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# Improved explicit formulation of bedload transport using a novel multi-level multi-model data-driven ensemble approach

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#### **ABSTRACT**

Estimation of bedload transport in rivers is a very complex and important river engineering challenge needs substantial additional efforts in pre-processing and ensemble modeling to derive the desired level of prediction accuracy. This paper aims to develop a new framework for the formulation of bedload transport in rivers using multi-level Multi-Model Ensemble (MME) approach to derive improved explicit formulations hybridized with multiple pre-processed-based models. Three pre-processing techniques of feature selection by Gamma Test (GT), dimension reduction by principal component analysis (PCA), and data clustering by subset selection of maximum dissimilarity (SSMD) are utilized at level 0. The multi-linear regression (MLR), MLR-PCA, artificial neural network (ANN), ANN-PCA, Gene expression programming (GEP), GEP-PCA, Group method of data handling (GMDH) and GMDH-PCA are used to develop individual explicit formulations at level 1, and the inferred formulas are hybridized with the MME approach at level 2 by Pareto optimality. A newly revised discrepancy ratio (RDR) for error distributions in conjunction with several statistical and graphical indicators were used to evaluate the strategy's performance. Results of MME showed that the

proposed framework acted as an efficient tool in explicit equation induction for bedload transport (i.e., 33–96% reduction of RMSE; 2-29% increase of R<sup>2</sup>, 2-138% increase of NSE and 38-98% reduction of RAE in testing step in comparison with the best individual model) and clearly outperformed estimations made by other models. The current study highlights the importance of pre-processing and multi-modelling techniques in deep learning models to encounter the challenges of function finding for complex bedload transport estimations in multiple observed datasets. 

Keywords: Multi-model ensembles approach, bedload transport, function finding, equation optimization, machine learning

#### 1- Introduction

Sediment transport in river flows leads to several challenges for the water resources tasks and is crucial in the context of reservoir sedimentation, flood control, river morphology changes, stable channel design, fish and wild life habitat, and watershed management (Van Rijn, 1993; Bhattacharya et al., 2007; Dey, 2014; Elkurdy et al., 2021; Ahmadianfar et al., 2021). In sediment transport, the coarse-grains conveyed by higher discharges and floods immediately above the bed are known as bedload (Barry, 2007).

Sediment transport is a highly complex, stochastic phenomenon with somewhat unknown theory. It is hard to measure in the field due to the time and cost-intensive process. These features of sediment transport produce high uncertainty in predictive

- 51 equations that made their applicability questionable and makes limitation on
- employing them (Bhattacharya et al., 2007; Riahi-Madvar and Seifi, 2018).
- The predictive methods of bedload transport are generally categorized into physical
- and data driven models (Kitsikoudis et al., 2014; Gholami et al., 2018 & 2019).
- 55 Considering challenges of phenomenon complexity, inaccuracies in the predictive
- 56 equations of bedload and measuring difficulties with the physical methods,
- 57 development of new data-driven-based models with an appropriate determination of
- 58 effective parameters of bedload having easily accessible field variables is vital
- 59 (Ghani et al., 2011; Gao, 2011; Ebtehaj et al., 2021).
- With the emerging applications of machine learning (ML) models, producing
- effective results in formulation of complex nonlinear challenges in river engineering,
- researchers have endeavoured to use these new techniques to cope with the
- 63 complicated nature of bedload transport in parallel with the experimental and
- physical-based studies (Bhattacharya et al., 2007; Safari et al., 2020).
- Various ML methods were implemented for sediment transport modelling such as
- artificial neural network: ANN (Afan et al., 2016; Bhattacharya et al., 2007;
- Kitsikoudis et al., 2014), fuzzy logic and adaptive neural fuzzy inference system:
- 68 ANFIS (Kitsikoudis et al., 2014; Qasem et al., 2017), support vector machine: SVM

(Roushangar and Shahnazi, 2020; Sahraei et al., 2017), genetic expression 69 programming: GEP (Danandeh Mehr et al., 2018; Ghani and Azamathulla, 2014). 70 Montes, et al., (2021), Noori et al. (2010a-c, 2011) and Liu et al (2020) figured out 71 that these techniques suffer from the generalization capabilities of the results due to 72 inappropriate selection of training set, inaccuracy issues with limited extrapolation 73 abilities when applied to unseen data set extensive than the data used in training 74 phase. They suggested pre-processing techniques such as data clustering for subset 75 selection in train and testing steps. The studies in the literature of bedload prediction, 76 have neglected mathematical-based clustering of train and test sets, while this study 77 considered a subset selection of maximum dissimilarity (SSMD) to overcome these 78 challenges. 79 The data-driven models developed so far for bedload transport estimations are 80 basically black-box type tools such as ANN, ANFIS and suffer from limited 81 interpretability of physical importance of the input parameters and their interactions 82 to the model outputs, inability to capture physical processes (Noori et al., 2010a-c, 83 2011; Montes et al., 2021; Seifi and Soroush, 2020; Madvar et al., 2020). Therefore, 84 the derivation of explicit accurate equations for bedload transport in rivers based on 85 AI models, remains challenging. 86

