

## Extended information

Machine learning-based identification of key biotic and abiotic drivers of mineral weathering rate in a complex enhanced weathering experiment

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## A Data processing

### A.1 Calculation of change in amount of dissolved inorganic carbon, solid inorganic carbon, total inorganic carbon and total alkalinity

This section outlines the methodology used to calculate the changes in the amount of dissolved inorganic carbon (*DIC*), solid inorganic carbon (*SIC*), total inorganic carbon (*TIC*), and total alkalinity (*TA*). Detailed equations and relevant measurements are provided to ensure clarity and reproducibility.

#### A.1.1 Change in dissolved inorganic carbon

The change in amount of DIC ( $\Delta DIC$ ) is defined as the difference between the final ( $DIC_f$ ) and the initial *DIC* present at the beginning of the experiment ( $DIC_0$ ):

$$\Delta DIC [mmol] = DIC_f [mmol] - DIC_0 [mmol] \quad (A.1)$$

Table A1: Concentrations of dissolved inorganic carbon ( $DIC$ ) and total alkalinity ( $TA$ ), measured in leached water of blank columns, with a solid bed consisting only of sand, biochar and organic amendments.

	Solid bed	$V_{irr}$ [l]	$DIC$ [ $\frac{mg}{l}$ ]	$TA$ [ $\frac{\mu mol}{l}$ ]
$B_1$	400g sand	5.6	11.875	1434.9
$B_2$	400g sand	5.6	11.088	1354.5
$B_3$	400g sand	8.4	11.783	1427.3
$B_{straw,1}$	400g sand + 10g straw	5.6	35.169	1981.1
$B_{straw,2}$	400g sand + 10g straw	8.4	13.589	1632.9
$B_{digestate,1}$	400g sand + 10g digestate	5.6	10.502	1445.4
$B_{alfalfa,1}$	400g sand + 10g alfalfa mixture	5.6	26.387	2304.8
$B_{biochar,1}$	400g sand + 20g biochar	5.6	15.452	1761.3
$B_{biochar,2}$	400g sand + 40g biochar	5.6	18.387	2053.5
$B_{biochar,3}$	400g sand + 20g biochar	8.4	14.708	1698.6

Here,  $DIC_f$  is calculated using the  $DIC$  concentration measured after 8 weeks ( $DIC_{conc}$ ) and the irrigated volume ( $V_{irr}$ ):

$$DIC_f [mmol] = DIC_{conc} \left[ \frac{mg}{l} \right] \cdot V_{irr} [l] \cdot \frac{1 [mmol]}{12.001 [mg]} \quad (A.2)$$

The  $DIC_0$  is the sum of the  $DIC$  present in the irrigated water and the  $DIC$  stemming from the organic amendments and biochar. To calculate the  $DIC$  stemming from the irrigated water, the  $DIC$  concentration of columns with a solid bed consisting only of 400g sand (coined  $B$ ), was measured ( $DIC_{conc}^B$ ) and averaged (Tab. A1).

$$DIC_{conc}^{water} \left[ \frac{mg}{l} \right] = \overline{DIC_{conc}^{B_i} \left[ \frac{mg}{l} \right]} \quad (A.3)$$

The  $DIC$  stemming from the organic amendments and biochar was based on  $DIC$  concentration measurements of columns with a solid bed made out of only sand and straw ( $B_{straw}$ ), digestate ( $B_{digestate}$ ), alfalfa-dominated mixture ( $B_{alfalfa}$ ) or biochar ( $B_{biochar}$ ) (Tab. A1):

$$DIC_{conc}^{matter} \left[ \frac{mg}{g \text{ matter}} \right] = \frac{(DIC_{conc}^{B_{matter,i}} \left[ \frac{mg}{l} \right] - DIC_{conc}^{water} \left[ \frac{mg}{l} \right]) \cdot V_{irr}^{B_{matter,i}} [l]}{m_{org,i}^{B_{matter,i}} [g]} \quad (A.4)$$

Here, *matter* can be either straw, digestate, the alfalfa mixture and biochar. Finally, the initial  $DIC$  was calculated using:

$$DIC_0 [mmol] = \left( DIC_{conc}^{water} [mg] \cdot V_{irr} [l] + \sum_{matter} DIC_{conc}^{matter} \left[ \frac{mg}{g \text{ matter}} \right] \cdot m_{matter} [g] \right) \cdot \frac{1 [mmol]}{12.001 [mg]} \quad (A.5)$$

Here, the summation goes over straw, digestate, the alfalfa mixture and biochar.

