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Assessing the ecotoxicity of multicomponent nanomaterials using a classification SAR approach†

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The emerging applications of nanotechnology have led to the synthesis, production and use of a continuously increasing number of nanomaterials. In recent years, the focus is being shifted to multicomponent nanomaterials (MCNMs), due to the control over their functional properties. At the same time, the increasing exposure of ecosystems to such materials has raised concerns over their environmental hazard, with several *in vivo* and *in vitro* studies used to assess the ecotoxicity of MCNMs. The demanding nature of such methods has also led to the increasing development of *in silico* methods, such as structure–activity relationship (SAR) models. Although such approaches have been developed for single component nanomaterials, models for the ecotoxicity of MCNMs are still sparse in scientific literature. In this paper, we address the case of MCNM ecotoxicity by developing an *in silico* classification SAR computational framework. The models are built over a dataset of 652 ecotoxicity measurements for 214 metal and metal oxide MCNMs, towards bacteria, eukaryotes, fish, plants and crustaceans. This dataset is, to the best of the authors' knowledge, the largest dataset used for MCNM ecotoxicity. It is found that two descriptors can adequately classify different MCNMs based on their ecotoxicity over the whole heterogeneous dataset. These descriptors are the hydration enthalpy of the metal ion and the energy difference between the MCNM conduction band and the redox potential in biological media. Although the classification does not allow a quantitative ecotoxicity assessment, the heterogeneous nature of the dataset can reveal key MCNM features that induce toxic action, allowing a more holistic understanding of MCNM ecotoxicity, as well as the nature of interaction between the different MCNM components.

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Environmental significance

In recent years, nanotechnology research has focused on multi-component nanomaterials (MCNMs), which allow increased control over nanomaterial properties. However, concern has been raised regarding the safety of extended ecosystem exposure to such materials. The demanding nature of *in vitro* and *in vivo* methods has led to emerging *in silico* techniques, such as structure–activity relationship (SAR) models, for nanomaterial safety assessment. Nevertheless, such approaches for the case of MCNM ecotoxicity are limited, and are built using limited datasets. In this work, a classification approach is presented for MCNM ecotoxicity towards bacteria, eukaryotes, fish, plants, and crustaceans. The use of the heterogeneous datasets allows a more holistic understanding of MCNM toxicity, assisting the synthesis of MCNMs that are safe-by-design towards ecosystems.

1. Introduction

The wide spectrum of engineered nanomaterial (ENM) applications, ranging from catalysis¹ and energy² to electronics³ and nanomedicine⁴ has rendered the synthesis of ENMs of varying composition and forms^{5–10} a key part of nanotechnology research. This increasing use of ENMs has

led to the demand for control and tunability of ENM properties, depending on the targeted application.^{11,12} These demanding requirements have led to the synthesis of multi-component nanomaterials (MCNMs),^{13,14} such as bimetallic nanomaterials¹⁵ or alloys,¹⁶ surface functionalized nanomaterials,¹⁷ coated nanoparticles,¹⁸ as well as doped nanomaterials,¹⁹ among others.

The emergence of ENM and MCNM applications has also raised concern regarding the safety of extended exposure to such materials. While such exposure can be intentional, with the use of these materials as drug carriers,^{20,21} non-intentional exposure to ENMs and MCNMs can occur in varying environments.^{22–25} Ecosystems can also be exposed to

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ENMs and MCNMs, throughout their manufacturing process, application and disposal stages, *via* different exposure routes.^{26–28}

The safety of ENMs and MCNMs is usually assessed by measuring their toxicity. In this context, *in vitro* toxicity methods are used for a faster and cost effective initial toxicity assessment,²⁹ while *in vivo* methods, which are the most reliable, are used at later stages of regulatory risk assessment,³⁰ as they are time consuming and are characterized by higher cost and ethical concerns regarding animal testing. However, both types of toxicity assessment methods cannot keep up with the innovation, synthesis, and application of novel nanomaterials.³¹ For this reason, *in silico* methods have emerged to assess the safety of nanomaterials, with computational models developed for biodistribution^{32,33} and toxicity,^{31,34,35} toward different cells and organisms.

During the last decades, the increasing applications of nanotechnology have led to a wide number of research studies regarding nanomaterial toxicity assessment using *in vitro*^{36–38} and *in vivo*^{39–41} methods, as well as comparative reviews between the different toxicity assessment methods.^{42,43} In a similar way, interest has shifted towards *in silico* methods, and in particular structure–activity relationship (SAR) models^{44,45} based on the correlation of structural characteristics of the materials under study, known as descriptors, to biological activity endpoint data.⁴⁶ Both quantitative (QSAR) and qualitative (classification SAR)⁴⁷ models have been developed, especially for the case of metal oxide nanoparticles (NPs),^{47–49} but also for other types of ENMs.^{35,50} *In silico* models for such materials have been comprehensively reviewed by Buglak *et al.*,⁴⁴ and Li *et al.*⁵¹ However, most models are developed using a limited dataset,^{47,48,52,53} using a lower number of toxicity data. This limitation can severely hinder the extension of the use of SAR approaches, as models developed using a limited and homogeneous dataset apply only to a range of data similar to their training datasets. This means that the model precision may be severely decreased when comparing with measurements with a slight difference in the experimental design, or the NP properties, such as size and shape, thus damaging the model reproducibility. Furthermore, when larger and more heterogeneous datasets are used for the model development,^{49,54–57} the extraction of mechanistic information from SAR models challenging due to the complexity of the descriptors used. Complex descriptors can be difficult to be computed for novel NPs or more complex nanomaterial structures. Furthermore, the lack of mechanistic understanding does not allow to extend the use of QSAR models as decision supporting tools for the design and synthesis of safe-by-design nanomaterials.⁵⁸ Nevertheless, theoretical frameworks have been developed to provide a more cohesive, consistent and mechanistic understanding of metal oxide toxicity, while also leading to predictive SAR models for toxicity.^{31,53,59–63} Based on a similar framework, we recently used an extensive and heterogeneous dataset of toxicity measurements towards a

wide range of cell lines and organisms, to develop a classification SAR model.⁶⁴

Although *in silico* methods have been developed for pure metal oxides, the case of MCNMs is not yet sufficiently covered. In particular, the case of metal-loaded TiO₂ MCNMs has been studied by means of QSAR modelling,^{65–69} while only the case of ZnO-based NPs has been covered besides the TiO₂ MCNMs.⁷⁰ Nevertheless, the limited number of MCNMs and toxicity data used to develop these models does not allow a more global understanding of MCNM toxicity mechanisms. In a recent work, we presented a classification SAR model using an extensive dataset for the case of cytotoxicity and antibacterial activity of metal and metal oxide MCNMs,⁷¹ which allowed a more mechanistic insight on the dominant MCNM toxicity pathways. Regarding the case of ecotoxicity, however, although there has been an increasing interest using *in vitro* and *in vivo* assessment methods,^{72–75} *in silico* studies with the development of SAR models are still missing for MCNMs.

In this work, a classification SAR approach for the prediction of MCNM ecotoxicity is presented. The model is developed using an extensive dataset of 652 half-maximal concentration measurements for the ecotoxicity of metal and metal oxide MCNMs. The MCNMs considered in the present work consist of doped metal oxides, composite metal oxides, bimetallic NPs, as well as surface-loaded metal oxide NPs. Different subsets of data were used to build different models, based on the target organisms. In particular, models were developed for MCNM ecotoxicity towards *E. coli*, *S. aureus*, *D. rerio*, *D. magna* and *C. albicans*. Furthermore, the approach was extended to more heterogeneous datasets, consisting of MCNM ecotoxicity measurements towards different organisms' groups, such as bacteria, eukaryotes, fish, crustaceans and plants. Finally, the complete heterogeneous dataset was used for the development of a SAR model, showing that the approach can offer a more general insight regarding MCNM ecotoxicity.

The novelty of the approach lies in the size and nature of the dataset used for the development of such a model. The present study is, to the best of the author's knowledge, the largest dataset of MCNM ecotoxicity measurements used for the development of a SAR model. Furthermore, the dataset is heterogeneous, consisting of ecotoxicity measurements towards bacteria, eukaryotes, fish, plants and crustaceans. Such a heterogeneous SAR approach has not been applied before for the case of MCNM ecotoxicity while the used approach to compute MCNM descriptors for surface-loaded MCNMs is also novel for ecotoxicity models. Finally, the aim of the present work is not restricted to the development of a predictive classification model, but also to unravel the key characteristics of MCNMs that induce ecotoxicity. The size and heterogeneous nature of the dataset used for the model development will also assist towards a more holistic and mechanistic understanding of the ecotoxic action of MCNMs and the interaction between their components.



Table 1 Summary of the classification scheme for the experimental measurements in the dataset

| Reported concentration | Condition | Classification |
|-------------------------------------|--|----------------|
| EC50 value reported | $\log(\text{EC50}) \leq -2.5$ | Toxic |
| | $\log(\text{EC50}) > -2.5$ | Non-toxic |
| C_{max} yields effect >50% | $\log(C_{\text{max}}) \leq -2.5$ | Toxic |
| | $\log(C_{\text{max}}) > -2.5$ | Omitted |
| C_{max} yields effect <50% | $\log(C_{\text{max}}) > -2.5$ | Non-toxic |
| | $\log(C_{\text{max}}) \leq -2.5$, $C_{\text{max}} \geq 0.5 \cdot C_{\text{threshold}}$ | Non-toxic |
| | $\log(C_{\text{max}}) \leq -2.5$, $C_{\text{max}} < 0.5 \cdot C_{\text{threshold}}$ | Omitted |

2. Methods

2.1. Toxicity endpoints, classification and datasets

The dataset for the model development consists of MCNM ecotoxicity measurements, retrieved from 102 papers published in scientific literature. In particular, the ecotoxicity endpoints consist of reported half maximal effective (EC50), inhibitory (IC50), and lethal (LC50) concentration measurements. For clarity, the different half-maximal concentration measurements are onwards referred to as EC50, as in the work of Simeone and Costa.³¹ The different EC50 values were derived from dose–response studies towards different fish, plants, and crustaceans. Furthermore, as bacteria and eukaryotic organisms such as yeast, fungi and common pathogens can survive in ecosystems,^{76–78} they are also included in the ecotoxicity dataset. Although the common practice is to develop SAR models based on a specific endpoint and target cell line or organism, heterogeneous datasets using multiple endpoints and cells/organisms have been previously used for the development of SAR models for ENM cytotoxicity,^{55,71} ecotoxicity,⁵⁴ as well as combined cytotoxic and ecotoxic predictions.^{49,56,64}

Regarding the classification scheme, a measurement is characterized as toxic or non-toxic based on the criteria

reported by Simeone and Costa,³¹ following the scheme presented in our previous works.^{64,71} Briefly, the measurement is classified as toxic if the logarithm of the concentration endpoint in molar units (mol L^{-1}) is lower than -2.5 ($\log(\text{EC50}) \leq -2.5$). However, in some works, EC50 concentrations are reported as being higher than the range of experimentally tested concentrations ($\text{EC}_{50} > C_{\text{max, tested}}$). In such cases, if the maximum concentration yielded an effect less than 50% and $\log(C_{\text{max}}) > -2.5$, then the measurement was classified as non-toxic. On the other hand, if $\log(C_{\text{max}}) \leq -2.5$, then the following scheme is applied: If C_{max} is more than 50% of the threshold concentration (C_{thres}), the measurement is classified as non-toxic. If, however, C_{max} is less than 50% of the threshold value, the measurement is omitted from the dataset. The value of 50% was arbitrarily chosen, so that a significant amount of MCNM has been exposed so that the measurement is classified as non-toxic, and to reduce the number of data omitted from the dataset.

