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Machine learning approaches to the prediction of powder flow behaviour of pharmaceutical materials from physical properties†

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Understanding powder flow in the pharmaceutical industry facilitates the development of robust production routes and effective manufacturing processes. In pharmaceutical manufacturing, machine learning (ML) models have the potential to enable rapid decision-making and minimise the time and material required to develop robust processes. This work focused on using ML models to predict the powder flow behaviour for routine, widely available pharmaceutical materials. A library of 112 pharmaceutical powders comprising a range of particle size and shape distributions, bulk densities, and flow function coefficients was developed. ML models to predict flow properties were trained on the physical properties of the pharmaceutical powders (size, shape, and bulk density) and assessed. The data were sampled using 10-fold cross-validation to evaluate the performance of the models with additional experimental data used to validate the model performance with the best performing models achieving a performance of over 80%. Important variables were analysed using SHAP values and found to include particle size distribution D10, D50, and aspect ratio D10. The very promising results presented here could pave the way toward a rapid digital screening tool that can reduce pharmaceutical manufacturing costs.

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Introduction

In recent years, the pharmaceutical industry has increasingly explored Industry 4.0 technologies with the goal of using digital design to improve the prediction of bulk materials properties and minimise the amount of time and material required in early-stage development.¹ Machine learning (ML) models can help inform and minimise extensive early-stage development experiments, water and power consumption.

Understanding powder flow of pharmaceutical materials is necessary when developing robust manufacturing processes.² Powder flow, typically characterised by the flow function coefficient (FFc), impacts the manufacturability of drug compounds, and optimising powder flow improves the likelihood that streamlined manufacturing processes can be developed successfully and operated consistently. For example, powder flow has a significant impact on steps involving tablet formation. Tablets can be manufactured using several techniques such as direct compression (DC), wet granulation (WG),

or roller compaction (RC).³ Using DC for tablet manufacture requires that material properties, such as blend uniformity, compactability, and lubrication are tightly controlled.⁴ By contrast, WG and RC are used to improve powder flow and compactability prior to tablet compression. However, these techniques have some disadvantages such as the use of heat in RC and the use of binding agents and secondary wetting in WG. Moreover, WG and RC are more expensive and time-consuming. Thus, DC offers a streamlined process with fewer steps than WG and RC for example, however, to use DC, powders must flow well.

The ability to predict flow properties of powders or powder blends using straightforward routine measurements is therefore of increasing importance.⁵ A variety of particle and bulk properties are known to affect flowability, powder behaviour and process performance in DC. For example, particle size distribution (PSD) has a significant impact on powder behaviour,⁶ and hence, PSD has traditionally been a key property considered when predicting powder behaviour.⁷

However other physical properties can also affect powder flow behaviour and process performance, including shape, surface texture, surface area, density, cohesivity, adhesivity, elasticity, plasticity, porosity, charge potential, hardness, and hygroscopicity.⁸ These physical properties can have complex effects on powder behaviour, which have been described in many publications.^{9–11}

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PSD has a significant impact on powder flowability, however the relationship between these properties is not directly predictable.^{7,12–14} The effect of PSD in manufacturing processes such as compression has been demonstrated previously, and therefore, the effects of PSD should be carefully studied to ensure good manufacturing properties to achieve the desired dosage form.^{15–17} The guidelines proposed by Leane *et al.* indicated that powders with a PSD D90 smaller than 1000 μm are ideal for direct compression, but no other PSD targets were established for other manufacturing techniques, such as WG or RC.¹⁸

Traditionally, powder flow has been measured by experimental methods, such as angle of repose, bulk density, Carr's compressibility index, Hausner ratio, ring shear tester or the use of a powder rheometer. However, these methods are time consuming and require reasonable amounts of material for each test carried out. Different approaches to estimating powder flow have been explored in the literature. Sandler and Wilson studied packing efficiency by measuring particle size of granular intermediates using Principal Component Analysis (PCA).¹⁹ Megarry *et al.* used a big-data approach using the shear cell test to better understand of the typical flow properties of pharmaceutical materials.²⁰ A Partial Least Square (PLS) approach using particle size and shape distributions⁷ determined the relevance of particle shape in powder flow prediction. Capece *et al.* explored how the granular Bond number correlates to the FFC and illustrated the complexity involved in predicting powder behaviour.^{21,22} Statistical modelling techniques published by Barjat *et al.* focused on the prediction of flowability for LIW feeders.²³ The studies described here

established the feasibility of the prediction of powder flow using digital design, but the models developed cannot be directly applied to real-world manufacturing challenges due to either particle attribute restrictions or limited data availability.

Here, we present an assessment of ML modelling for predicting FFC as a reliable, generally applicable method for a wide range of pharmaceutical powders. The proposed models aim to predict the FFC of new materials, using the simple to measure particle properties. Usually, materials that have a value of FFC greater than 10 are considered free-flowing,²⁴ and therefore, easy to manufacture. Here, by combining ML models with experimental measurements, the amount of material and time required to estimate powder flow was significantly decreased from 30 g and 2 hours to 2 g and 5 min. The intended application of this model is in die filling, where dynamic powder flow predominates. Implementing such models in the early stages of drug development could help target particle engineering or improve decision-making for formulation and processing technology selection while reducing the time and material required.