- 87 To address this problem, in the current study a hybridization of four mathematical
- models including ANN, GEP, group method of data handling (GMDH), and multi
- linear regression (MLR) are employed in deriving the explicit predictive equations
- 90 for bedload using a new multi-model-based strategy.
- The data-driven models are susceptible to the number of the input variable. To the
- best of our knowledge, few studies there are relating to the use of approach to reduce
- 93 the dimension of input data space and to astutely designate appropriate input
- variables for prediction of bedload in a multi-model ensemble approach.
- Generally, the bedload rate is chosen as a dependent parameter. The fluid properties,
- 96 flow conditions, sediment properties, and channel geometry are considered
- 97 independent parameters in data-driven model developments (Montes et al., 2021;
- 98 Qasem et al., 2017). In conventional ML-based models usually rely on the
- 99 researchers' subjective "suggesting" the input variables that will result in a poor
- prediction (Liu et al., 2020).
- Hence, proposing a sophisticated approach such as principal component analysis
- 102 (PCA) (Snieder et al., 2020) and Gamma test (GT) to reduce the dimension of the
- input space leading to choose proper input parameters of the model, is valuable. The
- studies in the literature neglected input vector manipulation and data dimension
- reduction for ML prediction of bedload. In contrast, the present study used PCA and

- GT techniques for dimension reduction and effective variable selection. In the current study pre-processing techniques of GT and PCA as dimension reductions are used in conjunction with ANN, GEP, GMDH and MLR.
- This literature review confirms that, there are three main challenges and questionable problems in the ML techniques developments for bedload rate including:
- 1- the input feature selection (Dehghani et al., 2019), input dimension reduction to 112 infer most effective variables (Snieder et al., 2020),
- 2- optimized subset selection of train and test data sets to avoid overfitting (Riahi-Madvar et al., 2019 & 2021), and
- 3- multi-model procedure to overcome the weakness of single models using ensembles modeling strategy (Khatibi et al., 2020).
- 4- This study aims to address these challenges and efforts to improve the estimation
   of bedload transport rate through considering techniques implemented in a multi model-based approach. As powerful ML models, MLR, ANN, GMDH and GEP
   in conjunction with SSMD, PCA and GT techniques are utilized for modeling.
- The bedload prediction challenges are improved by a successive strategy including

  Multiple Models (MM) in three levels as follows:

(i) Level 0: use pre-processing techniques, SSMD, GT and PCA in data manipulation, dimension reduction and input feature selection,

- (ii) Level 1: developing standalone ML models as base reuse and recursion techniques, that their results are reused as inputs in the next level inputs;
- (iii) Level 2: reuse and recursion of base models in a Pareto multi-gene framework by reusing the results of the previous level to the inputs of the present level and the bedload rate as a target for improved accuracy.

The main contribution of the current paper is four-fold. First, implementing the SSMD, GT, and PCA-based approaches in input vector manipulation, dimension reduction, and pre-processing of an extensive bedload transport database. Second, the utilized data set includes a wide range of low shear to high shear sediment transport observations. When combined with the pre-processing techniques, will improve the generalization issues of previous studies by dimension reduction. Third utilizing individual MLR, MLR-PCA, ANN, ANN-PCA, GEP, GEP-PCA, GMDH, GMDH-PCA models to derive explicit predictive equations for bedload. Fourth, integrating the output of individual models with the POMGGP procedure as a new multi-model strategy that utilizes individual models' power of and eliminates their weakness in bedload predictions.

To the best of the author's knowledge, the presented multi-model ensembles approach driven by the different techniques is a unique one in the literature concerning bedload rate prediction. This paper is organized as follows. Section 2 presents the material and method, including data, dimension analysis, preprocessing techniques, stand-alone, and multi-model strategy. Section 3 discusses the results of the study in three pre-defined levels. Section 4 provides summaries and conclusions.