In a similar way, if the concentration tested was lower than the threshold value ($\log(C_{\text{max}}) \leq -2.5$), and the effect was higher than 50%, then the EC50 value is set to the concentration tested and classified as toxic. Otherwise, for measurements with an effect higher than 50% where the concentration tested was higher than the threshold value ($\log(C_{\text{max}}) > -2.5$), the measurement is removed. The classification scheme is summarized in Table 1.

Based on the above classification scheme, a final dataset of 652 MCNM ecotoxicity measurements is developed, presented in the ESI† of the paper. The dataset consists of ecotoxicity measurements of bimetallic NPs, metal-doped metal oxide NPs, surface loaded metal oxide NPs, and composite metal oxide NPs. The total dataset is divided to different subsets of data, corresponding to the individual organism that was exposed to the MCNMs. Datasets corresponding to the different organism groups are also created. The dataset size (minimum of 30 measurements) and the number of toxic and non-toxic measurements

Table 2 Different subsets of data used in the present work

| Cell type/cell type | No of measurements | No of NPs | % of toxic measurements | % of non-toxic measurements |
|----------------------|--------------------|-----------|-------------------------|-----------------------------|
| Individual organisms | | | | |
| <i>E. coli</i> | 92 | 65 | 50 | 50 |
| <i>S. aureus</i> | 70 | 45 | 50 | 50 |
| <i>D. rerio</i> | 88 | 36 | 21.6 | 78.4 |
| <i>D. magna</i> | 37 | 26 | 64.9 | 35.1 |
| <i>C. albicans</i> | 36 | 31 | 66.7 | 33.3 |
| Organism groups | | | | |
| Bacteria | 252 | 82 | 49.2 | 50.8 |
| Eukaryotes | 123 | 69 | 57.7 | 42.3 |
| Fish | 101 | 42 | 26.7 | 73.3 |
| Crustaceans | 74 | 58 | 64.9 | 35.1 |
| Plants | 102 | 40 | 30.4 | 69.6 |
| Complete dataset | 652 | 214 | 46.2 | 53.8 |



(minimum of 20% of both data classes) served as the criteria for the creation of a data subset. Based on the above scheme, 5 data subsets are created for individual organisms, as well as 5 datasets for organism groups. The complete dataset is also used for the model development. The final datasets are presented in Table 2.

2.2. Descriptors

2.2.1. Individual component descriptors. The MCNM descriptors are calculated using the descriptors of their individual components. The components consist of either metals or metal oxides. The individual components descriptors are calculated as described in our previous works,^{64,71} based on previous theoretical frameworks.^{60,61,63,79–81}

2.2.2. MCNM descriptors. The calculation of MCNM descriptors is realized using the descriptors of individual components. As the individual components of the MCNMs are either metal or metal oxides, it could be assumed that they have similar modes of action, as reported in previous works.⁵¹ Based on this assumption, an additive mixture approach is used for the calculation of MCNM descriptors, as suggested by Mikolajczyk *et al.*,⁶⁵ and used in earlier published works regarding nanomaterial mixtures,^{82–85} as well as multicomponent nanomaterials:^{66–68,71,86}

$$D_{\text{MCNM},j} = \sum_{i=1}^n x_i \cdot D_{i,j} \quad (1)$$

where $D_{\text{MCNM},j}$ is the j th descriptor in the MCNM descriptor matrix, x_i is the molar fraction of component i in the MCNM, n is the number of components, and $D_{i,j}$ is the j -th descriptor of the descriptor matrix of component i .

A different computational scheme is used for surface-loaded MCNMs, as presented in our previous work.⁷¹ Briefly, the composition of the MCNM surface, which is the area of interaction with the biological media, is computed. This is done by computing the total mass and molecular amount (q_i , in moles) for each of the particle components in a single particle, using the nominal densities (25 °C) of each component (from online handbooks/databases) and mass/molar fractions, as well as the particle volume:

$$V_{\text{part}} = \frac{4\pi R^3}{3} \quad (2)$$

where R is the particle radius.

The surface components are all assumed to be situated on the particle surface ($q_{s,i} = q_i$). The amount of the core component on the particle surface ($q_{s,\text{core}}$) is computed based on the core component's unit cell:

$$q_{s,\text{core}} = \frac{n_{\text{at}} \cdot \left(\frac{\text{SA}_{\text{part}}}{\text{SA}_{\text{unit cell}}} \right)}{N_{\text{av}}} \quad (3)$$

where n_{at} is the number of atoms or molecules in the unit cell, N_{av} is Avogadro's constant, $\text{SA}_{\text{part}} (= 4\pi R^2)$ is the surface

area of the particle, and $\text{SA}_{\text{unit cell}}$ is the surface area of the unit cell, based on the corresponding unit cell parameters. The surface composition is hence computed as a molar fraction of each component:

$$x_{s,i} = \frac{q_{s,i}}{\sum_i q_{s,i}} \quad (4)$$

Using the above approach, an estimation of the surface composition of surface-loaded MCNMs is obtained. This composition is then used in eqn (1) using the additive mixture approach to obtain the MCNM descriptors.

2.2.3. Descriptor selection. Once the descriptor matrix is created, several feature selection methods are employed in order to limit the number of descriptors for the SAR model development. This is done by removing redundant descriptors in order to increase the interpretability of the results by reducing the complexity of the model. First, intercorrelated descriptors for the individual components were identified using Pearson's correlation coefficient.⁸⁷ This is done to ensure that the descriptors used for the model development are not simple correlations of other descriptors, as such a case would damage the interpretability of the results. After the removal of the intercorrelated descriptors, the MCNM descriptor matrix is used to undergo further processing. Different feature selection methods, namely ReliefF⁸⁸ and chi-square⁸⁹ are used to rank the descriptors based on their relevance to the endpoints, with the 4 most relevant descriptors being kept for the SAR model development. Finally, the different combinations of the most relevant descriptors were tested, in order to identify the optimal models for the MCNM toxicity classification.

2.3. Model implementation, training and validation

As shown in Table 2, five datasets consist of MCNM ecotoxicity measurements towards individual organisms, while five different datasets were created for organism groups, namely bacteria, eukaryotes, fish, plants and crustaceans. Finally, the total set of data, consisting of all the MCNM ecotoxicity endpoints, was also used for SAR model development.

For datasets with a lower number of measurements ($n < 100$), a five-fold cross validation was used. For larger datasets ($n \geq 100$), a hold-out validation is used with 80% of the data as a training set and 20% as a validation set. In the hold-out validation scheme, the model is trained using a five-fold cross validation on the training set, while the validation is performed by comparing the trained model predictions to the toxic class assigned to the measurements in the validation set. The data splitting is random, and performed using MATLAB®. The model training is performed using the classification learner toolkit, by implementing Support Vector Machines (SVM), k-Nearest-Neighbors (kNN) and Random Forests (RF). The optimal models were identified with the use of different statistical metrics based on the resulting



different descriptors, while the threshold value to consider the descriptors intercorrelated was set to 0.95.⁹³ Only one of the intercorrelated descriptors is kept, while the other descriptors are discarded. From the remaining features, descriptors that were non-continuous over the dataset, as well as redundant descriptors with similar physical meaning, were subsequently removed from the descriptor matrix.

As a second step, the feature selection methods, namely ReliefF and chi-square, are used to rank the remaining descriptors based on their relevance to the response variable (ecotoxic class), as described in section 2.2.3. The four highest ranked descriptors are kept, following the feature selection analysis. Representative results for the four highest ranked descriptors, as derived from the two methods, using the complete dataset, are shown in Fig. 2. The descriptors rank using the two feature selection methods, for the different data subsets, is presented in Table 3. It is mentioned that in Table 3, only the descriptors that were ranked among the four most relevant descriptors for at least one dataset are shown.

The descriptor ranking presented in Fig. 2 and Table 3 shows that seven descriptors are ranked within the four highest ranked descriptors at least once, for all the data subsets, using the two feature selection methods. Interestingly, HE (hydration enthalpy^{61,63,64,71}), Dbio (energy difference between the conduction band of the MCNM and redox potential of biological media^{64,71}), and IP (ionic potential of the metal component⁸¹) are ranked amongst the four highest ranked descriptors for all 11 datasets, using at

As mentioned in section 2.2.3, the intercorrelation analysis of the individual component descriptors is the first step of the descriptor selection process. The results of this analysis for different metal and metal oxide descriptors are presented in Fig. 1, below. The Pearson coefficient has been selected to analyze the intercorrelation between the



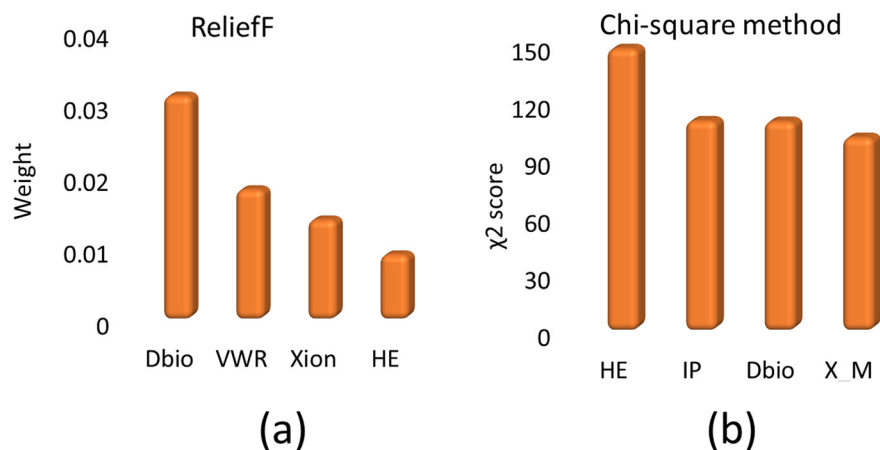


Fig. 2 The four most relevant descriptors, for the complete dataset, as derived from a) ReliefF and b) chi-square methods.

Table 3 Descriptor ranking for the different datasets, using the ReliefF and chi-square feature selection methods

| | Relief F | | | | | | | Chi-square | | | | | | |
|--------------------|----------|------|------|----|-----|-----|------|------------|------|------|----|-----|-----|------|
| | HE | Dbio | S_IP | IP | X_M | VWR | Xion | HE | Dbio | S_IP | IP | X_M | VWR | Xion |
| <i>E. coli</i> | 3 | — | 2 | — | — | 1 | 4 | 2 | 1 | — | 3 | — | 4 | — |
| <i>S. aureus</i> | 3 | — | 1 | 2 | 4 | — | — | 4 | 1 | — | — | 3 | 2 | — |
| <i>D. rerio</i> | — | 2 | — | 4 | — | 1 | 3 | 1 | — | 3 | — | 2 | 4 | — |
| <i>D. magna</i> | 2 | 4 | — | — | — | 1 | 3 | 1 | — | 2 | 3 | — | 4 | — |
| <i>C. albicans</i> | 3 | 1 | 2 | — | — | — | 4 | 2 | 1 | 3 | 4 | — | — | — |
| Bacteria | 2 | 4 | 1 | — | — | — | 3 | 2 | — | — | 4 | 1 | 3 | — |
| Eukaryotes | 3 | 1 | 4 | 2 | — | — | — | 2 | 1 | — | — | 3 | — | 4 |
| Fish | — | 1 | — | 3 | 2 | 4 | — | 2 | 1 | — | — | — | 4 | 3 |
| Crustaceans | 1 | 3 | — | 4 | 2 | — | — | 1 | 2 | 3 | 4 | — | — | — |
| Plants | 4 | 2 | — | — | 3 | 1 | — | 2 | — | 3 | 1 | — | — | 4 |
| Complete dataset | 4 | 1 | — | — | — | 2 | 3 | 1 | 3 | — | 2 | 4 | — | — |

least one feature selection method. Based on the above observations, the three abovementioned descriptors are kept for the model development, with different descriptor combinations being tested. The results are presented in the following sections, for the models developed for the different datasets.

