash layer makes no contribution to the pressure drop, such that the contribution of the ash to the pressure drop arises from the ash deposits in the wall and the ash plug for furnace temperatures above 800 °C. Comparison of the model predictions without the cracking sub-model and without the ash layer sub-model enabled (red dotted lines versus yellow dashed lines in Fig. 15) shows that the sharp decrease in pressure drop is not observed if crack formation is not considered, and that the difference in the pressure drop between the two cases is nearly independent of the furnace temperature. This implies that sintering has negligible effect on the flow resistance of the ash layer, such that sintering of the ash deposits in the wall and the ash plug is responsible for the decrease in pressure drop above 800 °C.

3.4 Discussion

Crack formation in the ash layer is responsible for a significant proportion of the reduction in the pressure drop observed in thermally treated filters. It is highly desirable for crack formation to occur because its full advantage can be achieved without excessive heating, minimising both the energy demand for the thermal treatment and the risk of damaging the filter. van de Lagemaat et al. [63] suggested that nanoparticle layers with porosity over 75% would be structurally unstable. Since ash layers typically have porosities greater than 85% [66], it is likely that crack formation will always occur when the ash layer is thermally treated. Future work should seek to substantiate this and establish the critical conditions for crack formation to occur in the ash layer. In addition, the change of filtration efficiency due to crack formation in the ash layer should be investigated.

Sintering of the ash deposits in the filter walls and the ash plug can further reduce the ash-induced pressure drop after the ash layer has cracked. Since the sintering behaviour of the ash depends on the composition of the lubricant oil, lubricants which leads to ash with low sintering onset temperatures may be favourable, such that a greater reduction in the filter pressure drop can be achieved for a given thermal energy input. On the other hand, severely sintered ash deposits may lead to permanent damage to the filter. Good control over the thermal treatment would be required if the ash sintered too easily.

Several aspects of the model merit further scrutiny. The formation of the ash plug was not considered in the simulations of ash loading. An accurate description of the ash plug formation process will likely improve the predictive power of the model. Experimental studies have shown that ash plug is formed by migration of ash deposits down the channel [38]. Sappok et al. [54] observed that ash migration can be induced by either gas flow in the channels or fragmentation of the soot cake layer. Whilst flow-induced ash migration has been modelled [54], no adequate quantitative model exists to fully describe ash plug formation. A deeper understanding of ash migration will benefit the development of ash management strategies.

Experimental data show considerable channel-to-channel variation in ash layer profiles [49]. While a multi-channel model may be able to capture the variations between chan-

nels, the development of such a model requires a good understanding of phenomena occurring within each individual channel. Future experiments may consider using a downsized filter with fewer channels to limit the variation in measurements across different channels. This would support the development of more accurate single-channel models and aid the development of multi-channel models.

Information about the change in the bulk volume of ash deposits is critical as substantiated in Section 3.3. In this work, measurements made using a dilatometer were used to describe the change in bulk volume of the ash deposits as a function of the temperature of the thermal treatment. Although this led to good agreement between the simulation results and experimental data, a dilatometer requires ash samples to be removed from the filter before testing. This risks altering the properties of the ash deposits and increases the uncertainty of the dilatometer results because the sintering behaviour depends on the initial morphology of the ash deposits [40]. The sintering behaviour of ash deposits may be better understood if measurements by X-ray chromatography [17] or Magnetic Resonance Imaging [4] were available. The application of non-destructive imaging techniques would eliminate the uncertainty introduced by removing ash from the filter.

4 Conclusions

A particulate filter model has been developed to simulate the change in the behaviour of ash-contaminated particulate filters when the filters are thermally treated by exposing them to high temperatures. The model was calibrated using experimental data for a clean filter, followed by data for a soot-loaded filter. The model was applied to simulate the accumulation of ash in the filter in order to estimate the spatial distribution of the ash deposits within the filter wall and the diameter of the particles in the ash layer. Finally, the impact of heat-induced sintering and cracking of the ash deposits was examined using the model.