### A.1.2 Change in amount of solid inorganic carbon

Analogously to  $\Delta DIC$ , change in amount of SIC ( $\Delta SIC$ ) is the difference between the final ( $SIC_f$ ) and the initial  $SIC$  that is added to the system through the input materials ( $SIC_0$ ):

$$\Delta SIC [mmol] = SIC_f [mmol] - SIC_0 [mmol] \quad (A.6)$$

Before measuring the  $SIC$  concentration ( $SIC_{conc}$ ) of the solid samples (using *C analysis, Flash 2000 CN Soil Analyser, Interscience, Antwerp, Belgium*), loss of ignition (LOI) is performed, which lead to mass loss. Therefore,  $SIC_f$  is calculated using:

$$SIC_f [mmol] = 10 \cdot SIC_{conc} [mass\%] \cdot LOI \text{ correction} \cdot m_{solid} [g] \cdot \frac{1 [mmol]}{12.001 [mg]} \quad (A.7)$$

where  $m_{solid}$  is the mass of the added minerals, organic amendments and biochar; and the *LOI correction* factor accounts for sample mass loss during LOI:

$$LOI\ correction = 1 - \frac{m_{solid\ sample}^{post-LOI} [g] - m_{crucible} [g]}{m_{solid\ sample}^{pre-LOI} [g] - m_{crucible} [g]} \quad (A.8)$$

with  $m_{solid\ sample}^{pre-LOI}$  and  $m_{solid\ sample}^{post-LOI}$  the gross sample mass before and after LOI, and  $m_{crucible}$  the mass of the crucible holding the solid sample.

$SIC_0$  is the *SIC* present at the start of the experiment and is calculated based on *SIC* measurements of the added minerals, organic amendments and biochar (Tab. A2) and their masses in the column ( $m_{sample,X}$ ):

$$SIC_0 [mmol] = 10 \cdot \sum_X SIC_{conc}^X [mass\%] \cdot LOI\ correction_X \cdot m_{sample,X} [g] \cdot \frac{1 [mmol]}{12.001 [mg]} \quad (A.9)$$

where the summation goes over the added minerals, organic amendments and biochar; and the *LOI correction* factor again accounts for sample mass loss during LOI (Eq. A.8).

Table A2: *Specific surface area measurements of minerals, loss of ignition and solid inorganic carbon measurements of samples of minerals, biochar and organic amendments. SSA = Specific surface area, LOI = loss of ignition, SIC = solid inorganic carbon. SSA measurements of minerals are obtained using Brunauer-Emmett-Teller method.  $m_{solid\ sample}^{pre-LOI}$  and  $m_{solid\ sample}^{post-LOI}$  are the gross sample masses before and after LOI,  $m_{crucible}$  is the mass of the crucible holding the sample and  $SIC$  is the measured solid inorganic carbon concentration.*

Matter		SSA [m <sup>2</sup> /g]	$m_{sample,X}^{pre-LOI}$ [g]	$m_{sample,X}^{post-LOI}$ [g]	$m_{crucible,X}$ [g]	$SIC_{conc}^X$ [mass%]
Lava	≤ 0.063 mm	1.896	37.5	37.4	20.9	0.009
	0.063-2 mm	0.513	41.9	41.9	20.2	0.006
Basalt	≤ 0.063 mm	10.577	38.4	38.3	19.7	0.134
	0.063-3 mm	6.972	40.2	40.0	21.3	0.068
Dunite	≤ 0.063 mm	7.050	36.2	35.8	18.8	0.031
	0.063-0.125 mm	0.959	45.2	45.1	19.8	0.005
	0.125-1 mm	1.134	51.2	51.1	20.0	0.008
Steelslag	≤ 0.1 mm	13.246	32.6	31.8	18.8	1.922
	0.1-0.5 mm	9.154	39.9	38.9	19.3	1.858
	0.5-3 mm	4.747	38.8	38.3	18.8	1.359
	mixed (≤ 3 mm)	10.271	29.6	29.2	19.0	0.992
Digestate	-	-	22.2	20.2	20.0	2.272
Straw	-	-	21.0	20.0	20.0	4.957
Alfalfa mixture	-	-	23.7	23.3	23.3	3.339
Biochar	-	-	23.8	20.5	19.5	0.263