Materials and methods

Materials

The materials in this study, including APIs and excipients, are listed in Table 1 below. These materials were included in the training dataset as both individual materials and blends, resulting in 112 observations used in ML model development.

Blends were made for ibuprofen 50, paracetamol powder, paracetamol granular special, mefenamic acid, and ibuprofen

Table 1 Materials included in the training data set

Material	Supplier	Material	Supplier
4-Aminobenzoic acid	Sigma-Aldrich	Ibuprofen 70	Sigma-Aldrich
Ac-Di-Sol	Dupont	Lactose	Sigma-Aldrich
Acetazolamide	Sigma-Aldrich	Lidocaine	Sigma-Aldrich
Affinisol	Dupont	Lubritose AN	Kerry
Aspirin	Sigma-Aldrich	Lubritose mannitol	Kerry
Avicel PH-101	Dupont	Lubritose MCC	Kerry
Avicel PH-102	Dupont	Lubritose PB	Kerry
Benecel K100M	Dupont	Lubritose SD	Kerry
Benzoic acid	Sigma-Aldrich	Magnesium stearate	Roquette
Benzylamine hydrochloride	Sigma-Aldrich	Magnesium stearate	Sigma-Aldrich
Bromhexine hydrochloride	Sigma-Aldrich	Mefenamic acid	Sigma-Aldrich
Caffeine	Sigma-Aldrich	Methocel MC2	Colorcon
Calcium carbonate	Sigma-Aldrich	Microcel MC-102	Roquette
Calcium phosphate dibasic	Sigma-Aldrich	Microcel MC-200	Roquette
Cellulose	Sigma-Aldrich	Nimesulide	Sigma-Aldrich
Croscarmellose Na	Dupont	Paracetamol granular special	Sigma-Aldrich
D-Glucose	Sigma-Aldrich	Paracetamol powder	Sigma-Aldrich
D-Mannitol	Sigma-Aldrich	Pearlitol 300DC	Roquette
D-Sorbitol	Sigma-Aldrich	Plasdone povidone	Ashland
Dropropizine	Sigma-Aldrich	Plasdone K29/32	Ashland
FastFlo 316	Dupont	Phenylephedrine	Sigma-Aldrich
FlowLac 90	Meggle Pharma	Roxithromycin	Sigma-Aldrich
Granulac 140	Meggle Pharma	S-Carboxymethyl-L-cysteine	Sigma-Aldrich
Granulac 230	Meggle Pharma	Soluplus	BASF
HPMC	Sigma-Aldrich	Span 60	Sigma-Aldrich
Ibuprofen 50	BASF	Stearic acid	Sigma-Aldrich



Table 2 The composition of binary blends. All binary blends included Fast Flo 316 and one of the following APIs: ibuprofen 50, paracetamol granular special, paracetamol powder, mefenamic acid, calcium carbonate

Binary mixture	Drug loading	Fast Flo 316
Low drug dosage	5%	95%
Medium drug dosage	20%	80%
High drug dosage	40%	60%

Table 3 The composition of multi-component blends. All multi-component blends included Fast Flo 316, one of the following APIs: ibuprofen 50, paracetamol granular special, paracetamol powder, mefenamic acid, calcium carbonate, and the remaining 25% of a combination of 20% Avicel PH-102, 3.5% croscarmellose sodium, and 1.5% magnesium stearate

Multicomponent mixture	Drug Loading	Fast Flo 316	Other excipients
Low drug dosage	5%	70%	25%
Medium drug dosage	20%	55%	25%
High drug dosage	40%	35%	25%

sodium salt at different drug loadings (5%, 20%, 40%) for binary mixtures with FastFlo 316 and multicomponent mixtures including FastFlo316, croscarmellose sodium, Avicel PH-102, and magnesium stearate. The blends were prepared using a 1 L bin blender (Pharmatech AB-105, UK). The composition of the blends is described in Tables 2 and 3.

Experimental methods

Particle size and shape – QICPIC, Sympatec: image analysis characterisation. Particle size and shape were analysed using QICPIC, Sympatec. QICPIC captures the physical properties of the particles by using a high-speed camera that performs dynamic image analysis. The measurements were done in triplicate. PSD is represented with the values D10, D50, D90, and Sauter Mean Diameter (SMD). Among the particle shape descriptors that can be analysed using this equipment, aspect ratio and sphericity were selected as they were proven to have the biggest impact on powder flow.

The instrument used to characterise particle size and shape presents some limitations such as a lower sensitivity in the detection of the shape of fine particles.

Surface area and energy measurements – Surface Energy Analyser. The surface area and energy of 35 materials were measured using inverse gas chromatography (iGC – Surface Energy Analyser, Surface Measurement Systems Ltd.). For the surface energy measurements, the method selected was Dorris-Gray.