#### 2- Material and methods

#### 2-1- Experimental data and dimensional analysis

Literature review revealed that bedload material properties, cross-section geometry features, and flow conditions are the main properties that affect the sediment transport in streams (Safari et al., 2020; Ghani ,1993;) and bedload transport in rivers can be defined by the following set of effective parameters in the form of unknown  $f_1$  function

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$$q_b = f_1(U, H, W, R, D_s, S, g, \rho_s, \rho_w, \mu, u_*, u_{*c})$$

Where  $q_b$  is bedload transport, U is flow velocity, H is flow depth, W is river width, P is hydraulic radius,  $P_s$  is sediment size, P is bed slope, P is gravity acceleration, P and P are sediments and water mass density respectively, P is dynamic viscosity,

u\* is shear velocity, and u\*<sub>c</sub> is critical shear velocity. The dimensionless form of bedload transport rate can be written in unknown  $f_2$  functional form as

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$$\emptyset = f_2(S, D_{gr}, \frac{R}{D_S}, \frac{U}{u_{*c}}, \frac{H}{D_S}, \frac{H}{W}, F_r, F_{rg}, R_e, R_{e*}, \theta, \frac{U}{u_*})$$
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in which the dimensionless parameters are particle mobility parameter  $\emptyset$ , Slope S, dimensionless grain diameter  $D_{gr}$ , relative depth  $\frac{R}{D_S}$ , critical velocity ratio  $\frac{U}{u_{*c}}$ , depth ratio  $\frac{H}{D_S}$ , aspect ratio  $\frac{H}{W}$ , Froud number  $F_r$ , densimetric Froud number  $F_{rg}$ , Reynold number  $R_e$ , densimetric Reynold number  $R_{e*}$ , shields parameter  $\theta$ , velocity ratio  $\frac{U}{u_*}$ , defined by

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$$\emptyset = \frac{q_b}{D_{es}\sqrt{g(s-1)D_e}}, D_{gr} = D_s \left[ \frac{(S-1)g}{\vartheta^2} \right]^{1/3}, R_{e*} = \frac{u_*D_s}{\vartheta}, R_e = \frac{UH}{\vartheta},$$
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$$F_r = \frac{U}{\sqrt{gH}}, F_{rg} = \frac{U}{\sqrt{gD_S(s-1)}}, \theta = \frac{\gamma HS}{gD_S(s-1)}$$

In order to develop the models, several datasets available in the literature were extracted, pre-processed, and utilized (Cao, 1997; Meyer-Peter and Müller, 1948; Recking et al., 2004). A total of 1280 data sets are used in this current study that are provided in the paper's supplementary material. The sediment diameter ranges from 0.274 mm to 44.3 mm. The bed sloped varies from 0.01 % to 20 %, flow depth from 0.00084 m to 1.0921 m, flow velocity from 0.193 m/s to 2.88 m/s, Froud number from 0.41 to 5.19 and bed material load from 0.01 g/m³ to 1356 g/m³.

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#### 2-2- Level 0: Pre-Processing techniques of bedload data

## 2-2-1-Feature selection using Gamma test

In the current study the GT is used to select the best input variables in ML-based bedload predictions. The GT stands on the hypothesis that when two points of x' and x are close together in input space, their corresponding bedload rate in output space of  $\varphi$ ' and  $\varphi$  should be close, else their difference is due to noise. In each data set of  $\{(x_i, \emptyset_i) \in \mathbb{R}^m, 1 \le i \le M\}$  by only supposition of the functional form of bedload transport  $\emptyset = \emptyset(x_1, ... x_m) + r$ , where  $\emptyset$  is a smooth function, and r is a random variable that shows noise with the bounded variance of noise Var(r). In mathematics a function could be considered "smooth" if it is differentiable everywhere (hence continuous) and in the Gamma test procedure the Ø is smooth if it has constrained first partial derivatives. For a function to be smooth, it must have continuous derivatives up to a certain order, say k. We say that function is kth order smooth. Now the domain of possible predictive model is constrained to the smooth functions  $\emptyset$  that have constrained first partial derivatives. The Gamma indicator  $\Gamma$  is an estimation of that portion of the variance of the predictions that cannot be achieved by a smooth model (Remesan et al., 2009). By calculating the Euclidean distance