### A.1.3 Change in total inorganic carbon

In this study, total inorganic carbon is defined as the sum of solid inorganic carbon and dissolved inorganic carbon. Therefore, the final TIC, initial TIC and change in TIC are calculated as:

$$TIC_f [mmol] = SIC_f [mmol] + DIC_f [mmol] \quad (A.10)$$

$$TIC_0 [mmol] = SIC_0 [mmol] + DIC_0 [mmol] \quad (A.11)$$

$$\Delta TIC [mmol] = \Delta SIC [mmol] + \Delta DIC [mmol] \quad (A.12)$$

#### A.1.4 Change in total alkalinity

The change in total alkalinity ( $\Delta TA$ ) is the difference between the final ( $TA_f$ ) and the initial  $TA$  that results from non-weathering processes ( $TA_0$ ):

$$\Delta TA [mmol] = TA_f [mmol] - TA_0 [mmol] \quad (A.13)$$

The final  $TA$  is calculated from the measured  $TA$  concentration at the end of the experiment ( $TA_{conc}$ ) and the irrigated volume:

$$TA_f [mmol] = 10^{-3} TA_{conc} \left[ \frac{\mu mol}{l} \right] \cdot V_{irr} [l] \quad (A.14)$$

The initial  $TA$  is calculated analogously to the initial  $DIC$  (Equations A.3-A.5), i.e., based on  $TA$  concentration measurements of batches with a solid bed consisting only of 400g sand ( $B$ ) and batches with a solid bed made out of only organic amendments and biochar ( $B_{straw}$ ,  $B_{digestate}$ ,  $B_{alfalfa}$ ,  $B_{biochar}$ ) (Tab. A1):

$$TA_{conc}^{water} \left[ \frac{mmol}{l} \right] = 10^{-3} \overline{TA_{conc}^{B_i} \left[ \frac{\mu mol}{l} \right]} \quad (A.15)$$

$$TA_{conc}^{matter} \left[ \frac{mmol}{g \text{ matter}} \right] = 10^{-3} \frac{(TA_{conc}^{B_{matter,i}} \left[ \frac{\mu mol}{l} \right] - TA_{conc}^{water} \left[ \frac{\mu mol}{l} \right]) \cdot V_{irr}^{B_{matter,i}} [l]}{m_{matter,i}^{B_{matter,i}} [g]} \quad (A.16)$$

$$TA_0 [mmol] = TA_{conc}^{water} \left[ \frac{mmol}{l} \right] \cdot V_{irr} [l] + \sum_{matter} TA_{conc}^{matter} \left[ \frac{mmol}{g \text{ matter}} \right] \cdot m_{matter} [g] \quad (A.17)$$

where the summation goes over straw, digestate, the alfalfa mixture and biochar.

## A.2 Transformation of model features

Some features were transformed before being used as input for the machine learning (ML) models. The relevant equations for such features are provided below.

The total surface area of mineral type  $M$  is calculated using:

$$SA_M [m^2] = \sum_G^{grain\ sizes} mass_{MG} [g] \cdot SSA_{MG} \left[ \frac{m^2}{g} \right] \quad (A.18)$$

where  $mass_{MG}$  is the added mass, and  $SSA_{MG}$  is the specific surface area of mineral type  $M$  with grain size  $G$  (Table A2).

The feature *chlorotica ratio* is defined as the ratio of initially live *A. chlorotica* earthworms with respect to the total number of initially live earthworms (i.e., the sum of the number of *A. chlorotica* earthworms and the number of *A. caliginosa* earthworms):

$$chlorotica\ ratio = \frac{number\ of\ initially\ live\ A.\ chlorotica\ earthworms}{number\ of\ initially\ live\ earthworms} \quad (A.19)$$

And analogously, the feature  $\dagger$  *chlorotica ratio* is defined as:

$$\dagger\ chlorotica\ ratio = \frac{number\ of\ initially\ dead\ A.\ chlorotica\ earthworms}{number\ of\ initially\ dead\ earthworms} \quad (A.20)$$

### A.3 Analysis of noise in data

To quantify the noise in the data, some of the input combinations of the columns was repeated. This resulted in 28 combinations that were repeated at least three times for  $\Delta DIC$ , 30 for  $\Delta SIC$ , 27 for change in amount of TIC ( $\Delta TIC$ ) and 27 for change in amount of TA ( $\Delta TA$ ). For each set of repeats, the standard deviation of the weathering indicators ( $\Delta DIC$ ,  $\Delta SIC$ ,  $\Delta TIC$  and acDTA) was calculated and rescaled by dividing by the interquartile range of the indicator of the full dataset (Fig. A1). As steel slag generally lead to different scales of the indicators, this rescaling was done separately for the subset of columns with and without steel slag. For all indicators, the resulting rescaled standard deviation was relatively high, especially for  $\Delta SIC$  and, to a lesser extent,  $\Delta TIC$  (Fig. A1).