Powder flow and bulk density – Powder Rheometer FT4. An FT4 Powder Rheometer (Freeman Technology, Malvern, UK) was used to carry out the shear cell test to measure the FFC and the bulk density of the different materials. The FT4 Powder Rheometer differs from other powder flow testers due to the

industrial value that it provides by assessing dynamic powder flow, bulk, and shear properties. The consolidation stress was set at 9 kPa, and the normal stress for shearing was set at 7, 6, 5, 4, and 3 kPa. The sample is sheared to obtain five yield points at different normal and shear stress. The 25 mm × 10 ml split-vessel was selected to carry out triplicates of each sample using the Freeman Technology user manual.²⁵ The output data is the powder's yield locus which is calculated from the relationship between the shear stress and the normal stress. By fitting Mohr stress circles to the yield locus, the major principal stress (σ_1) and unconfined yield strength (σ_c) are defined; and the ratio between them is the flow function, that can be used to rank flowability.

Powders with a value of flow function coefficient below 4 have poor flow; between 4 and 10, they are fairly flowable; and above 10, free-flowing.²⁴ The flow function coefficient has been correlated with the manufacturing process by the Manufacturing Classification System,^{18,26} assigning to each flow function coefficient and drug loading a suitable manufacturing process.

For this work, the consolidated bulk density was also measured using the FT4 Powder Rheometer (Freeman Technology Ltd.). The results of bulk density calculated using this method are generally more accurate and reproducible than the conventional measurements, such as the measurement in a graduated cylinder, or in a volumeter.^{25,27} The test was repeated at least 2 more times, using different samples each time, until the results were consistent, and the average value was calculated and taken as the result.

Machine learning methods

Unsupervised and supervised ML models were built to investigate FFC prediction from particle size distribution and particle shape.

Unsupervised learning approaches. For unsupervised learning, Principal Component Analysis (PCA), hierarchical clustering and Louvain clustering were applied to the data. PCA was done using Anaconda Spyder (Scientific Python Development Environment), matplotlib, and sci-kit learn. Louvain clustering was done using Orange Data Mining, an open-source data visualisation toolkit written in Python, Cython, C++, and C. For Louvain clustering, the data were normalised and a 3-component PCA was applied as pre-processing. To plot the graph, the distance metric used was Euclidean and 30 neighbours (details in Section 1.1 of ESI†).

Classification models. Support Vector Machines (SVM), Random Forests (RF), neural networks, Naïve Bayes (NB), k-Nearest Neighbours (kNN), Logistic Regression (LR), and AdaBoost (AB) were all investigated for classification capabilities of powder flow into three categories: cohesive, easy-flowing, free-flowing (as defined in the Experimental methods section). Python was used to write the code for the algorithms, using libraries including pandas, NumPy, matplotlib, and sci-kit learn (details in Section 1.2 of ESI†).

The performance of each algorithm was evaluated using the following: area under the curve receiver operating



characteristics (AUC–ROC), precision and recall.²⁸ These metrics were calculated from the corresponding model's confusion matrix. As classification accuracy (CA) can be misleading when a class imbalance is present,^{29,30} AUC–ROC was used to evaluate model performance with a maximum possible of 1 (details in Section 1.3 of ESI†).

112 pharmaceutical powders were included in these models, sampled using 10-fold cross-validation to test the performance.

Regression models. Linear regression (LR),³¹ Gradient Boosting (GB),^{32,33} Random Forest (RF)^{34,35} and AdaBoost (AB)³¹ were used for FFC value prediction. Python, sci-kit learn and Orange Data Mining software were used to implement these models. 106 observations were included with an 80 : 20 train-test split.

Model interpretability

Shapley Additive Explanation (SHAP)^{36,37} values were used to increase the interpretability of the models here and move away from the lack of understanding behind model decision making. This method has been used in this paper for both global interpretability, *i.e.* to identify the most important variables during training, and for local interpretability, *i.e.*, to understand how the model made the prediction for a selected test powder.

External validation

External validation was used to assess the performance of both the classification and regression models as standard practice to demonstrate the applicability of the model in unseen data. Prior to the test/train split, 8 materials were removed from the data set. These 8 “unseen” pharmaceutical powders were used for external validation of the highest performing classification and regression models (neural network and RF for classification and CATboost GB for regression). External validation was done for both the neural network and RF classification models due to possible overfitting of the data for the neural network classification model (see Results and discussion section, Fig. 7 and 9).

Results and discussion

Particle size and shape – QICPIC image analysis characterisation

Particle size distribution (PSD) by QICPIC image analysis. 112 systems were analysed using the QICPIC instrument including 30 active pharmaceutical ingredients, 43 excipients and 40 blends (list in ESI†) (Table 4).

Particle shape by QICPIC image analysis: aspect ratio and sphericity distribution. The aspect ratio distribution and

Table 4 Particle size distribution results, including the range of values and the median value for each parameter

Parameter	Range of values (μm)	Median (μm)
D10	9–225	54.84
D50	25–644	149.19
D90	53–1892	328.87
Sauter Mean Diameter (SMD)	19–393	94.63



Fig. 1 The distribution of aspect ratio values across the materials included in the training data set. Values presented here are the mean values from three measurements.



Fig. 2 The sphericity distribution of the materials included in the training dataset. Values presented here are the mean values from three measurements.

sphericity distribution were included as these parameters were found to have high feature importance scores in the ML models. Fig. 1 and 2 show the distribution of aspect ratio and sphericity for the materials tested in this work. As seen in Fig. 1, most of the materials have an aspect ratio between 0.6 and 0.8.