 $\delta$  and  $\gamma$  of  $k^{th}$  nearest neighbour  $x_N[i,k]$  from  $x_i(1 \le i \le M)$ ,  $(1 \le k \le p)$  the  $\Gamma$  is computed from least-square fit between  $\delta$  and  $\gamma$  as:  $\gamma = A\delta + \Gamma$ . The slope of regression A represents the complexity of bedload transport phenomenon under investigation. In the GT, if the  $\Gamma$  in comparison with the variance of  $\emptyset$  as  $V_{ratio}$  were high, the probability of predicting  $\emptyset$  using selected inputs is low, when the  $V_{ratio}$  is small or near zero, the probability of predicting  $\emptyset$  by selected inputs is high. So, using the mask tests, the most effective parameters on  $\emptyset$  can be determined. Also, the GT using M-test can determine the appropriate number of data records in modelling bedload transport (Dehghani et al. ,2019). In this study the WinGamma software is sued for GT, freely available at:

http://users.cs.cf.ac.uk/O.F.Rana/Antonia.J.Jones/GammaArchive/Gamma%20Soft ware/Mathematica/GammaTestMathematicaFiles.htm

# 2-2-2-Data Clustering and Subset Selection by SSMD

According to Montes et al. (2021) and Safari (2020), the range of dissimilarity in the training dataset directly influences the model generality, overfitting problem, extrapolation ability and accuracy. The SSMD is used to avoid the overfitting of data-driven models. Suppose that X is the dataset as  $X = (x_1, x_2, ..., x_p)$  and a collection of m = 1, 2, ..., N points defined as a selected subset for the training stage. If the squared distance between  $i^{th}$  and  $j^{th}$  point define as  $D_{ij}^2$ , and k points have

already been selected  $(k \le p)$ , then the minimum distance from applicant point of N to k points define as (Memarzadeh et al., 2020)

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$$D_{ij}^2 = ||x_i - x_j||^2 = \sum_{k=1}^p (x_{ki} - x_{kj})^2$$
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The  $(k+1)^{th}$  candidate point in train group is chosen from remaining (N-k) points in the dataset that has the highest distance from an existing point. In this study, the SSMD code is developed in MATLAB environment.

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# 2-2-3-Component selection and dimension reduction using PCA

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In the PCA pre-processing technique, the original input variables are converted and reduced to fewer independent principal components (PCs) through an orthogonal projection into uncorrelated PCs (Lu et al., 2003). Using this technique, combinations of the P primary variable,  $X_1, \ldots, X_p$ , are used to create P independent components,  $PC_1, \ldots, PC_p$  equal to the number of original variables.

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# 2-3- Level 1: Standalone predictive models

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# 2-3-1-Multiple linear regression (MLR)

If we have n observations of the p-dimensional independent variable X and want to establish a linear relationship with the response variable  $\emptyset$ , we can use the following MLR model (Zounemat-Kermani et al., 2020):

 $\emptyset = \beta_0 + \beta_1 x_1 + \beta_2 x_2 + \dots + \beta_p x_p + \varepsilon$ 

The parameters  $\beta_j$ , j = 0, 1, ..., p are called regression coefficients. The least-squares method is commonly used to estimate the regression coefficients.

#### 2-3-2-ANN-MLP

The Multi-Layer Perceptron (MLP) models are the most popular NN tools used in most of research and literature (Seifi and Soroush, 2020). By determining the weights and biases of NN architecture, and simplifying the MLP, the predictive equation of model can be derived. The ANN is developed using the MATLAB toolbox.

#### 2-3-3-Pareto Optimal GEP and MGEP

The innovative technique of gene expression programming (GEP) utilized with Darwinian theory of evolution by natural selection to automatically solve optimization problems based on its two main components, the chromosomes and expression trees. A new sophisticated version of GEP is the Multigene-GEP(MGEP), that the initial population is created by GP trees with different genes (a number selected from1 and  $G_{max}$ ). In the MGEP approach two conflict goals are considered. The first is the selection of the bedload predictive equation with lowest complexity and the second is the highest accuracy. These two conflict objects lead to a multi objective optimization problem. Here to solve the optimization problem

with two conflict goals the Pareto optimality is combined with multi-genetic programming. In the multi-model-based framework the Pareto optimization is sued in order to balance between the complexity of model and the accuracy. Suppose that  $X_1$  and  $X_2$  are two feasible solutions. In the dominance relationship, two solutions must satisfy the constraints of (Zhang et al., 2017):  $f_d(X_1) \le f_d(X_2), \forall d \in$  $\{1,2,\ldots,D\}$  and  $f_i(X_1) \leq f_i(X_2), \exists i \in \{1,2,\ldots,D\}$ , In which  $f_d$  is the fitness value of d solution, and D is the number of the optimization goals. If the feasible solution X\* satisfies the above conditions and there isn't any sequence solution X while  $X < X^*$ , so that the solution  $X^*$  will be preserved and is called the Pareto optimal solution. A collection of entire Pareto optimal solutions is entitled as the final Pareto optimal solutions set, and a set of values of the target function that are related to the disassembly sequence is called the Pareto optimal frontier. The complexity of each multi-gene is calculated simply by summation of individual gene complexity. In individual genes, the complexity is determined by counting the nodes, the subtrees, leaves. A tradeoff between model accuracy and complexity would result in Pareto optimal selection of the best equation. The POMGGP is used in the MATLAB software with GPTIPS toolbox.