The model suggests that crack formation in the ash layer leads to significant reduction of the pressure drop at relatively low temperatures. Most of the benefits due to crack formation were attained using a treatment temperature of 800 °C. Further reduction of the pressure drop at higher temperatures is controlled by the sintering of ash deposits within the porous wall of the filter and the ash plug.

The model has been shown to be capable of predicting the pressure drop of ash-loaded filters when the bulk volume of the ash deposits is known. The accuracy of the model prediction for thermally treated filters relies heavily on accurate measurements of the change in the bulk volume of the ash deposits. In this work it is shown that crack formation in the ash layer is critical to reduce the pressure drop across the filter. Therefore it is crucial to guarantee crack formation during the thermal treatment. Further work is required to determine in more detail the operating conditions and ash properties that induce crack formation.

Research data

Research data supporting this publication is available in the University of Cambridge data repository ([doi:10.17863/CAM.56872](https://doi.org/10.17863/CAM.56872)).

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Nomenclature

Upper-case Roman

A, B	Cracking model parameter (literature)	-
C	Stoke-Cunningham factor	-
D	Channel side length	m
D_f	Fractal dimension	-
E	Type space	-
F	Friction factor	-
Kn	Knudsen number	-
L	Length	m
N	Particle number density	$1/\text{m}^3$
N_{cord}	Coordination number	-
\dot{N}	Particle number density flow rate	$1/\text{m}^3\text{s}$
P	Pressure	Pa
Pe	Peclet number	-
Q	Filtration efficiency	-
R	Interception parameter	-
S	Index set	-
V	Volume	m^3
V_f	Volume filling factor	-
X	Progress variable for sintering	-

Lower-case Roman

d	Diameter	m
f	Kuwabara geometric function	-
g	Kuwabara geometric function	-
i, j	Index of volume element	-
m	Mass	kg
n	Number of particles	-
n_{ax}	Number of axial discretisation	-
n_w	Number of through-wall discretisation	-
t	Time	s
u, v	Velocity	m/s
w	Thickness	m
z	Index of particle size class	-

Upper-case Greek

Φ	Partition coefficient	-
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Lower-case Greek

$\alpha, \beta, \gamma, \theta$	Cracking model parameter	-
ε	Porosity	-
η	Single collector efficiency	-
κ	Permeability	m ²
μ	Dynamic viscosity	Pa s
ρ	Mass density	kg/m ³
ρ_w	Particle packing density in filter wall	kg/m ³
ψ	Percolation constant	-

Subscripts

agg	Aggregate
ash	Ash
cake	Soot cake
c	Denotes the cracking sub-model
ci	Inlet channel
co	Outlet channel
coll	Collector
D	Denotes diffusional filtration
d	Denotes damaged layer
frac	Denotes fractal diameter
mob	Denotes mobility diameter
p	Particle
plug	Ash plug
pm	Porous media

pore	Pore
pri	Primary particle
R	Denotes interceptional filtration
s	Denotes the sintering sub-model
w	Denotes parameters in the filter wall
0	Denotes an initial condition

Abbreviations

ACT	Asymmetric Channel Technology
DPF	Diesel Particulate Filter
GDI	Gasoline Direct Injection
GPF	Gasoline Particulate Filter
ICE	Internal Combustion Engine
PM	Particulate Matter
PN	Particle Number
WFM	Wall Flow Monolith
XRT	X-ray tomography

A Derivation of the sintering model

This section describes the derivation of eq. (31)

$$\epsilon_{\text{ash}} = 1 - \frac{1 - \epsilon_{\text{ash}0}}{d_{\text{ash}0}^2} d_{\text{ash}}^2,$$

which relates the porosity of the ash layer ϵ_{ash} to the size of the ash particles d_{ash} .