Fig. 2 shows that most of the materials had sphericity greater than 0.5 with 68 of the materials having a sphericity value between 0.6 and 0.8.³⁸

Surface area and energy measurements – Surface Energy Analyser

A representative sample of 35 powders were selected for surface area and surface energy measurement in the training dataset. Surface energy parameters measured were specific surface energy (mJ m^{-2}), surface energy (mJ m^{-2}), and dispersive surface energy (mJ m^{-2}) at 0%, 3%, 5%, and 10% of coverage (see ESI†) (Table 5).

Powder flow and bulk density

The bulk density and powder flow of the 112 pharmaceutical powders were analysed using Powder Rheometer FT4 – Freeman



Table 5 The surface area and surface measurements for the 35 materials analysed

Parameter	Range of values	Mean
Surface area	0.17 to 2.76 m ² g ⁻¹	0.64 m ² g ⁻¹
Specific surface energy	2.94 to 16.81 mJ m ⁻²	7.07 ± 0.48 mJ m ⁻²
Surface energy (com)	0.06 to 140.73 mJ m ⁻²	41.62 ± 0.66 mJ m ⁻²

**Fig. 3** The distribution of bulk density of the pharmaceutical powders of study.**Table 6** The number of observations in each range of interest of flow function coefficient

Flow function coefficient	Powder behaviour	Number of observations
FFc < 4	Cohesive	29
4 < FFc < 10	Easy-flowing	32
FFc > 10	Free-flowing	51

Technology Ltd. (see Fig. 3 and Table 6). Both types of measurements were done in triplicate.

Using unsupervised and supervised learning methods to predict powder flow

Unsupervised algorithms. PCA, Louvain and hierarchical clustering analysis were performed using Orange Data Mining software to determine if the resulting clusters corresponded with powder flow behaviour. In PCA, two principal components only accounted for 45% of the variance. The number of components was increased incrementally to 6 where 88% of the variance was accounted for. Louvain clustering showed the data clustered into 4 groups (see Fig. 4), and hierarchical clustering resulted in 3 groups of data. For all methods, groups did not correlate with powder flow behaviour.

Powder flow classification by a variety of supervised learning methods. Classification models were also implemented to predict powder flow as described in the Methods section. Classes were defined according to the FFc, as follows: cohesive (FFc ≤ 4); easy flowing (4 < FFc < 10); and free-flowing (FFc ≥ 10). Model features included particle size distribution, particle

**Fig. 4** Louvain clustering analysis.

shape distribution, bulk density, and powder flow. Surface area and surface energy data were later added, and the performance of the two different feature sets was compared.

Two types of models, namely a single-step and two-step classification, were investigated using supervised algorithms described in the Machine learning methods section. The first model developed a single-step classification in which materials were classified into one of the three FFc classes described above. The performance of this classification was assessed by calculating AUC-ROC (see Fig. 5). The highest performance achieved was by the multilayer perceptron neural network model (0.823). For classes 1 and 3, over 60% of the instances were correctly classified; however, for class 2, less than 45% of the materials were correctly classified by the model. The model therefore appeared to be better at predicting the FFc classes of cohesive and free-flowing materials but struggled to classify the easy-flowing²⁶ materials across the transition from cohesive to free flowing.

As the MLP neural network confusion matrix indicated that easy-flowing materials were difficult to distinguish from free-flowing materials (see Section 4 of ESI[†]). A two-step classification model was developed as following Jenike's classification of powder flow:²⁴ Step 1 classified materials into free-flowing (FFc

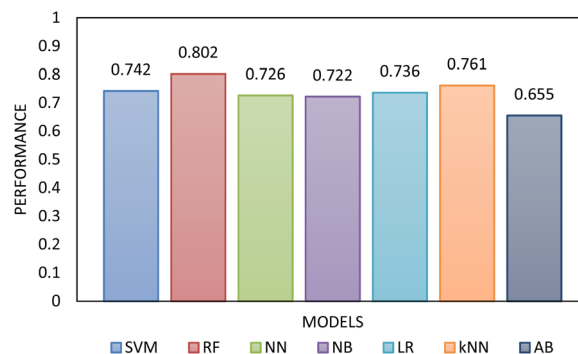
**Fig. 5** The AUC-ROC performance analysis of the single-step model compared to each other evaluated using 10-fold cross-validation. The highest performance was achieved by RF.



Fig. 6 The performance of the classification algorithms included in step 1 of the two-step model evaluated by 10-fold cross-validation.

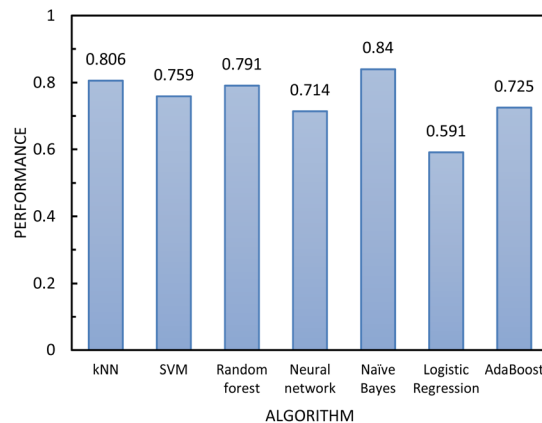


Fig. 8 The performance of the classification algorithms included in step 2 of the two-step model evaluated using 10-fold cross-validation.