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#### 2-3-4- Group Method of Data Handling (GMDH)

GMDH is one of the meta-heuristic data-driven models based on multivariate 276 analysis for complex systems without the need to have a special basic knowledge. 277 The GMDH develops an analytic function using a progressive network with 278 binomial transfer functions (Shaghaghi et al., 2018). The mathematical form of 279 GMDH that maps inputs  $(x_1, x_2, x_3, ..., x_n)$  to the predicted output  $(\widehat{\emptyset})$  is written as 280  $\widehat{\emptyset} = a_o + \sum_{i=1}^n a_i x_i + \sum_{i=1}^n \sum_{j=1}^n a_{ij} x_i x_j + \sum_{i=1}^n \sum_{j=1}^n \sum_{k=1}^n x_j a_{ijk} x_i x_j x_k + \cdots$ 281 The least-squares error rule is utilized for coefficient determination of GMDH in 282 MATLAB environment. 283

## 2-4- Level 2: Multi-model ensembles (MME) approach

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In the present study, in addition to the individual predictive models, an innovative multi-model ensembles approach is presented. This feeds the output of standalone 286 models into the POMGGP as a multi-model technique to improve the predictive capability of models. 288 This new contribution in bedload rate prediction as an ensembles approach consists of two primary levels: Level 1 in which the original input variables or pre-processed 290 (PCs) are used to estimate bedload transport rate in standalone models of MLR, 291 MLR-PCA, ANN, ANN-PCA, GMDH, GMDH-PCA, GEP, GEP-PCA; Level 2 in 292 which the outputs of level 1 models are used as inputs to the POMGGP along with 293 the original bedload rate as output. 294

In this framework, as presented in Fig.1 the POMGGP is used to run models at level 1 based on Pareto optimality analysis. Observed values of bedload rate serve as the target output in both levels. The strength of the developed framework is learning at two levels, automatic individual model selection by natural evolution in multi-gene GEP, balancing surrogate model complexity and accuracy via Pareto optimality.

The idea behind the multi-model ensembles approach has been inspired by the hierarchical recursion of models, that teamworking of models in parallel can help achieve a more accurate prediction (Khatibi et al., 2020).

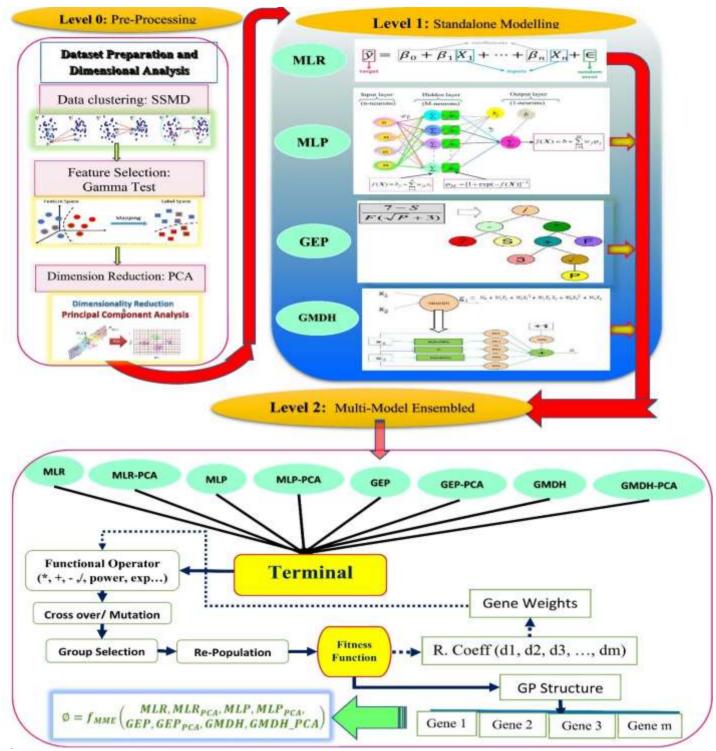
The models in level 1 and 2 are comparatively evaluated using performance metrics coefficient of determination (R<sup>2</sup>), root mean square error (RMSE), mean absolute error (MAE), Nash Sutcliffe efficiency (NSE), and graphical analysis including scatter plots, importance probability, Pareto front and Taylor diagrams.