3.2. MCNM ecotoxicity models for individual organisms

Following the descriptor selection step, the analysis continues with the development of SAR models for the datasets consisting of MCNM ecotoxicity measurements towards individual organisms. Different combinations of the three descriptors identified in the descriptor selection step

(HE, Dbio, IP) are tested, and the optimal model for each dataset is kept. The inclusion of a descriptor is accepted when the model accuracy is significantly improved. The accuracy of the models built using the different descriptor combinations, for the individual organism datasets, is presented in Table 4.

As seen from the results of Table 4, amongst the single descriptor models, HE is the most predictive descriptor, for most datasets, with the exception of the model developed for the measurements towards *E. coli* where IP is the most predictive descriptor. The inclusion of a second descriptor improved the model accuracy, with the combination of HE and Dbio being the most predictive set of descriptors, except from the case of the *D. magna* dataset, where the accuracy

Table 4 Accuracy percentage of models developed for the individual organism datasets, using the different descriptor combinations

| Dataset | Descriptor combination | | | | | | |
|--------------------|------------------------|----------|--------|----------|------|------|------|
| | HE, Dbio, IP | HE, Dbio | HE, IP | Dbio, IP | HE | Dbio | IP |
| <i>E. coli</i> | 95.2 | 94.6 | 85.9 | 90.2 | 70.7 | 53.3 | 78.3 |
| <i>S. aureus</i> | 97.1 | 97.1 | 81.4 | 95.7 | 84.3 | 51.4 | 77.1 |
| <i>D. rerio</i> | 89.8 | 93.2 | 87.5 | 81.8 | 83.0 | 71.6 | 78.4 |
| <i>D. magna</i> | 86.5 | 89.2 | 89.2 | 86.5 | 89.2 | 78.4 | 83.8 |
| <i>C. albicans</i> | 91.7 | 94.4 | 88.9 | 88.9 | 86.1 | 83.3 | 80.6 |



Table 5 Statistical metrics of the classification models built for the datasets of individual organisms

| Organism | Descriptors | Validation scheme | Acc (%) | Prec (%) | Sens (%) | Sel (%) |
|--------------------|-------------|-------------------------|---------|----------|----------|---------|
| <i>E. coli</i> | HE, Dbio | 5-Fold cross validation | 94.6 | 97.7 | 91.3 | 97.8 |
| <i>S. aureus</i> | HE, Dbio | 5-Fold cross validation | 97.1 | 94.6 | 100 | 94.3 |
| <i>D. rerio</i> | HE, Dbio | 5-Fold cross validation | 93.2 | 100 | 68.4 | 100 |
| <i>D. magna</i> | HE | 5-Fold cross validation | 89.2 | 100 | 83.3 | 100 |
| <i>C. albicans</i> | HE, Dbio | 5-Fold cross validation | 94.4 | 95.8 | 95.8 | 91.7 |

did not improve with the addition of Dbio as a descriptor. Finally, the inclusion of all three descriptors produced a non-significant or lower accuracy than the combination of HE and Dbio. Hence, in order to reduce the complexity of the model and enhance the interpretability of the results, the minimum number of descriptors that produce the highest accuracy are kept. This means that the two-descriptor combination of HE and Dbio is kept for the different models, except from the case of the *D. magna* model, where the single descriptor of HE is kept.

The statistical metrics for the optimal models developed using the ecotoxicity measurements within the individual organism datasets are presented in Table 5.

The statistical metrics presented in Table 5 show that the models developed have acceptable values for accuracy, towards all the individual model datasets. The lowest accuracy is obtained by the *D. magna* model, which, however, has an acceptable value of 89.2%. The developed models also show high values for precision, sensitivity and selectivity, with the exception of the *D. rerio* model, which shows a

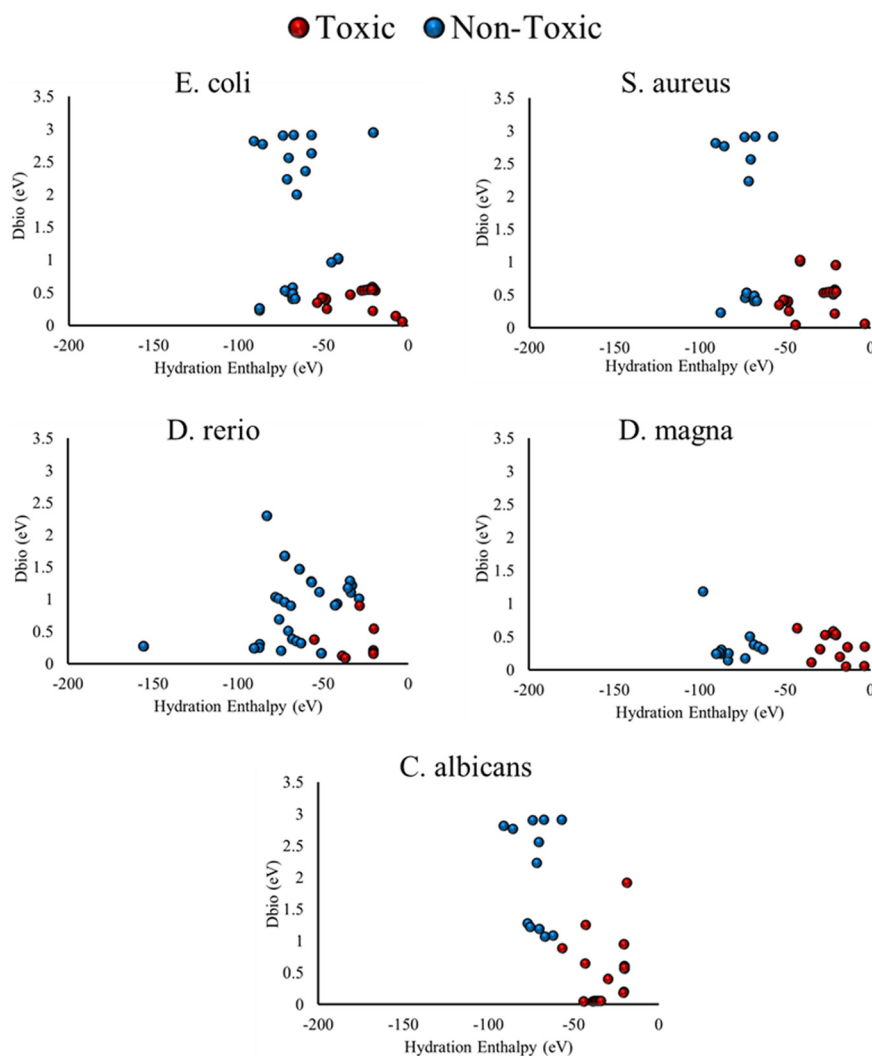


Fig. 3 SAR model predictions of the ecotoxic class of the different MCNMs, towards the different datasets, as a function of the two descriptors (HE, Dbio).



sensitivity of 68.4%. This lower sensitivity could be assigned to the more imbalanced nature of the dataset, as only 21.6% of the measurements are classified as toxic within the *D. rerio* ecotoxicity dataset. The very high values of precision and selectivity could also be assigned to this imbalance, for the *D. rerio* dataset. In any case, the different models were able to classify the ecotoxicity measurements in the different datasets successfully, using the same descriptor combination of HE and Dbio, except from the *D. magna* model which used the HE descriptor. However, as seen in the results of Table 4, the addition of Dbio did not decrease or increase the accuracy.

The results presented in Tables 4 and 5 show that the classification SAR approach presented can predict the MCNM ecotoxicity towards the different organisms with acceptable accuracy, using similar descriptors. This could hint towards similar underlying mechanisms being responsible for the ecotoxicity of the different MCNMs towards the different organisms. Furthermore, the same results show that the additive mixture approach, as suggested by Mikolajczyk *et al.*,⁶⁵ and the approach used for surface-loaded MCNMs previously presented,⁷¹ is able to produce predictive descriptors for MCNM ecotoxicity. This can assist towards the understanding of the interaction between the various MCNM components.

The small number of descriptors used for the classification SAR model development, together with the fact that similar descriptors are used for all the individual organism datasets, allows the mapping of the classification results over the descriptor space. The predicted ecotoxic class for the different MCNMs, towards the different organisms, is presented in Fig. 3, as a function of the two descriptors, HE and Dbio. It is noted that the results for the *D. magna* model are also presented over the same descriptor space for comparison, even though the model is developed using only HE as a descriptor.

Results of Fig. 3 show that similar results are obtained for the ecotoxic class prediction over the descriptor space, for all the different datasets. Specifically, the ecotoxic MCNMs are characterized by a less negative HE and a lower Dbio value, for all the different classification SAR models developed. Specifically, a threshold value of HE close to -50 eV and a Dbio value close to 1 eV separate the two classes, with the exception of the *C. albicans* model, where ecotoxic

classification is obtained for higher Dbio values. This similar behavior may hint towards similar dominating mechanisms for ecotoxic action among the MCNMs of the different datasets, towards different organisms. Such results are consistent with previous findings for the case of pure metal oxides,^{61–64} and may assist towards a more global understanding of ecotoxic action of MCNMs. Further discussion regarding the ecotoxic mechanisms and the interpretation of model results will be presented in a subsequent section of the present paper.

3.3. MCNM ecotoxicity models for heterogeneous datasets

Based on the results of the previous section, where similar descriptors were able to predict the ecotoxic class of the different MCNMs in the individual organism datasets, the approach is extended for more heterogeneous datasets in terms of the target organism. For this reason, the datasets consisting of ecotoxicity measurements towards similar organism groups are used for the SAR model development. The complete dataset is also used to develop a SAR model, in order to test whether more holistic ecotoxicity predictions can be obtained with the present approach.

As in the previous section, the different combinations of the descriptors identified in the selection step (HE, Dbio, IP) are used for SAR model development, in order to identify the optimal descriptor combination. The accuracy of the models built using the different descriptor combinations, for the organism groups and the complete dataset, is presented in Table 6. It is noted that for the descriptor combination tests, all the models were developed using five-fold cross validation.