> 10) or non-free-flowing ($FFc < 10$), cohesive and easy-flowing powders were included in the latter category. Step 2 classified the material into cohesive materials ($FFc < 4$) and non-cohesive materials ($FF > 4$), easy-flowing and cohesive powders were included in the latter category. According to the literature, easy and free-flowing powders are suitable for manufacturing with free-flowing powders being most suitable for direct compression.¹⁸ The performance of the algorithms included in step 1 and step 2 was again assessed using AUC-ROC (see Fig. 6 and 8). The results showed that by separating the classification decisions, the two-step model was able to perform better than the previous model. This improvement in the performance of the two-step model could be explained by considering that the imbalanced training dataset used for the single-step model affected the performance of the model, and when the dataset was split into subconsequent steps, the detrimental impact of the imbalanced data was minimised.

In determining which algorithm should be used for external validation, we prioritised model performance in step 1 as this classification step (free- vs. non-free-flowing) is more impactful for determining manufacturability than the classification in step 2 (cohesive vs. non-cohesive). For step 1, the neural network model had the highest performance (0.835), followed by RF (0.817). The neural network model was initially used for external validation (see ESI†). However, since the external validation classification accuracy was significantly worse (62.5%) than the classification accuracy for the test set, we hypothesize that the

		Predicted		Σ
		Non-free-flowing	Free-flowing	
Actual	Non-free-flowing	5	0	5
	Free-flowing	1	2	3
Σ		6	2	8

Fig. 7 External validation performed for the RF model. 87.5% of the materials were correctly classified.

		Predicted			Σ
		Cohesive	Easy-flowing	Free-flowing	
Actual	Cohesive	16	0	13	29
	Easy-flowing	0	29	3	32
	Free-flowing	6	13	32	51
Σ		22	42	68	112

Fig. 9 Step 1 and step 2 RF model confusion matrices combined as evaluated by 10-fold cross validation.

neural network algorithm was overfitting the data. As the model with the next highest performance, the RF model for both step 1 and 2 was used for all remaining external validation.

The RF confusion matrices for step 1 and step 2 have been combined to have a better overview of the performance of the two-step model (see Fig. 9).

Surface area and surface energy data were also added to the training set of the single-step and the two-step models because

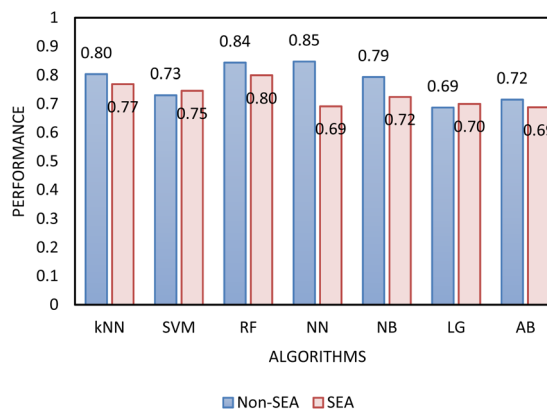


Fig. 10 Comparison of the performance evaluated by AUC-ROC of the classification learning algorithms for the two-step models with (SEA) and without (non-SEA) the inclusion of surface area and surface energy in the training dataset.



it has been previously shown that surface parameters have a significant influence on powder behaviour.^{39,40} The addition of these parameters resulted in a decrease in model performance for all algorithms, except kNN in step 1, and SVM and LR in step 2. A previous publication also showed that when improved particle size and shape data are available, the addition of surface area and surface energy data does not translate into an improvement of the performance of the model for the prediction of powder flow.²³ The decrease in performance due to the addition of more data can be a result of the small correlation between surface area and surface energy with powder flow, because the information introduced to the model is effectively noise. This result suggests that powder flowability is more strongly dependent on size and shape than it on surface area and surface energy. As the addition of these parameters did not improve performance, they were not included in later training datasets (Fig. 10).

Regression models. For regression models the dependent variable (or predicted response) was the FFC value or the reciprocal of the FFC. All regression models performed poorly

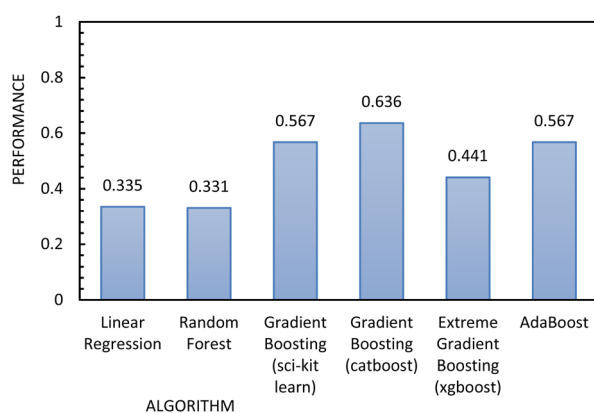


Fig. 11 Regression metrics to evaluate the performance of the algorithms used to build the regression model, using FFC as the independent variable.