Furthermore, a newly revised discrepancy ratio (RDR) for error distributions developed by the authors (Riahi et al., 2020) is used to overcome non-normality, zero or negative value predictions with a rectified linear unit (ReLu) function (Ramachandran et al., 2018). The RDR is calculated by:

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$$RDR = Sign(\emptyset_{p,i} - \emptyset_{o,i}) \left| log \left| \frac{\emptyset_{p,i}}{\emptyset_{o,i}} \right| \right|$$
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In which the  $\emptyset_{o,i}$  is measured value  $\emptyset_{p,i}$  is the estimated model output. In the case of over-predictions by POMGGP, the value of RDR>0 and in the case of under-

prediction RDR<0 and for exact predictions RDR is equal to zero. The multimodel ensemble is developed using MATLAB environment.



**Fig. 1.** Flowchart of the developed multi-model ensembles approach for function finding in bedload prediction

#### 3- Results and discussion

#### 3-1- Level 0: Pre-processing

The results obtained using the pre-processing techniques are presented in Table 1. The train (80%) and test (20%) sets are selected using the SSMD approach. For a sizeable natural data bank like those used in this study, the SSMD expandes the envelope range of training sets, improves the applicability of developed predictive models and encompasses outlier data in the training set.

Table 1. Descriptive statistics of parameters in all, train and test subsets categorized by SSMD.

First Third

						First		Third	
	Parameter	Mean	Mode	SD	Min	quartile	Median	quartile	Max
	S	0.02	0.01	0.04	0.00	0.00	0.01	0.02	0.20
	Dgr	163.08	12.88	210.77	6.98	35.97	80.70	221.92	1150.14
A 11	U/u*c	11.73	11.21	3.38	3.26	9.86	12.12	14.37	18.99
All	H/W	0.21	0.13	0.14	0.01	0.09	0.18	0.28	0.85
(1280	Fr	1.19	1.13	0.60	0.41	0.79	1.09	1.35	5.19
data	Re*	944.92	28.00	1834.21	21.00	55.00	199.50	1103.00	15086.00
points)	θ	0.20	0.05	0.30	0.01	0.05	0.07	0.29	3.70
	U/u*c	11.69	11.87	3.43	3.26	9.86	12.13	14.37	18.98
	Φ	1.84	0.00	11.59	0.00	0.00	0.02	0.67	264.05
	S	0.03	0.07	0.04	0.00	0.00	0.01	0.02	0.20
	Dgr	192.86	12.88	223.84	6.98	51.78	107.35	262.17	1150.14
	U/u*c	11.08	11.21	3.36	3.26	9.00	11.25	13.42	18.99
Train	H/W	0.22	0.04	0.15	0.01	0.10	0.20	0.30	0.85
(1024	Fr	1.21	1.13	0.64	0.41	0.83	1.08	1.33	5.19
data points)	Re*	1151.67	30.00	1992.09	21.00	98.00	362.50	1265.00	15086.00
points)	θ	0.19	0.05	0.32	0.01	0.05	0.07	0.16	3.70
	U/u*c	11.04	3.26	3.42	3.26	9.00	11.24	13.42	18.98
	Φ	1.95	0.00	12.93	0.00	0.00	0.01	0.14	264.05
	S	0.01	0.00	0.00	0.00	0.00	0.01	0.01	0.02
	Dgr	43.95	12.88	63.24	12.86	12.88	12.88	51.78	608.49
Tr. 4	U/u*c	14.31	11.47	1.89	7.19	13.46	14.55	15.66	17.68
Test	H/W	0.15	0.07	0.11	0.01	0.08	0.12	0.18	0.58
(256 data	Fr	1.12	0.69	0.41	0.45	0.72	1.13	1.44	2.16
	Re*	117.91	28.00	309.44	25.00	29.00	35.00	81.00	3528.00
points)	θ	0.25	0.06	0.21	0.02	0.05	0.30	0.43	0.87
	U/u*c	14.31	14.24	1.89	7.20	13.46	14.55	15.66	17.68
	Φ	1.42	0.00	1.79	0.00	0.00	0.94	2.32	9.07
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The GT is used for feature selection and determining the proper input vector that characterizes the complex process in bedload transport. At first, the datasets are normalized [-1 1] and then GT is utilized via mask test procedure, and GT results for different input configurations are shown in Table 2. In Table 2, 12 dimensionless variables are used as the input variables with varying combinations to the GT.