The results of Table 6 show that among the single descriptor models, HE is the most predictive descriptor for the fish and crustaceans models, as well as for the complete dataset model. IP is the most predictive descriptor for the bacteria and plants models, while Dbio is the most predictive for the eukaryotes model. As for the case of individual organism models, the addition of a second descriptor improved the accuracy in all the heterogeneous models, except from the crustacean model, where the addition of a second descriptor to HE did not affect the model accuracy. Interestingly, the most predictive descriptor pair was found

Table 6 Accuracy percentage of models developed for the heterogeneous organism group datasets and the complete dataset, using the different descriptor combinations

| Dataset | Descriptor combination | | | | | | |
|------------------|------------------------|----------|--------|----------|------|------|------|
| | HE, Dbio, IP | HE, Dbio | HE, IP | Dbio, IP | HE | Dbio | IP |
| Bacteria | 92.9 | 92.9 | 85.3 | 90.9 | 75.4 | 75.0 | 84.9 |
| Eukaryotes | 93.5 | 92.7 | 81.3 | 87.0 | 71.5 | 85.4 | 72.4 |
| Fish | 87.1 | 88.1 | 78.2 | 84.2 | 76.2 | 74.3 | 72.3 |
| Crustaceans | 93.2 | 93.2 | 93.2 | 82.4 | 93.2 | 78.4 | 74.3 |
| Plants | 87.3 | 87.3 | 85.3 | 83.3 | 80.4 | 69.6 | 85.3 |
| Complete dataset | 89.9 | 89.6 | 75.6 | 80.8 | 76.4 | 64.0 | 72.9 |



Table 7 Statistical metrics of the optimal classification models built for the datasets of organism groups and the complete dataset

| Organism | Descriptors | Validation scheme | Acc (%) | Prec (%) | Sens (%) | Sel (%) |
|------------------|-------------|--|---------|----------|----------|---------|
| Bacteria | HE, Dbio | Train. (<i>n</i> = 202) Val. (<i>n</i> = 50) | 94.0 | 90.5 | 95.0 | 93.3 |
| Eukaryotes | HE, Dbio | Train. (<i>n</i> = 99) Val. (<i>n</i> = 24) | 91.7 | 92.9 | 92.9 | 90.0 |
| Fish | HE, Dbio | Train. (<i>n</i> = 81) Val. (<i>n</i> = 20) | 85.0 | 80.0 | 66.7 | 92.3 |
| Crustaceans | HE | 5-Fold cross validation | 93.2 | 97.8 | 91.7 | 96.2 |
| Plants | HE, Dbio | Train. (<i>n</i> = 82) Val. (<i>n</i> = 20) | 90.0 | 100 | 75.0 | 100 |
| Complete dataset | HE, Dbio | Train. (<i>n</i> = 522) Val. (<i>n</i> = 130) | 89.2 | 83.3 | 92.6 | 86.9 |

to be the HE and Dbio combination, as in the individual organism models. Finally, the addition of IP to the HE and Dbio combination led to non or insignificant increase of the model accuracy. From the above results, it is concluded that

the optimal descriptor set is the combination of HE and Dbio, with the exception of the crustacean model, where the single descriptor model developed using HE was found to be optimal.

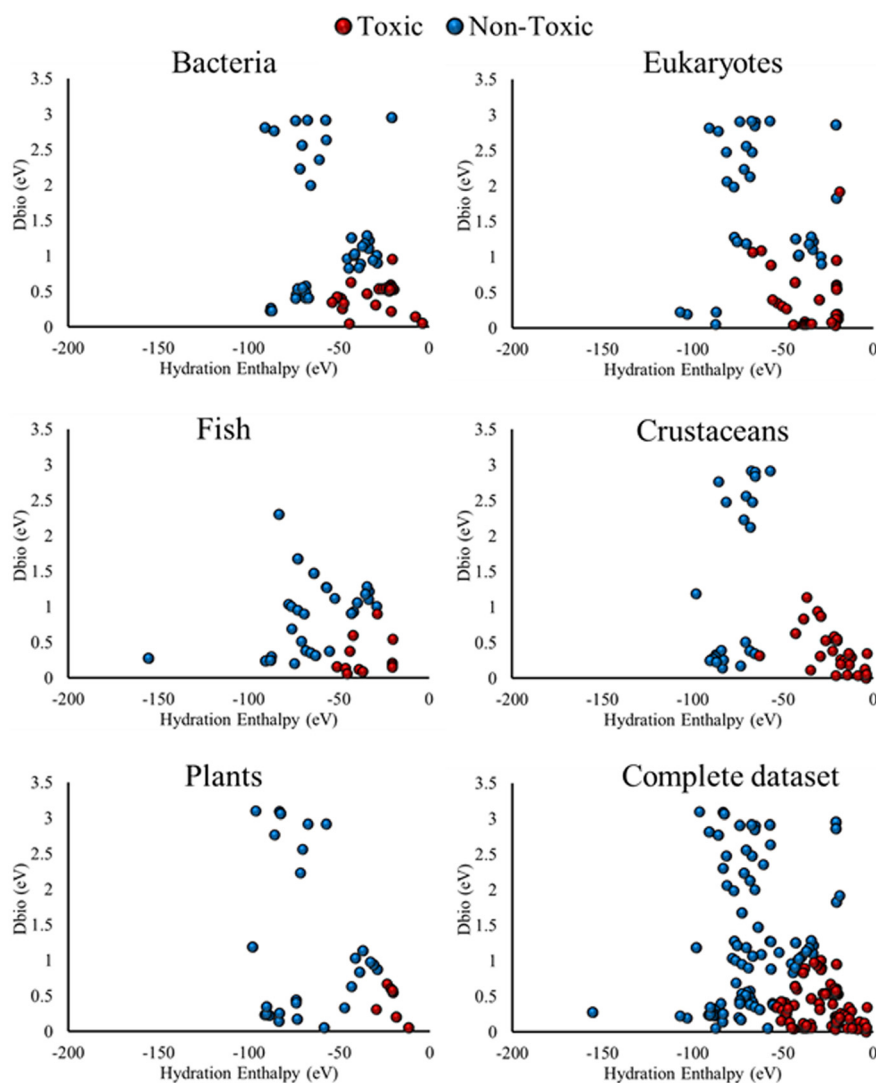


Fig. 4 SAR model predictions of the ecotoxic class of the different MCNMs, towards the different organism groups and complete datasets, as a function of the two descriptors (HE, Dbio).



Using the abovementioned descriptor combinations, the optimal models are developed. The resulting statistical metrics for the optimal models using the ecotoxicity measurements within the heterogeneous datasets of organism groups and the complete dataset are presented in Table 7. It is mentioned that the models presented in Table 7 display different accuracy than in Table 6 (except from the crustaceans model), due to the fact that they are developed using a hold-out validation scheme, contrary to the models of Table 6, where a five-fold cross validation scheme was used for all models.

Results of Table 7 show that acceptable accuracy is obtained for all developed models using the heterogeneous datasets for the organism groups, as well as the complete dataset. Furthermore, acceptable values are obtained for all statistical metrics along the different models, except from the sensitivity obtained by the models developed using the fish and plants datasets. This lower model sensitivity could be assigned to the more imbalanced nature of these datasets (26.7% and 30.4% of toxic measurements, respectively). Nevertheless, the fact that the developed models can predict the ecotoxic class of the various MCNMs in the different validation sets of the different organism group and complete datasets, using similar descriptors, enhances the notion that similar mechanisms dominate the MCNM ecotoxic action. As in the case of the individual organism models, HE and Dbio are found to be the most predictive set of descriptors, except from the case of the crustacean model, which was developed using only the HE descriptor. Moreover, the additive mixture approach used to compute the MCNM descriptors was again able to produce predictive descriptors for the different classification SAR models, allowing a more global understanding of the nature of the interaction between MCNM component.

The ecotoxic class predicted by the different classification SAR models, mapped over the space defined by the two model descriptors, is presented in Fig. 4. This mapping allows more clear understanding of the effect of the different descriptors on the model results, and is possible due to the low number of descriptors used during the model development.

As in the case of the models developed for the individual organism datasets (Fig. 3), the results of Fig. 4 present a similar ecotoxic class prediction over the descriptor space, for all models developed for the organism group and complete datasets. Ecotoxic class is predicted for MCNMs that exhibit a less negative HE and a lower Dbio value. For all the different classification SAR models developed. A similar trend is observed across all the SAR models, both for the case individual organisms (Fig. 3) and organism groups (Fig. 4). This behavior enhances the notion of similar mechanisms being dominant for the ecotoxic action of MCNMs towards different organisms. Such results may lead to a more holistic understanding of ecotoxic action of MCNMs, which in turn can assist in the development of MCNMs with properties that increase or decrease the interaction of MCNMs with the

abovementioned organism, according to the desired application.

3.4. Applicability domain

Based on the guidelines for (Q)SAR model development, as set by the OECD, a validated SAR model should have a defined applicability domain, which is one of the five principles of SAR validation.⁹⁰ In this way, the descriptor space within which SAR model can be applied is clearly defined. For this reason, the applicability domain identification methods usually consist of geometric, range-based, and distance-based methods.⁹² In the present work, the bounding box PCA, convex hull and centroid distance methods are employed to identify the applicability domain of the developed classification SAR models, for the different datasets. The number of measurements that fell outside the applicability domain of the model, using the different methods, is presented in Table 8. It is noted that the applicability domain is defined using the training set of each dataset.

The results of Table 8 show that the Bounding box PCA method included all the MCNM measurements within the applicability domain of all the developed classification SAR models. The convex hull method also includes the total set of MCNM measurements within the applicability domain of the models developed for most datasets, with the exception of the fish dataset, where two measurements were deemed to be outside the applicability domain. A single measurement was also deemed to be outside the applicability domain for the complete dataset model, by the convex hull method. On the other hand, the centroid distance method defined narrower applicability domains, which did not include a number of MCNM measurements, for all the different ecotoxicity datasets. Such results show the dependency of the applicability domain on the method used for its definition. This dependency has been identified and discussed in scientific literature.⁹² The analysis of the applicability domain results reproduces the findings of previous SAR model development, where metal oxides with similar

Table 8 Number of MCNMs toxicity measurements that are deemed to be outside the applicability domain

| Dataset | Bounding box PCA | Convex hull | Centroid distance |
|--------------------|---------------------|----------------|----------------------|
| <i>E. coli</i> | — | — | 2 |
| <i>S. aureus</i> | — | — | 3 |
| <i>D. rerio</i> | — | — | 2 |
| <i>D. magna</i> | — | — | 2 |
| <i>C. albicans</i> | — | — | 2 |
| Bacteria | — | — | 12 |
| Eukaryotes | — | — | 6 |
| Fish | — | 2 | 2 |
| Crustaceans | — | — | 4 |
| Plants | — | — | 5 |
| Complete dataset | — | 1 | 27 |



descriptor values are outside the applicability domain of the models.^{62,64}

3.5. Overall predictions and mechanistic information

The results presented in the previous sections showed that the classification SAR models developed for the different datasets consisting of MCNM ecotoxicity measurements towards individual organisms, as well as for heterogeneous datasets for organism groups, were developed using similar sets of descriptors as the optimal combination (HE and Dbio descriptors). Moreover, the optimal model developed for the complete dataset also used the same combination of two descriptors. This similar descriptor combination allows the mechanistic interpretation of results over the whole set of MCNM ecotoxicity measurements, assisted by an overall ecotoxicity prediction by the SAR model.

The complete dataset model is used to predict the ecotoxic class of the different MCNMs, and the results are presented in Fig. 5a, over the space defined by the two descriptors used for the model development. In order to compare the model

predictions to the actual experimental measurements, Fig. 5b is used to present the corresponding percentage of toxic measurements for each MCNM (as several MCNMs have multiple measurements within the dataset).

The results of Fig. 5a show that the ecotoxic MCNMs are characterized by a lower Dbio value, as well as a less negative HE value. Hence, using the physical interpretation of the two descriptors, MCNMs that have a conduction band energy close to the redox potential of biological pairs (lower Dbio value), together with a less exothermic hydration of their respective metal cations (less negative hydration enthalpy, HE), are more probable to be ecotoxic. Similar conclusions have been drawn from previous experimental and modelling works for metal oxide NPs.^{61–63} In the same direction with those works, our previous works for the case of pure metal oxides toxicity,⁶⁴ as well as metal oxide MCNMs cytotoxicity,⁷¹ have also identified these descriptors as predictive. The measurements presented in Fig. 5b show that the percentage of ecotoxic experimental measurements for the different MCNMs follows the same trend over the descriptor space. Besides the misclassification of some MCNMs, the experimental classification follows the model predictions, as the vast majority of ecotoxic MCNMs are situated in the space bounded by low Dbio and less negative HE values. Similarly, MCNMs that are characterized by a high Dbio or a highly negative HE value exhibit less ecotoxic measurements. The good agreement between the model classification and the measurements in the dataset does not only show the accuracy and predictive ability of the classification model, but also enhances the notion that similar MCNM characteristics may induce ecotoxic action towards the different organisms taken into account in the dataset. These key characteristics can be adequately quantified by the two descriptors taken into account for the classification SAR model development.