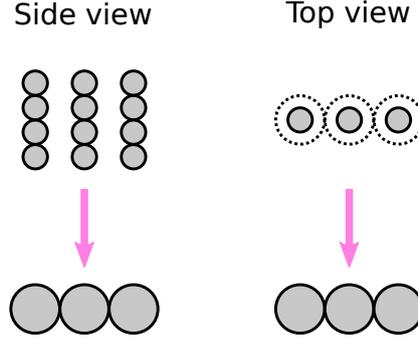


Figure A.1: Side view and top view of the particle columns representing the ash layer in the sintering model.

The sintering model approximates the ash layer as columns of equally sized particles as shown in Fig. A.1. It is assumed that the particles merge as sintering proceeds, until they eventually form a mono-layer of particles in point contact. The mono-layer is the maximum extent of the sintering that can be considered by this analogy. This state is denoted by the subscript “final” in the following equations. The initial state where sintering has not occurred is denoted by the subscript “0”. The porosity of the mono-layer ϵ_{final} is independent of the size of the particles

$$\epsilon_{\text{final}} = 1 - \frac{\frac{\pi}{6} d_{\text{p,final}}^3}{d_{\text{p,final}}^3} = 1 - \frac{\pi}{6}, \quad (\text{A.1})$$

where $d_{\text{p,final}}$ is the final particle diameter. The initial particle diameter $d_{\text{p}0}$ can be calculated based on conservation of the volume of the ash

$$d_{\text{p}0} = d_{\text{p,final}} n_{\text{p}0}^{-\frac{1}{3}}, \quad (\text{A.2})$$

where $n_{\text{p}0}$ is the initial number of particles in a column.

The initial porosity ϵ_0 can be derived by considering the particle columns from top view shown in Fig. A.1. The porosity of the particle columns scales proportionally with the cross-section area occupied by the ash particles. As a result, ϵ_0 can be expressed as

$$\epsilon_0 = 1 - \frac{\pi}{6} \frac{d_{\text{p}0}^2}{d_{\text{p,final}}^2} = 1 - \frac{\pi}{6} n_{\text{p}0}^{-\frac{2}{3}}. \quad (\text{A.3})$$

The following general equation is assumed to describe intermediate levels of sintering

$$\varepsilon = 1 - \frac{\pi}{6} \frac{d_p^2}{d_{p,\text{final}}^2} = 1 - \frac{\pi}{6} \frac{1}{d_{p0}^2 n_{p0}^{\frac{2}{3}}} d_p^2. \quad (\text{A.4})$$

An expression of n_{p0} can be found by rearranging eq. (A.3)

$$n_{p0}^{\frac{2}{3}} = \frac{\pi}{6} \frac{1}{1 - \varepsilon_0} \quad (\text{A.5})$$

Eq. (31) can be derived from eq. (A.4) by substituting eq. (A.5)

$$\varepsilon_{\text{ash}} = 1 - \frac{1 - \varepsilon_{\text{ash}0}}{d_{\text{ash}0}^2} d_{\text{ash}}^2. \quad \square$$

B Axial distribution of ash deposits within the wall

Fig. B.1 shows the amount of ash accumulated within the filter wall during the first four ash accumulation cycles of the simulations discussed in Section 3.2.3. It can be observed that the ash deposition in the wall is practically uniform in the axial direction for every cycle. The maximum deviation between the volume element with the highest ash load and that with the lowest ash load is 0.9% after the first cycle. This deviation decreases as more cycles are executed. The maximum deviation after four cycles is 0.6%. This substantiates the assumption in the sintering model for the ash plug (see Section 2.4.1) that the axial variation of the ash deposits in the wall is negligible.

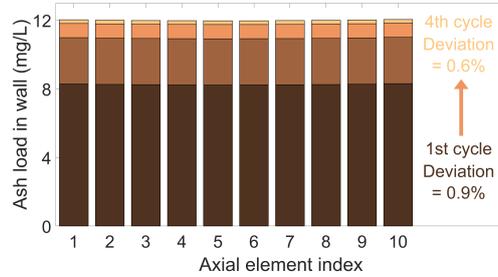


Figure B.1: *Ash load accumulated in the porous wall over the first four ash accumulation cycles. The ash deposit distribution is practically uniform. The uniformity increases with increased ash accumulation.*

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