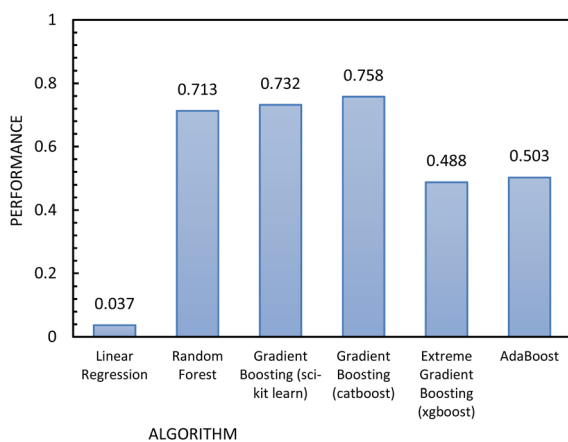


Fig. 12 Regression metrics to evaluate the performance of the algorithms used to build the regression model, using 1/FFC as the independent variable.

		Predicted			Σ
		Cohesive	Easy-flowing	Free-flowing	
Actual	Cohesive	0	1	1	2
	Easy-flowing	0	2	1	3
	Free-flowing	0	0	3	3
Σ		0	2	5	8

Fig. 13 External validation confusion matrix for the combined step 1 and step 2 RF models.

for FFC prediction (see Fig. 11). The reciprocal of the FFC was then calculated and used as the dependent variable. This transformation was carried out to emphasise small differences in low FFC values. For example, the difference between the behaviour of two materials with FFC 3 and 6 is more significant than the difference in the behaviour between two materials with an FFC of 13 and 16.²⁶

Fig. 12 shows the results of the regression models that have 1/FFC as the dependent variable with performances evaluated by 10-fold cross-validation. From these results, we see that using 1/FFC decreased prediction error compared to the models that predict FFC directly (see Fig. 11). For these models, CatBoost exhibited the best performance, with an R^2 value of 0.758, and an RMSE of 0.069.

External validation: classification models. As the two-step model had the best performance, external validation was done for this model (see Fig. 13). For the validation, 8 previously unused materials were analysed, including 2 cohesive, 3 easy-flowing, and 3 free-flowing materials. The external validation resulted in 5 of the 8 materials being correctly classified.

SHAP values in the RF classification model. SHAP values were calculated to help understand the predictions from the external validation.³⁷ This method improves the interpretability of the predictive models by showing feature importance



Fig. 14 Feature importance analysis for the RF model in step 1. The features are ranked based on their absolute mean SHAP score. The impact of each feature on each class is represented using colours and hence, the impact of the prediction on the output non-free flowing is represented in blue and the impact of the output free flowing is represented in red.





Fig. 15 Feature importance analysis for the RF model in step 2. The features are ranked based on their absolute mean SHAP score. The impact of each feature on each class is represented using colours and hence, the impact of the prediction on the output non-free flowing is represented in blue and the impact of the output free flowing is represented in red.

analysis. SHAP values can also be calculated to improve interpretability for individual predictions, *i.e.* for a given powder, what factors affected the model's predicted classification.

The feature that had the biggest impact on model performance in step 1 PSD D10 (*i.e.*, 10% of the particles have a particle size smaller than this value; Fig. 14). Therefore, this analysis indicated that the model's prediction between free-flowing and non-free-flowing powders was impacted significantly by the presence of fines in the material as captured in the elevated impact of the D10 value.

For step 2, the feature importance analysis is also calculated for the neural network model. Fig. 15 shows that, as for the step 1 classifier, PSD D10 is also the most important material feature. Other features that had high importance scores were PSD D50 and PSD D90 (particle size distribution 50% and 90% percentile, respectively). These were also important parameters for Step 1.

SHAP values were calculated using SHAP python library for one of the powder samples that were misclassified in the external validation to examine why the prediction was incorrect. The specific powder was a cohesive material that the external validation classified as free-flowing. This powder's prediction was chosen for SHAP local analysis since the misclassification of a non-free flowing powder as free flowing would result in a waste of both time and material in investigating the manufacturability of this powder. Step 1 classified this powder with a 67% probability (see Fig. 16) as free-flowing, but its measured FFc was 1.9.



Fig. 16 SHAP force plot of the prediction of the cohesive material from the external validation that was classified as free-flowing by step 1. The red bars of the plot show the drivers that are pushing the classification toward free-flowing and the blue bars show the drivers that are pushing the classification towards non-free flowing.

Table 7 Results of the external validation performed with the regression model, setting the target variable first as "FFc", and then as "1/FFc"

Actual FFc	Predicted FFc	Actual 1/FFc	Predicted 1/FFc
1.90	15.94	0.526	0.127
2.28	15.85	0.438	0.203
7.42	8.24	0.135	0.148
7.46	9.73	0.134	0.053
8.17	15.83	0.122	0.088
32.14	13.78	0.031	0.079
38.21	27.84	0.026	0.075
23.00	17.27	0.043	0.040

Fig. 16 also shows that the feature that had the biggest impact on the model output was again the PSD D10 value of the sample (225.11 μm). This value of PSD D10 is significantly higher than the mean value of PSD D10 of the training set (shown in Table 4). Furthermore, the material displayed high sphericity with a high value of aspect ratio D90 (0.98). Therefore, since the powder had large and spherical particles, these properties may have resulted in this misclassification. Adding additional training data with a wider range of different combinations of particle size and shape with varying bulk properties would help avoid such misclassifications in future models.