It should be noted however, that the exact boundary values of descriptors that define whether a MCNM is ecotoxic is not possible using the present approach, as it does not aim towards a quantitative ecotoxicity prediction, but rather towards a qualitative assessment. Hence, the model results are sensitive towards the classification scheme used. Furthermore, the models developed for heterogeneous datasets (Table 7 and Fig. 4) in terms of the tested organisms do not take into account the varying cell morphology of the different target organisms. Although the cell morphology has an influence on the ecotoxic action of the different MCNMs, all the different models were developed using similar descriptors, with a good accuracy towards the ecotoxicity measurements, which may hint that the MCNMs have similar ecotoxic modes of action. However, these results do not mean that the toxicity mechanisms are unaffected by the morphology of the target cells, but rather that the dominating toxicity mechanisms are of similar nature. Specifically, the cell morphology could severely affect the ecotoxic action in a more quantitative way, with higher or lower uptake and MCNM-cell interaction rates. Due to the

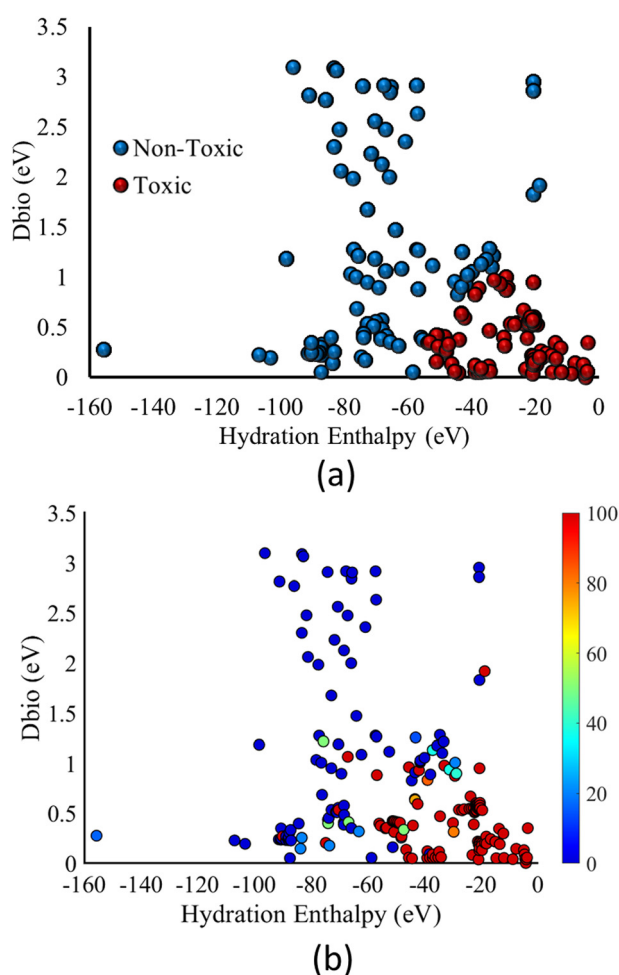


Fig. 5 a) Ecotoxic class prediction for the different MCNMs by the complete dataset model, b) ecotoxic measurements percentage for the different MCNMs in the complete dataset.



qualitative nature of the model presented in this work, the extraction of quantitative information is not possible by the present approach. However, similar mechanisms for the toxic action of metal oxides, towards different morphologies of target cells or organisms have been concluded to occur in previous nanotoxicology studies,^{61,63} where the conduction band of nanoparticles has been found to induce electron transfer and toxicity towards BEAS-2B and RAW264.7 cells,⁶³ as well as *E. coli*.⁶¹ On the basis of such observations, classification SAR approaches for nanomaterial toxicity using heterogeneous morphology datasets have been developed,^{49,54–57,64,71} assessing the toxicity of nanomaterials over a wide range of cell morphologies. In a similar way, the general trends seen in Fig. 5 are reproduced over a large and populous dataset of heterogeneous MCNM measurements, which encourages a more general understanding of MCNM ecotoxicity mechanisms.

As previously mentioned, Dbio expresses the energy difference between the conduction band and the mean redox potential of pairs in biological media.^{60,61,63,94,95} Hence, a low Dbio value means that electron transfer between the MCNM and the cell is more probable. This electron transfer, previously identified as a toxicity mechanism for the case of metal oxides, can increase the oxidative stress on the cells, by unbalancing its reducing capacity.^{59,60} Furthermore, overlapping metal oxide conduction bands and biological redox potentials have been correlated with the production of reactive oxygen species (ROS).⁹⁶ Such ROS can include hydrogen peroxides and OH radicals.^{63,97–103} Similar quantities have been used for the development of QSAR,⁵³ classification SAR^{61–64,71} models, as well as for nanomaterial toxicity grouping,³¹ towards cells and organisms of different kind, showing a more holistic metal oxide toxicity pathway.

The hydration enthalpy (HE) denotes the energy released during the hydration of a respective metal ion, that is released from the MCNM. In this way, the descriptor expresses the affinity of these metal ions to water molecules.⁶¹ A more negative HE value means that water molecules will be more strongly attracted to the metal ions, leading an increase of the ion's hydration shell. In turn, a larger hydration cell hinders the ion's permeability through the cell membrane.^{104–108} As in previous works, HE is computed using Latimer's equation:⁶¹

$$HE = -\frac{637.184 \cdot z^2}{(r + 50)} \quad (5)$$

where z is the oxidation number, or metal cation charge and r its ionic radius. The term of z^2/r , used in eqn (5), has shown predictive capabilities as a descriptor for the toxicity classification of metal oxides.⁶² Furthermore, this term can be seen as the inner product of oxidation number or cation charge (z) and ionic potential (z/r). The ionic potential has previously been correlated with metal oxide toxicity,¹⁰⁹ with this correlation explained by the damaging charge potential of metal ions.^{31,51} On the other hand, the cation charge is a predictor for the metal/metal oxide metal ion release and

solubility.^{31,51,110} Using the above physical interpretations, it is concluded that the descriptor HE is a complex descriptor, that expresses the contributions of the release of metal ions, their charge, and their interaction with aqueous media.

The release of metal ions has been concluded to be a dominant initial pathway of metal and metal oxide nanoparticle toxicity.⁹⁷ Several underlying mechanisms regarding the toxic interaction between these ions and different cells have been identified, such as enzyme inactivation,^{111,112} cell membrane damage^{63,113} and the increase of oxidative stress.^{111,112,114–116} DNA damage has also been attributed to the interaction with certain metal ions.^{111,117,118}

The predictive capability of the two descriptors towards the various and heterogeneous datasets could hint that the abovementioned mechanisms could be the dominant pathways towards the metal and metal oxide MCNM ecotoxicity. However, it cannot be concluded whether one of the two pathways prevails over the other. As seen from the results of sections 3.2 and 3.3, single descriptor models were less accurate than models developed using the combination of the two descriptors, except from the *D. magna* and crustaceans models. For the rest of the models, the ecotoxic MCNMs exhibit a lower Dbio value and a less negative HE. This could mean that both electron exchange between the MCNM and the organism cells, as well as metal ion release that permeate the cell must take place for the MCNM to induce ecotoxic action, as defined with the present classification scheme.

Regarding the nature of interaction between the different components of the MCNMs in the different datasets, the additive mixture approach introduced by Mikolajczyk *et al.* for the case of MCNMs,⁶⁵ was able to calculate descriptors that were predictive towards the whole dataset range. The additive approach assumes that the different MCNM components have similar modes of action. While other approaches have been used for the descriptor calculation for mixture of chemicals,^{119,120} QSAR model development for nanomaterial mixtures has mainly employed the additive mixture approach.^{82,83,121} Works that have developed QSARs for smaller MCNMs datasets have also used similar approaches for the descriptors, showing high predictivity towards toxicity endpoints.^{65–68,70} In our previous work, we used the additive mixture approach, together with a novel approach to calculate descriptors for surface loaded MCNMs, to calculate predictive descriptors for the cytotoxicity classification over a large MCNM dataset, leading to the development of high accuracy models.⁷¹

With the results of the present work, it is seen that the exact same approach can be used to calculate descriptors that are also able to classify MCNMs based on their ecotoxicity, towards a wide range of organisms, such as bacteria, eukaryotes, fish, plants, and crustaceans. As the additive mixture approach assumes, this could hint towards similar ecotoxic pathways of the different MCNM components. The components in the present work are either metals or metal



oxides, which have previously been reported to have similar modes of action.^{51,64} However, the qualitative nature of the classification approach presented in this work does not allow the quantitative interpretation of the individual component impact in certainty. Nevertheless, quantitative methods have deemed that the compositional ratio of MCNM components dictates their respective impact, showing an additive effect to toxicity.^{66,67} The large and heterogeneous nature of the MCNM ecotoxicity dataset used in the present work, along with the high accuracy of the developed models, enhances the notion that this additive effect also occurs for MCNM ecotoxicity. In any case, this should be tested by developing quantitative models for more homogeneous datasets, under similar experimental conditions, which will be a subject of future work.

To summarize, the results presented in this study can show the potential of data-based models, such as classification SAR approaches, to be used not only as predictive models, but also as tools for a more general understanding of toxicity mechanisms and modes of action. With their use as an inductive tool to extract scientific information from large and heterogeneous datasets of toxicity measurements, such models can potentially be used to assist the synthesis of safe-by-design nanomaterials. A potential example of such an approach is presented in the work of Feng *et al.*,⁵⁸ where based on the results of previous research regarding the dependence of metal oxide toxicity on the conduction band energy, the authors managed to synthesize MCNMs with control over their biological activity, by adequately tuning the conduction band energy of the produced nanomaterials, which is consistent with the findings of the present work.

Conclusions

In the present paper, a dataset of 652 measurements is used to develop classification SAR models for MCNM ecotoxicity. This dataset is, to the best of the author's knowledge, the largest dataset of MCNM ecotoxicity measurements used for the development of such a model. Furthermore, the dataset is heterogeneous, consisting of ecotoxicity measurements towards bacteria, eukaryotes, fish, plants and crustaceans. The models developed for the ecotoxicity towards a specific organism classified the MCNMs in the dataset with accuracy exceeding 89%, based on their ecotoxicity towards the different organisms and organism groups. Models were also able to classify the ecotoxic MCNMs towards different organism groups with an accuracy exceeding 90% towards bacteria, eukaryotes, plants and crustaceans, while the accuracy for fish was 85%. Finally, the complete heterogeneous dataset was used to build a classification model for the MCNM ecotoxicity, with an accuracy of 89.2%. The same descriptor set was used for the classification of the different subsets of data, allowing a more holistic understanding of the underlying mechanisms that dictate MCNM ecotoxicity.

In particular, the electron transfer between the MCNM and the biological pairs, as well as the release and transport of metal ions from the MCNMs, were deemed to be the dictating ecotoxic pathways for the different MCNMs. These findings are consistent with previous works for the cytotoxicity of metal oxide nanoparticles and MCNMs. However, the identification of these descriptors for the case of ecotoxicity is novel for the case of MCNM. The two descriptors that expressed the abovementioned mechanisms were computed based on the additive mixture approach, shedding light on the nature of the interaction between the MCNM components. These findings are consistent with previous works regarding the QSAR modelling of MCNM toxicity, where the additive mixture approach was found to produce predictive descriptors.