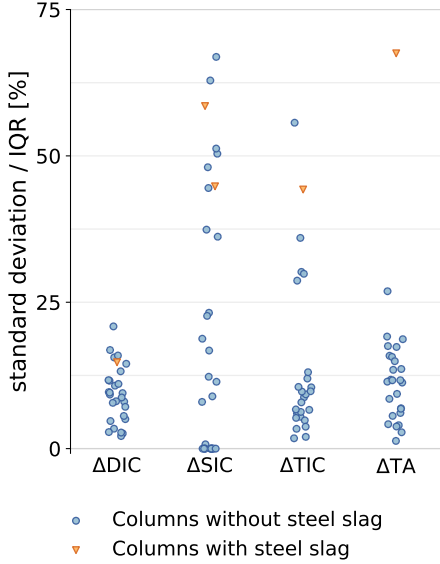


Figure A1: Quantification of the noise in data of the changes of dissolved inorganic carbon ( $\Delta DIC$ ), solid inorganic carbon ( $\Delta SIC$ ), total inorganic carbon ( $\Delta TIC$ ) and total alkalinity ( $\Delta TA$ ). The y axis depicts the standard deviations of columns that were repeated at least three times, rescaled by interquartile range (IQR) of the full dataset. The rescaling was done separately for the datasets with (orange) and without steel slag (blue).

To assess whether this noise resulted from certain inputs of the columns, we used a Random Forest (RF) to predict the standard deviations of the target values of repeated combinations for each of the indicators ( $\Delta DIC$ ,  $\Delta SIC$ ,  $\Delta TIC$ ,  $\Delta TA$ ). As the number of repeated columns was limited, leave one out cross validation (CV) was used. However, the model's performances were poor for all indicators, except for the prediction of  $\Delta SIC$  (Tab. A3). A subsequent Shapley Additive exPlanations (SHAP) analysis revealed that high standard deviations were predicted for columns with steel slag (A2). Therefore, the prediction of  $\Delta SIC$  was repeated for columns without steel slag, which resulted in poor performances (Tab. A3). This suggests that, while steel slag leads to noisy measurements for  $SIC$ , the remaining noise of the indicators likely arises from other factors than the column inputs, such as sampling inconsistencies from imperfectly homogenized columns or irregular irrigation.

		MAE	MSE	RMSE	$R^2$
$\Delta DIC$	RF	0.37	0.45	0.20	-0.50
	naive	0.30	0.38	0.14	-0.08
$\Delta SIC$	RF	3.2	8.4	70	0.73
	naive	8.2	16.5	272	-0.07
$\Delta TIC$	RF	2.8	7.9	62	-0.03
	naive	3.5	8.1	65	-0.08
$\Delta TA$	RF	1.6	5.1	26	-0.20
	naive	1.8	4.8	23	-0.08
$\Delta SIC$ (without steel slag)	RF	0.97	1.2	1.4	0.23
	naive	1.21	1.4	2.0	-0.08

Table A3: Performance of the Random Forest prediction of the standard deviation of  $\Delta DIC$ ,  $\Delta SIC$ ,  $\Delta TIC$  and  $\Delta TA$  of repeated measurements; and of  $\Delta SIC$  of repeated measurements without steel slag. Depicted performance metrics are mean absolute error (MAE), mean squared error (MSE), root mean squared error (RMSE), and determination coefficient ( $R^2$ ). The performance of the naive regressor, which predicts the mean target value for the whole dataset, is added as a baseline.

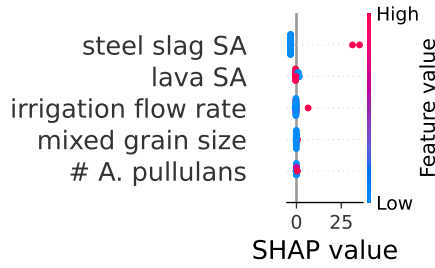


Figure A2: SHapley Additive exPlanation (SHAP) values of the top 5 most important features in the random forest prediction of the standard deviation of the change in solid inorganic carbon of repeated measurements.