From the SHAP value analysis, the model here may slightly inflate the importance of D10 when compared with sphericity



Fig. 17 (a) Correlation of the actual FFc values and the predicted FFc values; (b) correlation between the actual 1/FFc values and the predicted 1/FFc.



values, as the size and aspect ratio were the most important factors in both the correct and incorrect classifications of the materials. Thus, retraining models with more materials with a training set with a greater variance in sphericity could improve the performance.

External validation: regression models. Although the regression models did not perform as well as the classification models, a further exploration of the regression models was carried out to better understand how the performance of these models could be improved. The same new, external dataset that was used in the previous section was used to validate the regression models. Here, the FFC and 1/FFC values for the 8 materials were predicted using CatBoost GB (the regression model with the highest performance) (see Fig. 12 and Table 7).

The prediction against 1/FFC as dependent variable ($R^2 = 0.5$) was better than the prediction of FFC ($R^2 = 0.37$), although neither result was satisfactory (Fig. 17).

Conclusions

Implementing ML models in the early stages of drug development can help determine suitable manufacturing strategies for a given material and provide rapid digital screening tools for advanced pharmaceutical development. In this work, FFC classes of pharmaceutical materials were predicted from routine, widely available, material-sparing analytical measurements. The 112 materials analysed exhibited a wide range of particle size distributions, particle shape distributions, and bulk densities and covered 3 classes of FFC that reflect what is captured in the literature.²⁴

This work suggests that particle size and shape distribution measured with dynamic image analysis are sufficient to enable the prediction of flow properties. The best performing model presented in this work was achieved by the combination of RF models for step 1 and step 2, with over 80% probability of distinguishing between classes for each step. Further improvements to model performance could be made with more data from cohesive materials as this would help address class imbalance in the training dataset. Additionally, including training data with different combinations of particle size and shape with differing bulk behaviour could also reduce misclassifications in future models. The FFC boundaries of the classes of powder flow could also be adapted to specific industry needs; for example, optimal FFC values will vary depending on the different pieces of equipment that might be available. In this work, propagation of analytical measurement error has not been included in the model training, and this research angle could be interesting to explore in further work. Moreover, the model could be extended to inform formulation optimization or even to provide a performance target for particle engineering efforts to develop materials for direct compression.

The ML model's implementation enables the prediction of the material flow properties (FFC) from size and shape allowing early decision-making regarding manufacturing route selection. Although there are more sophisticated techniques to capture particle size and shape data, the consideration of the whole

particle size and particle shape distribution may allow a better understanding of the data and of the relationship between particle size and shape and powder flow, resulting in a better predictive model. Moreover, the model could be extended to inform formulation optimization or even to provide a performance target for particle engineering efforts to develop materials for direct compression. Implementation of the models presented here in industry applications could save time and effort in early-stage development. The work presented in this paper illustrates the benefits of implementing digital design workflows for the prediction of material properties in the pharmaceutical industry where the availability of data is often limited. This work highlighted multiple potential applications that could result from increasing the available FAIR data in this industry and how it can help to digitalise pharmaceutical manufacturing.

Data availability

All data underpinning this publication are openly available from the University of Strathclyde KnowledgeBase at <https://doi.org/10.15129/3005493b-a125-4a5d-8599-525ce952facf>. The code supporting this article has been uploaded as part of the ESI.†

Author contributions

Laura Pereira Diaz: writing – original draft, data generation, data curation, investigation, methodology, validation. Cameron Brown: review and editing, supervision. Ebenezer Ojo: samples of pharmaceutical mixtures. Chantal Mustoe: review and editing. Alastair Florence: review and editing, supervision.

Conflicts of interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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References