The novelty of the present study lies in the development of a SAR approach using a large, heterogeneous dataset for the prediction of MCNM ecotoxicity. Such an approach allows a more holistic understanding of the ecotoxic action, upon ecosystem exposure of different MCNMs and their constituting components towards various organisms. The mechanistic information extracted by the present approach regarding MCNM ecotoxicity and the interaction of the multiple components of MCNMs, can thus assist towards a more knowledge-driven MCNM ecotoxicity assessment, as well as the synthesis of safe-by-design MCNMs for various applications.

Data availability

The data supporting this article have been included as part of the ESI.†

Conflicts of interest

There are no conflicts to declare.

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References

- 1 S. Mitchell, R. Qin, N. Zheng and J. Pérez-Ramírez, Nanoscale engineering of catalytic materials for sustainable technologies, *Nat. Nanotechnol.*, 2021, **16**, 129–139.
- 2 F. Christian, E. Selly, D. Adityawarman and A. Indarto, Application of nanotechnologies in the energy sector: A brief and short review, *Front. Energy*, 2013, **7**, 6–18.
- 3 C.-Z. Ning, L. Dou and P. Yang, Bandgap engineering in semiconductor alloy nanomaterials with widely tunable compositions, *Nat. Rev. Mater.*, 2017, **2**, 17070.



- 4 X. Hu, Y. Zhang, T. Ding, J. Liu and H. Zhao, Multifunctional Gold Nanoparticles: A Novel Nanomaterial for Various Medical Applications and Biological Activities, *Front. Bioeng. Biotechnol.*, 2020, **8**, 990.
- 5 G. P. Gakis, H. Vergnes, F. Cristiano, Y. Tison, C. Vahlas, B. Caussat, A. G. Boudouvis and E. Scheid, In situ N₂-NH₃ plasma pre-treatment of silicon substrate enhances the initial growth and restricts the substrate oxidation during alumina ALD, *J. Appl. Phys.*, 2019, **126**, 125305.
- 6 G. P. Gakis, C. Vahlas, H. Vergnes, S. Dourdain, Y. Tison, H. Martinez, J. Bour, D. Ruch, A. G. Boudouvis, B. Caussat and E. Scheid, Investigation of the initial deposition steps and the interfacial layer of Atomic Layer Deposited (ALD) Al₂O₃ on Si, *Appl. Surf. Sci.*, 2019, **492**, 245–254.
- 7 G. P. Gakis, H. Vergnes, E. Scheid, C. Vahlas, A. G. Boudouvis and B. Caussat, Detailed investigation of the surface mechanisms and their interplay with transport phenomena in alumina atomic layer deposition from TMA and water, *Chem. Eng. Sci.*, 2019, **195**, 399–412.
- 8 R. Sharma, Newer Methods of Nanoparticle Synthesis: Nitroimidazole properties with Nanometal oxides in Polymer Cages as Drug-Biomarker Monitors, *Nat. Preced.*, 2009, DOI: [10.1038/npre.2009.3952.1](https://doi.org/10.1038/npre.2009.3952.1).
- 9 G. P. Gakis, S. Termine, A.-F. A. Trompeta, I. G. Aviziotis and C. A. Charitidis, Unraveling the mechanisms of carbon nanotube growth by chemical vapor deposition, *Chem. Eng. J.*, 2022, **445**, 136807.
- 10 G. P. Gakis, E. N. Skountzos, I. G. Aviziotis and C. A. Charitidis, Multi-parametric analysis of the CVD of CNTs: Effect of reaction temperature, pressure and acetylene flow rate, *Chem. Eng. Sci.*, 2023, **267**, 118374.
- 11 Q. Tang, Z. Zhou and Z. Chen, Graphene-related nanomaterials: tuning properties by functionalization, *Nanoscale*, 2013, **5**, 4541–4583.
- 12 X. Y. Wong, A. Sena-Torralba, R. Álvarez-Diduk, K. Muthoosamy and A. Merkoçi, Nanomaterials for Nanotheranostics: Tuning Their Properties According to Disease Needs, *ACS Nano*, 2020, **14**, 2585–2627.
- 13 Q. Yao, Z. Yu, L. Li and X. Huang, Strain and Surface Engineering of Multicomponent Metallic Nanomaterials with Unconventional Phases, *Chem. Rev.*, 2023, **123**, 9676–9717.
- 14 G. Yang, L. Sun and Q. Zhang, Multicomponent chiral plasmonic hybrid nanomaterials: recent advances in synthesis and applications, *Nanoscale Adv.*, 2024, **6**, 318–336.
- 15 O. Vasylykiv and Y. Sakka, Nanoexplosion Synthesis of Multimetal Oxide Ceramic Nanopowders, *Nano Lett.*, 2005, **5**, 2598–2604.
- 16 Y. Zhang and S.-J. Park, Stabilizing CuPd bimetallic alloy nanoparticles deposited on holey carbon nitride for selective hydroxylation of benzene to phenol, *J. Catal.*, 2019, **379**, 154–163.
- 17 S. Alex and A. Tiwari, Functionalized Gold Nanoparticles: Synthesis, Properties and Applications; A Review, *J. Nanosci. Nanotechnol.*, 2015, **15**, 1869–1894.
- 18 S. Patil, R. Tandon and N. Tandon, A current research on silica coated ferrite nanoparticle and their application: Review, *Curr. Res. Green Sustainable Chem.*, 2021, **4**, 100063.
- 19 R. Medhi, M. D. Marquez and T. R. Lee, Visible-Light-Active Doped Metal Oxide Nanoparticles: Review of their Synthesis, Properties, and Applications, *ACS Appl. Nano Mater.*, 2020, **3**, 6156–6185.
- 20 M. A. Hamzawy, A. M. Abo-youssef, H. F. Salem and S. A. Mohammed, Antitumor activity of intratracheal inhalation of temozolomide (TMZ) loaded into gold nanoparticles and/or liposomes against urethane-induced lung cancer in BALB/c mice, *Drug Delivery*, 2017, **24**, 599–607.
- 21 A. K. Verma and S. Perlman, Lipid nanoparticle-mRNA: another step in the fight against COVID-19, *Cell Res.*, 2022, **32**, 421–422.
- 22 M. Manigrasso, C. Protano, M. L. Astolfi, L. Massimi, P. Avino, M. Vitali and S. Canepari, Evidences of copper nanoparticle exposure in indoor environments: Long-term assessment, high-resolution field emission scanning electron microscopy evaluation, in silico respiratory dosimetry study and possible health implications, *Sci. Total Environ.*, 2019, **653**, 1192–1203.
- 23 M. J. Bessa, F. Brandão, M. Viana, J. F. Gomes, E. Monfort, F. R. Cassee, S. Fraga and J. P. Teixeira, Nanoparticle exposure and hazard in the ceramic industry: an overview of potential sources, toxicity and health effects, *Environ. Res.*, 2020, **184**, 109297.
- 24 S. Sonwani, S. Madaan, J. Arora, S. Suryanarayan, D. Rangra, N. Mongia, T. Vats and P. Saxena, Inhalation Exposure to Atmospheric Nanoparticles and Its Associated Impacts on Human Health: A Review, *Front. Sustain. Cities*, 2021, **3**, 690444.
- 25 Y. Nazarenko, T. W. Han, P. J. Liou and G. Mainelis, Potential for exposure to engineered nanoparticles from nanotechnology-based consumer spray products, *J. Exposure Sci. Environ. Epidemiol.*, 2011, **21**, 515–528.
- 26 M. Mortimer and P. A. Holden, in *Exposure to Engineered Nanomaterials in the Environment*, ed. N. Marmiroli, J. C. White and J. Song, Elsevier, 2019, pp. 61–103, DOI: [10.1016/B978-0-12-814835-8.00003-0](https://doi.org/10.1016/B978-0-12-814835-8.00003-0).
- 27 G. V. Lowry, E. M. Hotze, E. S. Bernhardt, D. D. Dionysiou, J. A. Pedersen, M. R. Wiesner and B. Xing, Environmental Occurrences, Behavior, Fate, and Ecological Effects of Nanomaterials: An Introduction to the Special Series, *J. Environ. Qual.*, 2010, **39**, 1867–1874.
- 28 J. Blasco, I. Corsi and V. Matranga, Particles in the oceans: Implication for a safe marine environment, *Mar. Environ. Res.*, 2015, **111**, 1–4.
- 29 V. Kumar, N. Sharma and S. S. Maitra, In vitro and in vivo toxicity assessment of nanoparticles, *Int. Nano Lett.*, 2017, **7**, 243–256.
- 30 D. T. Savage, J. Z. Hilt and T. D. Dziubla, in *Nanotoxicity: Methods and Protocols*, ed. Q. Zhang, Springer New York, New York, NY, 2019, pp. 1–29, DOI: [10.1007/978-1-4939-8916-4_1](https://doi.org/10.1007/978-1-4939-8916-4_1).