In the first configuration, all 12 input parameters are used and GT indices calculated as given in the first row of Table 2. Then in the next GT run, the first input parameter is removed and masked and the GT results are recalculated, as given in the second row. Again, the removed variable is returned into the input vector and the second input variable is masked, and GT is performed in all the combinations. This method is continued for all selected variables in Table 2, one by one and in each step the  $\Gamma$  value is calculated.

The masking of the most influential variables in bedload prediction is associated with increases in the  $\Gamma$  value (V ratio) regarding the case that includes all variables (first row in Table 2). The highest  $\Gamma$  value indicates that the removed variable is essential and should be selected as the input variable of models.

Finally based on the results of GT in Table 2, the most important variables with the highest  $\Gamma$  value are S,  $D_{gr}$ ,  $U/u*_c$ , H/W,  $F_r$ ,  $Re*_$ , Q,  $U/u*_$  as shown in bold style. The input components reduced from 12 to 8 and the functional form simplified as

Table 2. The GT results on the selected 12 input masks for feature selection

	Removed	Gamma	Gradient	Standard Error	V-Ratio	Mask
0	None	0.04134632	0.24293793	0.02504585	0.165385281	111111111111
1	$\mathbf{S}$	0.043221076	0.25201855	0.02499108	0.172884302	011111111111
2	Dgr	0.041702514	0.246745969	0.025434014	0.166810055	101111111111
3	$R/D_s$	0.041275668	0.249138526	0.025293184	0.165102672	110111111111
4	U/u*c	0.042657031	0.251330176	0.025272607	0.170628125	111011111111
5	H/ D <sub>s</sub>	0.041493969	0.260073122	0.025051376	0.165975875	111101111111
6	H/W	0.044526813	0.273403925	0.025769505	0.178107253	111110111111
7	Fr	0.043472253	0.258844691	0.024823123	0.173889012	111111011111
8	Frg	0.040546081	0.271921358	0.025270117	0.162184324	111111101111
9	Re	0.040604003	0.258535669	0.024947276	0.162416013	111111110111
10	Re*	0.041900604	0.255306066	0.025551793	0.167602418	111111111011
11	θ	0.042757646	0.337648978	0.024902663	0.171030583	111111111101
12	U/u*	0.042685003	0.251320735	0.025282989	0.170740012	111111111110

$$\emptyset = f_3(S, D_{gr}, \frac{U}{u_{*c}}, \frac{H}{W}, F_r, R_{e*}, \theta, \frac{U}{u_{*}})$$

The PCA is used as a dimension reduction technique over the GT results. According to KMO=0.624, the PCA is applicable for dimension reduction and the input variables are reduced into three principal components which are a linear combination of primitive dimensionless variables as

355 
$$PC_{1} = 0.069S_{n} + 0.310D_{gr,n} - 0.256\left(\frac{U}{u_{*c}}\right)_{n} + 0.012\left(\frac{H}{W}\right)_{n} - 0.042(F_{r})_{n}$$

$$+ 0.293(R_{e*})_{n} - 0.151(\theta)_{n} - 0.253\left(\frac{U}{u_{*}}\right)_{n}$$
357 
$$PC_{2} = 0.314S_{n} - 0.143D_{gr,n} - 0.051\left(\frac{U}{u_{*c}}\right)_{n} + 0.064\left(\frac{H}{W}\right)_{n} + 0.39(F_{r})_{n} - 0.081(R_{e*})_{n} + 0.382(\theta)_{n} - 0.059\left(\frac{U}{u_{*}}\right)_{n}$$
358 
$$PC_{3} = -0.113S_{n} + 0.289D_{gr,n} + 0.267\left(\frac{U}{u_{*c}}\right)_{n} + 0.683\left(\frac{H}{W}\right)_{n} + 0.037(F_{r})_{n}$$
360 
$$+ 0.289(R_{e*})_{n} + 0.266(\theta)_{n} + 0.264\left(\frac{VU}{u_{*}}\right)$$

Here, the n footnote indicates the normalized parameters in PCA. These three PCs explained the 85 % of total variances in the bedload transport datasets. The PCA results are given in Table 3, and the Kaiser criterion shows that three components have eigenvalues of more than 1 with a cumulative total variance of 85 %.

Therefore, the 8 bedload transport parameters can be reduced to the three uncorrelated PCs while preserving 85 % of the information of primary variables. As this table shows, the PC1 has an eigenvalue of 3.526 that explains 44.071 % of the total variance, PC2 has an eigenvalue of 2.049 with 25.614 % of total variance presented and PC with an eigenvalue of 1.16 has an eigenvalue of 14.505 %.