- 1 J. Maier, *UK Industrial Digitalisation Review*, 2017.
- 2 H. Abe, S. Yasui, A. Kuwata and H. Takeuchi, *Chem. Pharm. Bull.*, 2009, **57**, 647–652.
- 3 R. F. Shangraw, *Pharm. Dosage Forms: Tablets*, 1989, **1**, 195–246.
- 4 B. E. Schaller, K. M. Moroney, B. Castro-Dominguez, P. Cronin, J. Belen-Girona, P. Ruane, D. M. Croker and G. M. Walker, *Int. J. Pharm.*, 2019, **566**, 615–630.
- 5 A. N. Trementozzi, C.-Y. Leung, F. Osei-Yeboah, E. Irdam, Y. Lin, J. M. MacPhee, P. Boulas, S. B. Karki and P. N. Zawaneh, *Int. J. Pharm.*, 2017, **523**, 133–141.
- 6 H. P. Goh, P. W. S. Heng and C. V. Liew, *Int. J. Pharm.*, 2018, **547**, 133–141.
- 7 W. Yu, K. Muteki, L. Zhang and G. Kim, *J. Pharm. Sci.*, 2011, **100**, 284–293.
- 8 U. V. Shah, V. Karde, C. Ghoroi and J. Y. Heng, *Int. J. Pharm.*, 2017, **518**, 138–154.
- 9 A. Guo, J. Beddow and A. Vetter, *Powder Technol.*, 1985, **43**, 279–284.
- 10 A. Crouter and L. Briens, *AAPS PharmSciTech*, 2014, **15**, 65–74.
- 11 K. Kunnath, L. Chen, K. Zheng and R. N. Davé, *Powder Technol.*, 2021, **377**, 709–722.
- 12 C. Sun and D. J. Grant, *Int. J. Pharm.*, 2001, **215**, 221–228.
- 13 J. S. Kaerger, S. Edge and R. Price, *Eur. J. Pharm. Sci.*, 2004, **22**, 173–179.
- 14 L. J. Bellamy, A. Nordon and D. Littlejohn, *Int. J. Pharm.*, 2008, **361**, 87–91.
- 15 H. Masuda, K. Higashitani and H. Yoshida, *Powder Technology Handbook*, CRC Press, 2006.
- 16 A. J. Hlinak, K. Kuriyan, K. R. Morris, G. V. Reklaitis and P. K. Basu, *J. Pharm. Innovation*, 2006, **1**, 12–17.
- 17 B. Y. Shekunov, P. Chattopadhyay, H. H. Tong and A. H. Chow, *Pharm. Res.*, 2007, **24**, 203–227.
- 18 M. Leane, K. Pitt, G. Reynolds and M. C. S. W. Group, *Pharm. Dev. Technol.*, 2015, **20**, 12–21.
- 19 N. Sandler and D. Wilson, *J. Pharm. Sci.*, 2010, **99**, 958–968.
- 20 A. J. Megarry, S. M. E. Swainson, R. J. Roberts and G. K. Reynolds, *Int. J. Pharm.*, 2019, **555**, 337–345.
- 21 M. Capece, K. R. Silva, D. Sunkara, J. Strong and P. Gao, *Int. J. Pharm.*, 2016, **511**, 178–189.
- 22 V. R. Nalluri and M. Kuentz, *Eur. J. Pharm. Biopharm.*, 2010, **74**, 388–396.
- 23 H. Barjat, S. Checkley, T. Chitu, N. Dawson, A. Farshchi, A. Ferreira, J. Gamble, M. Leane, A. Mitchell, C. Morris, K. Pitt, R. Storey, F. Tahir and M. Tobyn, *J. Pharm. Innovation*, 2021, **16**, 181–196.
- 24 A. W. Jenike, *Bulletin No. 123*, Utah State University, 1964.
- 25 F. T. Ltd, *Shear Testing*, <https://www.freemantech.co.uk/powder-testing/ft4-powder-rheometer-powder-flow-tester/shear-testing><https://www.freemantech.co.uk/powder-testing/ft4-powder-rheometer-powder-flow-tester/shear-testing>, accessed 01/02/2021.
- 26 J. Zegzulka, D. Gelnar, L. Jezerska, R. Prokes and J. Rozbroj, *Sci. Rep.*, 2020, **10**, 1–19.
- 27 W. H. Organization, *The International Pharmacopoeia*, 2012, vol. 6.
- 28 J. Lever, *Nat. Methods*, 2016, **13**, 603–605.
- 29 P. Branco, L. Torgo and R. P. Ribeiro, *ACM Computing Surveys (CSUR)*, 2016, vol. 49, pp. 1–50.
- 30 M. Galar, A. Fernandez, E. Barrenechea, H. Bustince and F. Herrera, *IEEE Trans. Syst. Man Cybern. C Appl. Rev.*, 2011, **42**, 463–484.
- 31 D. C. Montgomery, E. A. Peck and G. G. Vining, *Introduction to Linear Regression Analysis*, John Wiley & Sons, 2021.
- 32 F. Sigrist, *arXiv*, 2018, preprint, arXiv:1808.03064, DOI: [10.48550/arXiv.1808.03064](https://doi.org/10.48550/arXiv.1808.03064).
- 33 Y. Zhang and A. Haghani, *Transport. Res. C Emerg. Technol.*, 2015, **58**, 308–324.
- 34 J. K. Jaiswal and R. Samikannu, *World Congress on Computing and Communication Technologies (WCCCT)*, 2017, pp. 65–68.
- 35 Q. Liu, X. Wang, X. Huang and X. Yin, *Tunn. Undergr. Space Technol.*, 2020, **106**, 103595.
- 36 S. M. Lundberg and S.-I. Lee, *Adv. Neural Inf. Process. Syst.*, 2017, **30**, 4768–4777.
- 37 S. Lundberg, *Welcome to the SHAP Documentation*, accessed 19/05/2022.
- 38 T. S. Pinto, O. Lima and L. Leal Filho, *Min., Metall., Explor.*, 2009, **26**, 105–108.
- 39 F. Fichtner, D. Mahlin, K. Welch, S. Gaisford and G. Alderborn, *Pharm. Res.*, 2008, **25**, 2750–2759.
- 40 C. G. Jange and R. K. Ambrose, *Powder Technol.*, 2019, **344**, 363–372.