- 31 F. C. Simeone and A. L. Costa, Assessment of cytotoxicity of metal oxide nanoparticles on the basis of fundamental physical–chemical parameters: a robust approach to grouping, *Environ. Sci.: Nano*, 2019, **6**, 3102–3112.
- 32 G. P. Gakis, A. Krikas, P. Neofytou, L. Tran and C. Charitidis, Modelling the biodistribution of inhaled gold nanoparticles in rats with interspecies extrapolation to humans, *Toxicol. Appl. Pharmacol.*, 2022, **457**, 116322.
- 33 A. Krikas, P. Neofytou, G. P. Gakis, I. Xiarchos, C. Charitidis and L. Tran, Modeling of clearance, retention, and translocation of inhaled gold nanoparticles in rats, *Inhalation Toxicol.*, 2022, **34**, 361–379.
- 34 I. Xiarchos, A. K. Morozinis, P. Kavouras and C. A. Charitidis, Nanocharacterization, Materials Modeling, and Research Integrity as Enablers of Sound Risk Assessment: Designing Responsible Nanotechnology, *Small*, 2020, **16**, 2001590.
- 35 M. Kotzabasaki, I. Sotiropoulos, C. Charitidis and H. Sarimveis, Machine learning methods for multi-walled carbon nanotubes (MWCNT) genotoxicity prediction, *Nanoscale Adv.*, 2021, **3**, 3167–3176.
- 36 C. F. Jones and D. W. Grainger, In vitro assessments of nanomaterial toxicity, *Adv. Drug Delivery Rev.*, 2009, **61**, 438–456.
- 37 S. G. Ciappellano, E. Tedesco, M. Venturini and F. Benetti, In vitro toxicity assessment of oral nanocarriers, *Adv. Drug Delivery Rev.*, 2016, **106**, 381–401.
- 38 C. Hirsch, M. Roesslein, H. F. Krug and P. Wick, Nanomaterial cell interactions: are current in vitro tests reliable?, *Nanomedicine*, 2011, **6**, 837–847.
- 39 K. L. Aillon, Y. Xie, N. El-Gendy, C. J. Berkland and M. L. Forrest, Effects of nanomaterial physicochemical properties on in vivo toxicity, *Adv. Drug Delivery Rev.*, 2009, **61**, 457–466.
- 40 S. T. Stern, P. P. Adiseshaiah and R. M. Crist, Autophagy and lysosomal dysfunction as emerging mechanisms of nanomaterial toxicity, *Part. Fibre Toxicol.*, 2012, **9**, 20.
- 41 A. Kermanizadeh, D. Balharry, H. Wallin, S. Loft and P. Møller, Nanomaterial translocation—the biokinetics, tissue accumulation, toxicity and fate of materials in secondary organs—a review, *Crit. Rev. Toxicol.*, 2015, **45**, 837–872.
- 42 A. Rhazouani, H. Gamrani, M. El Achaby, K. Aziz, L. Gebrati, M. S. Uddin and F. Aziz, Synthesis and Toxicity of Graphene Oxide Nanoparticles: A Literature Review of *In Vitro* and *In Vivo* Studies, *BioMed Res. Int.*, 2021, **2021**, 5518999.
- 43 R. Landsiedel, U. G. Sauer, L. Ma-Hock, J. Schnekenburger and M. Wiemann, Pulmonary toxicity of nanomaterials: a critical comparison of published in vitro assays and in vivo inhalation or instillation studies, *Nanomedicine*, 2014, **9**, 2557–2585.
- 44 A. A. Buglak, A. V. Zherdev and B. B. Dzantiev, Nano-(Q)SAR for Cytotoxicity Prediction of Engineered Nanomaterials, *Molecules*, 2019, **24**(24), 4537.
- 45 C. Oksel, C. Y. Ma, J. J. Liu, T. Wilkins and X. Z. Wang, (Q) SAR modelling of nanomaterial toxicity: A critical review, *Particuology*, 2015, **21**, 1–19.
- 46 L. Lamon, D. Asturiol, A. Vilchez, R. Ruperez-Illescas, J. Cabellos, A. Richarz and A. Worth, Computational models for the assessment of manufactured nanomaterials: Development of model reporting standards and mapping of the model landscape, *Comput. Toxicol.*, 2019, **9**, 143–151.
- 47 T. Puzyn, B. Rasulev, A. Gajewicz, X. Hu, T. P. Dasari, A. Michalkova, H. M. Hwang, A. Toropov, D. Leszczynska and J. Leszczynski, Using nano-QSAR to predict the cytotoxicity of metal oxide nanoparticles, *Nat. Nanotechnol.*, 2011, **6**, 175–178.
- 48 S. Kar, K. Pathakoti, P. B. Tchounwou, D. Leszczynska and J. Leszczynski, Evaluating the cytotoxicity of a large pool of metal oxide nanoparticles to *Escherichia coli*: Mechanistic understanding through In Vitro and In Silico studies, *Chemosphere*, 2021, **264**, 128428.
- 49 V. V. Kleandrova, F. Luan, H. González-Díaz, J. M. Ruso, A. Speck-Planche and M. N. D. S. Cordeiro, Computational Tool for Risk Assessment of Nanomaterials: Novel QSTR-Perturbation Model for Simultaneous Prediction of Ecotoxicity and Cytotoxicity of Uncoated and Coated Nanoparticles under Multiple Experimental Conditions, *Environ. Sci. Technol.*, 2014, **48**, 14686–14694.
- 50 A. A. Toropov and A. P. Toropova, Quasi-QSAR for mutagenic potential of multi-walled carbon-nanotubes, *Chemosphere*, 2015, **124**, 40–46.
- 51 J. Li, C. Wang, L. Yue, F. Chen, X. Cao and Z. Wang, Nano-QSAR modeling for predicting the cytotoxicity of metallic and metal oxide nanoparticles: A review, *Ecotoxicol. Environ. Saf.*, 2022, **243**, 113955.
- 52 J. Cao, Y. Pan, Y. Jiang, R. Qi, B. Yuan, Z. Jia, J. Jiang and Q. Wang, Computer-aided nanotoxicology: risk assessment of metal oxide nanoparticles via nano-QSAR, *Green Chem.*, 2020, **22**, 3512–3521.
- 53 A. Gajewicz, N. Schaeublin, B. Rasulev, S. Hussain, D. Leszczynska, T. Puzyn and J. Leszczynski, Towards understanding mechanisms governing cytotoxicity of metal oxides nanoparticles: Hints from nano-QSAR studies, *Nanotoxicology*, 2015, **9**, 313–325.
- 54 V. V. Kleandrova, F. Luan, H. González-Díaz, J. M. Ruso, A. Melo, A. Speck-Planche and M. N. D. S. Cordeiro, Computational ecotoxicology: Simultaneous prediction of ecotoxic effects of nanoparticles under different experimental conditions, *Environ. Int.*, 2014, **73**, 288–294.
- 55 F. Luan, V. V. Kleandrova, H. González-Díaz, J. M. Ruso, A. Melo, A. Speck-Planche and M. N. D. S. Cordeiro, Computer-aided nanotoxicology: assessing cytotoxicity of nanoparticles under diverse experimental conditions by using a novel QSTR-perturbation approach, *Nanoscale*, 2014, **6**, 10623–10630.
- 56 R. Concu, V. V. Kleandrova, A. Speck-Planche and M. N. D. S. Cordeiro, Probing the toxicity of nanoparticles: a unified in silico machine learning model based on perturbation theory, *Nanotoxicology*, 2017, **11**, 891–906.



- 57 M. K. Ha, T. X. Trinh, J. S. Choi, D. Maulina, H. G. Byun and T. H. Yoon, Toxicity Classification of Oxide Nanomaterials: Effects of Data Gap Filling and PChem Score-based Screening Approaches, *Sci. Rep.*, 2018, **8**, 3141.
- 58 Y. Feng, Y. Chang, K. Xu, R. Zheng, X. Wu, Y. Cheng and H. Zhang, Safety-by-Design of Metal Oxide Nanoparticles Based on the Regulation of their Energy Edges, *Small*, 2020, **16**, 1907643.
- 59 E. Burello and A. Worth, in *Towards Efficient Designing of Safe Nanomaterials: Innovative Merge of Computational Approaches and Experimental Techniques*, The Royal Society of Chemistry, 2012, pp. 257–283, DOI: [10.1039/9781849735476-00257](https://doi.org/10.1039/9781849735476-00257).
- 60 E. Burello and A. P. Worth, A theoretical framework for predicting the oxidative stress potential of oxide nanoparticles, *Nanotoxicology*, 2011, **5**, 228–235.
- 61 C. Kaweeterawat, A. Ivask, R. Liu, H. Zhang, C. H. Chang, C. Low-Kam, H. Fischer, Z. Ji, S. Pokhrel, Y. Cohen, D. Telesca, J. Zink, L. Madler, P. A. Holden, A. Nel and H. Godwin, Toxicity of metal oxide nanoparticles in *Escherichia coli* correlates with conduction band and hydration energies, *Environ. Sci. Technol.*, 2015, **49**, 1105–1112.
- 62 R. Liu, H. Y. Zhang, Z. X. Ji, R. Rallo, T. Xia, C. H. Chang, A. Nel and Y. Cohen, Development of structure–activity relationship for metal oxide nanoparticles, *Nanoscale*, 2013, **5**, 5644–5653.
- 63 H. Zhang, Z. Ji, T. Xia, H. Meng, C. Low-Kam, R. Liu, S. Pokhrel, S. Lin, X. Wang, Y.-P. Liao, M. Wang, L. Li, R. Rallo, R. Damoiseaux, D. Telesca, L. Mädler, Y. Cohen, J. I. Zink and A. E. Nel, Use of Metal Oxide Nanoparticle Band Gap To Develop a Predictive Paradigm for Oxidative Stress and Acute Pulmonary Inflammation, *ACS Nano*, 2012, **6**, 4349–4368.
- 64 G. P. Gakis, I. G. Aviziotis and C. A. Charitidis, Metal and metal oxide nanoparticle toxicity: moving towards a more holistic structure–activity approach, *Environ. Sci.: Nano*, 2023, **10**, 761–780.
- 65 A. Mikolajczyk, N. Sizochenko, E. Mulkiewicz, A. Malankowska, M. Nischk, P. Jurczak, S. Hirano, G. Nowaczyk, A. Zaleska-Medynska, J. Leszczynski, A. Gajewicz and T. Puzyn, Evaluating the toxicity of TiO₂-based nanoparticles to Chinese hamster ovary cells and *Escherichia coli*: a complementary experimental and computational approach, *Beilstein J. Nanotechnol.*, 2017, **8**, 2171–2180.
- 66 A. Mikolajczyk, A. Gajewicz, E. Mulkiewicz, B. Rasulev, M. Marchelek, M. Diak, S. Hirano, A. Zaleska-Medynska and T. Puzyn, Nano-QSAR modeling for ecosafe design of heterogeneous TiO₂-based nano-photocatalysts, *Environ. Sci.: Nano*, 2018, **5**, 1150–1160.
- 67 A. Mikolajczyk, N. Sizochenko, E. Mulkiewicz, A. Malankowska, B. Rasulev and T. Puzyn, A chemoinformatics approach for the characterization of hybrid nanomaterials: safer and efficient design perspective, *Nanoscale*, 2019, **11**, 11808–11818.
- 68 F. Stoliński, A. Rybińska-Fryca, M. Gromelski, A. Mikolajczyk and T. Puzyn, NanoMixHamster: a web-based tool for predicting cytotoxicity of TiO₂-based multicomponent nanomaterials toward Chinese hamster ovary (CHO-K1) cells, *Nanotoxicology*, 2022, **16**, 276–289.
- 69 A. Rybińska-Fryca, A. Mikolajczyk and T. Puzyn, Structure–activity prediction networks (SAPNets): a step beyond Nano-QSAR for effective implementation of the safe-by-design concept, *Nanoscale*, 2020, **12**, 20669–20676.
- 70 T. C. Le, H. Yin, R. Chen, Y. Chen, L. Zhao, P. S. Casey, C. Chen and D. A. Winkler, An Experimental and Computational Approach to the Development of ZnO Nanoparticles that are Safe by Design, *Small*, 2016, **12**, 3568–3577.
- 71 G. P. Gakis, I. G. Aviziotis and C. A. Charitidis, A structure–activity approach towards the toxicity assessment of multicomponent metal oxide nanomaterials, *Nanoscale*, 2023, **15**, 16432–16446.
- 72 F. Zhang, Z. Wang, W. J. G. M. Peijnenburg and M. G. Vijver, Review and Prospects on the Ecotoxicity of Mixtures of Nanoparticles and Hybrid Nanomaterials, *Environ. Sci. Technol.*, 2022, **56**, 15238–15250.
- 73 J. Bozich, M. Hang, R. Hamers and R. Klaper, Core chemistry influences the toxicity of multicomponent metal oxide nanomaterials, lithium nickel manganese cobalt oxide, and lithium cobalt oxide to *Daphnia magna*, *Environ. Toxicol. Chem.*, 2017, **36**, 2493–2502.
- 74 A. M. Z. de Medeiros, L. U. Khan, G. H. da Silva, C. A. Ospina, O. L. Alves, V. L. de Castro and D. S. T. Martinez, Graphene oxide-silver nanoparticle hybrid material: an integrated nanosafety study in zebrafish embryos, *Ecotoxicol. Environ. Saf.*, 2021, **209**, 111776.
- 75 T. X. Trinh and J. Kim, Status Quo in Data Availability and Predictive Models of Nano-Mixture Toxicity, *Nanomaterials*, 2021, **11**(1), 124.
- 76 J. Jang, H. G. Hur, M. J. Sadowsky, M. N. Byappanahalli, T. Yan and S. Ishii, Environmental *Escherichia coli*: ecology and public health implications—a review, *J. Appl. Microbiol.*, 2017, **123**, 570–581.
- 77 M. Sautour, J.-P. Lemaître, L. Ranjard, C. Truntzer, L. Basmaciyan, G. Depret, A. Hartmann and F. Dalle, Detection and survival of *Candida albicans* in soils, *Environ. DNA*, 2021, **3**, 1093–1101.
- 78 H. K. Abbas, J. R. Wilkinson, R. M. Zablotowicz, C. Accinelli, C. A. Abel, H. A. Bruns and M. A. Weaver, Ecology of *Aspergillus flavus*, regulation of aflatoxin production, and management strategies to reduce aflatoxin contamination of corn, *Toxin Rev.*, 2009, **28**, 142–153.
- 79 J. Portier, G. Campet, J. Etourneau and B. Tanguy, A simple model for the estimation of electronegativities of cations in different electronic states and coordinations, *J. Alloys Compd.*, 1994, **209**, 285–289.
- 80 J. Portier, H. S. Hilal, I. Saadeddin, S. J. Hwang, M. A. Subramanian and G. Campet, Thermodynamic correlations and band gap calculations in metal oxides, *Prog. Solid State Chem.*, 2004, **32**, 207–217.



- 81 P. Ambure, R. B. Aher, A. Gajewicz, T. Puzyn and K. Roy, "NanoBRIDGES" software: Open access tools to perform QSAR and nano-QSAR modeling, *Chemom. Intell. Lab. Syst.*, 2015, **147**, 1–13.
- 82 B. Yuan, P. Wang, L. Sang, J. Gong, Y. Pan and Y. Hu, QNAR modeling of cytotoxicity of mixing nano-TiO₂ and heavy metals, *Ecotoxicol. Environ. Saf.*, 2021, **208**, 111634.
- 83 L. Sang, Y. Wang, C. Zong, P. Wang, H. Zhang, D. Guo, B. Yuan and Y. Pan, Machine Learning for Evaluating the Cytotoxicity of Mixtures of Nano-TiO₂ and Heavy Metals: QSAR Model Apply Random Forest Algorithm after Clustering Analysis, *Molecules*, 2022, **27**(18), 6125.
- 84 F. Zhang, Z. Wang, W. J. G. M. Peijnenburg and M. G. Vijver, Machine learning-driven QSAR models for predicting the mixture toxicity of nanoparticles, *Environ. Int.*, 2023, **177**, 108025.
- 85 T. X. Trinh, M. Seo, T. H. Yoon and J. Kim, Developing random forest based QSAR models for predicting the mixture toxicity of TiO₂ based nano-mixtures to *Daphnia magna*, *NanoImpact*, 2022, **25**, 100383.
- 86 T. X. Trinh and J. Kim, Status Quo in Data Availability and Predictive Models of Nano-Mixture Toxicity, *Nanomaterials*, 2021, **11**, 124.
- 87 I. Guyon and A. Elisseeff, An introduction to variable and feature selection, *J. Mach. Learn. Res.*, 2003, **3**, 1157–1182.
- 88 M. Eklund, U. Norinder, S. Boyer and L. Carlsson, Choosing Feature Selection and Learning Algorithms in QSAR, *J. Chem. Inf. Model.*, 2014, **54**, 837–843.
- 89 Y. Li, Z. Dai, D. Cao, F. Luo, Y. Chen and Z. Yuan, Chi-MIC-share: a new feature selection algorithm for quantitative structure–activity relationship models, *RSC Adv.*, 2020, **10**, 19852–19860.
- 90 OECD, Guidance Document on the Validation of (Quantitative) Structure-Activity Relationship [(Q)SAR] Models, 2014.
- 91 F. Sahigara, D. Ballabio, R. Todeschini and V. Consonni, Assessing the Validity of QSARs for Ready Biodegradability of Chemicals: An Applicability Domain Perspective, *Curr. Comput.-Aided Drug Des.*, 2014, **10**, 137–147.
- 92 F. Sahigara, K. Mansouri, D. Ballabio, A. Mauri, V. Consonni and R. Todeschini, Comparison of Different Approaches to Define the Applicability Domain of QSAR Models, *Molecules*, 2012, **17**, 4791–4810.
- 93 A. Rácz, D. Bajusz and K. Héberger, Interrelation Limits in Molecular Descriptor Preselection for QSAR/QSPR, *Mol. Inf.*, 2019, **38**, 1800154.
- 94 M. Auffan, J. Rose, M. R. Wiesner and J. Y. Bottero, Chemical stability of metallic nanoparticles: a parameter controlling their potential cellular toxicity in vitro, *Environ. Pollut.*, 2009, **157**, 1127–1133.
- 95 G. S. Plumlee, S. A. Morman and T. L. Ziegler, The Toxicological Geochemistry of Earth Materials: An Overview of Processes and the Interdisciplinary Methods Used to Understand Them, *Rev. Mineral. Geochem.*, 2006, **64**, 5–57.
- 96 Y. Li, W. Zhang, J. Niu and Y. Chen, Mechanism of Photogenerated Reactive Oxygen Species and Correlation with the Antibacterial Properties of Engineered Metal-Oxide Nanoparticles, *ACS Nano*, 2012, **6**, 5164–5173.
- 97 A. B. Djuricic, Y. H. Leung, A. M. Ng, X. Y. Xu, P. K. Lee, N. Degger and R. S. Wu, Toxicity of metal oxide nanoparticles: mechanisms, characterization, and avoiding experimental artefacts, *Small*, 2015, **11**, 26–44.
- 98 M. Premanathan, K. Karthikeyan, K. Jeyasubramanian and G. Manivannan, Selective toxicity of ZnO nanoparticles toward Gram-positive bacteria and cancer cells by apoptosis through lipid peroxidation, *Nanomed.: Nanotechnol. Biol. Med.*, 2011, **7**, 184–192.
- 99 H. L. Karlsson, P. Cronholm, J. Gustafsson and L. Möller, Copper Oxide Nanoparticles Are Highly Toxic: A Comparison between Metal Oxide Nanoparticles and Carbon Nanotubes, *Chem. Res. Toxicol.*, 2008, **21**, 1726–1732.
- 100 G. Applerot, A. Lipovsky, R. Dror, N. Perkash, Y. Nitzan, R. Lubart and A. Gedanken, Enhanced Antibacterial Activity of Nanocrystalline ZnO Due to Increased ROS-Mediated Cell Injury, *Adv. Funct. Mater.*, 2009, **19**, 842–852.
- 101 T. Xia, M. Kovochich, J. Brant, M. Hotze, J. Sempf, T. Oberley, C. Sioutas, J. I. Yeh, M. R. Wiesner and A. E. Nel, Comparison of the Abilities of Ambient and Manufactured Nanoparticles To Induce Cellular Toxicity According to an Oxidative Stress Paradigm, *Nano Lett.*, 2006, **6**, 1794–1807.
- 102 T. C. Long, N. Saleh, R. D. Tilton, G. V. Lowry and B. Veronesi, Titanium Dioxide (P25) Produces Reactive Oxygen Species in Immortalized Brain Microglia (BV2): Implications for Nanoparticle Neurotoxicity, *Environ. Sci. Technol.*, 2006, **40**, 4346–4352.
- 103 A. L. Neal, What can be inferred from bacterium–nanoparticle interactions about the potential consequences of environmental exposure to nanoparticles?, *Ecotoxicology*, 2008, **17**, 362.
- 104 F. Thevenod, Catch me if you can! Novel aspects of cadmium transport in mammalian cells, *BioMetals*, 2010, **23**, 857–875.
- 105 S. Sugiharto, T. M. Lewis, A. J. Moorhouse, P. R. Schofield and P. H. Barry, Anion-cation permeability correlates with hydrated counterion size in glycine receptor channels, *Biophys. J.*, 2008, **95**, 4698–4715.
- 106 R. Epsztein, E. Shaulsky, M. Qin and M. Elimelech, Activation behavior for ion permeation in ion-exchange membranes: Role of ion dehydration in selective transport, *J. Membr. Sci.*, 2019, **580**, 316–326.
- 107 S. Adapa and A. Malani, Role of hydration energy and cations association on monovalent and divalent cations adsorption at mica-aqueous interface, *Sci. Rep.*, 2018, **8**, 12198.
- 108 Z. Qu and H. C. Hartzell, Anion permeation in Ca(2+)-activated Cl(−) channels, *J. Gen. Physiol.*, 2000, **116**, 825–844.
- 109 Y. Mu, F. Wu, Q. Zhao, R. Ji, Y. Qie, Y. Zhou, Y. Hu, C. Pang, D. Hristozov, J. P. Giesy and B. Xing, Predicting toxic potencies of metal oxide nanoparticles by means of nano-QSARs, *Nanotoxicology*, 2016, **10**, 1207–1214.



- 110 G. E. Brown, V. E. Henrich, W. H. Casey, D. L. Clark, C. Eggleston, A. Felmy, D. W. Goodman, M. Grätzel, G. Maciel, M. I. McCarthy, K. H. Nealson, D. A. Sverjensky, M. F. Toney and J. M. Zachara, Metal Oxide Surfaces and Their Interactions with Aqueous Solutions and Microbial Organisms, *Chem. Rev.*, 1999, **99**, 77–174.
- 111 M. Balali-Mood, K. Naseri, Z. Tahergorabi, M. R. Khazdair and M. Sadeghi, Toxic Mechanisms of Five Heavy Metals: Mercury, Lead, Chromium, Cadmium, and Arsenic, *Front. Pharmacol.*, 2021, **12**, 643972.
- 112 A. Manuja, B. Kumar, R. Kumar, D. Chhabra, M. Ghosh, M. Manuja, B. Brar, Y. Pal, B. N. Tripathi and M. Prasad, Metal/metal oxide nanoparticles: Toxicity concerns associated with their physical state and remediation for biomedical applications, *Toxicol. Rep.*, 2021, **8**, 1970–1978.
- 113 V. Forest, J.-F. Hochepeid, L. Leclerc, A. Trouvé, K. Abdelkebir, G. Sarry, V. Augusto and J. Pourchez, Towards an alternative to nano-QSAR for nanoparticle toxicity ranking in case of small datasets, *J. Nanopart. Res.*, 2019, **21**, 95.
- 114 M. Jaishankar, T. Tseten, N. Anbalagan, B. B. Mathew and K. N. Beeregowda, Toxicity, mechanism and health effects of some heavy metals, *Interdiscip. Toxicol.*, 2014, **7**, 60–72.
- 115 S. J. Stohs and D. Bagchi, Oxidative mechanisms in the toxicity of metal ions, *Free Radical Biol. Med.*, 1995, **18**, 321–336.
- 116 A. Abdal Dayem, M. K. Hossain, S. B. Lee, K. Kim, S. K. Saha, G.-M. Yang, H. Y. Choi and S.-G. Cho, The Role of Reactive Oxygen Species (ROS) in the Biological Activities of Metallic Nanoparticles, *Int. J. Mol. Sci.*, 2017, **18**, 120.
- 117 R. R. Crichton, in *Metal Chelation in Medicine*, The Royal Society of Chemistry, 2017, pp. 1–23, DOI: [10.1039/9781782623892-00001](https://doi.org/10.1039/9781782623892-00001).
- 118 N. Mei, Y. Zhang, Y. Chen, X. Guo, W. Ding, S. F. Ali, A. S. Biris, P. Rice, M. M. Moore and T. Chen, Silver nanoparticle-induced mutations and oxidative stress in mouse lymphoma cells, *Environ. Mol. Mutagen.*, 2012, **53**, 409–419.
- 119 F. Chen, L. Wu, X. Xiao, L. Rong, M. Li and X. Zou, Mixture toxicity of zinc oxide nanoparticle and chemicals with different mode of action upon *Vibrio fischeri*, *Environ. Sci. Eur.*, 2020, **32**, 41.
- 120 D. Wang, S. Wang, L. Bai, M. S. Nasir, S. Li and W. Yan, Mathematical Modeling Approaches for Assessing the Joint Toxicity of Chemical Mixtures Based on Luminescent Bacteria: A Systematic Review, *Front. Microbiol.*, 2020, **11**, 1651.
- 121 M. Na, S. H. Nam, K. Moon and J. Kim, Development of a nano-QSAR model for predicting the toxicity of nano-metal oxide mixtures to *Aliivibrio fischeri*, *Environ. Sci.: Nano*, 2023, **10**, 325–337.