A scree graph of the amount of variance explained versus PCs and eigenvalues is shown in Fig.2, indicates that a break of the line occurred after PC3 and shows that only first three PCs maintain useful information. The selected PCs are rotated to determine their importance relative to each of 8 dimensionless parameters, as given in Table 4. A high value for each parameter's PC loading indicates a reasonable correlation between the parameter and corresponding PC.

Table 3. The PCA results on bedload transport data

Component	Eigenvalues						
Component	Total	% of Variance	Cumulative %				
1	3.526	44.071	44.071				
2	2.049	25.614	69.685				
3	1.160	14.505	84.190				
4	0.800	10.006	94.196				
5	0.297	3.712	97.908				
6	0.129	1.610	99.518				
7	0.037	0.462	99.980				
8	0.002	0.020	100.000				

Table 4. Rotated PC loading of bedload effective parameters

D	Component						
Parameter	1	2	3				
S	0.423	0.825	-0.159				
$D_{gr}$	0.865	-0.171	0.345				
$\frac{U}{u_{*c}}$	-0.834	-0.317	0.320				
$\frac{H}{W}$	0.068	0.106	0.791				
$F_r$	0.126	0.925	0.011				
$R_{e*}$	0.856	-0.028	0.339				
$\theta$	-0.222	0.815	0.281				
$\frac{U}{u_*}$	-0.830	-0.334	0.317				

As these results show, the first component is explained by  $D_{gr}$  and  $R_{e*}$  and includes the highest level of information and describes the sediment material properties. The second PC is explained by S, ,  $F_r$  and  $\theta$  that describes the flow properties, and the third PC is explained by  $\frac{U}{u_{*c}}$ ,  $\frac{H}{W}$ , and  $\frac{U}{u_*}$ , which this PC describes the geometry and

friction properties of the bedload transport. These three relevant PCs will be used as an input vector to the multi-models as follows

$$\emptyset = f_4(PC_1, PC_2, PC_3)$$

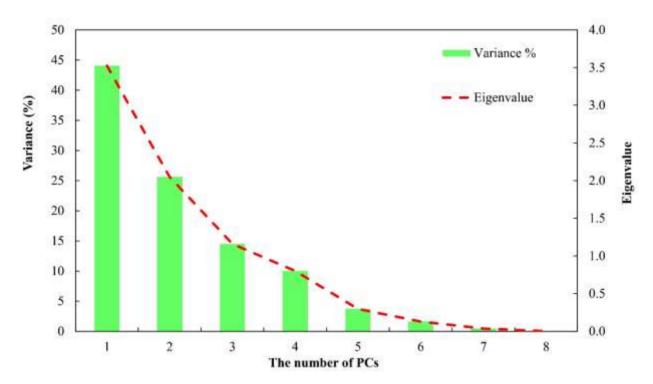


Fig. 2. Scree plot showing the variance of all components

#### 3-2- Level 1: Performance of standalone models

The results obtained by using the presented standalone models are presented and discussed here. A comprehensive evaluation of the model predictions should include at least 'goodness-of-fit' such as R<sup>2</sup>, NSE and error indices such as RMSE, or RAE. The comprehensive comparison of the best single model results using the selected input variables by the GT and the quantitative values of performance evaluation

- 400 indices of the MLR, MLR-PCA, ANN, ANN-PCA, GEP, GEP-PCA, GMDH,
- 401 GMDH-PCA are presented in Table 5.
- In the training step, the ensembles ANN-PCA model showed a relatively accurate
- estimation of bedload with ( $R^2=1\approx0.996$ ), RMSE=0.71 when compared with the
- 404 ANN ( $R^2$ =0.98, RMSE=1.66), GEP-PCA ( $R^2$ =0.95, RMSE=2.94) and the others.
- Based on the classification of model performances by the R<sup>2</sup> metric, all models in
- Table 5 had an outstanding performance  $(0.7>R^2>1)$  in bedload predictions, except
- 407 the MLR-PCA. In the test stag, the same performance trend and accuracy
- improvement when combined the standalone models with the PCA were declared.
- The best results were comparatively obtained by the ANN-PCA, GEP-PCA and
- 410 ANN models. In this regard the ANN-PCA model with R<sup>2</sup>=0.96, RMSE=0.38,
- RAE=0.16 and NSE=0.95 have the best predictions for the bedload in the test stage.
- The NSE values of the GEP-PCA, ANN-PCA, GEP and ANN models in the train
- and testing steps confirmed excellent predictions for the bedload transport in the test
- stage with NSE>0.75. The best accuracy of the GEP-PCA and ANN PCA-based
- models confirmed their ability in the emerging non-linear system indentation when
- combined GT and PCA's pre-processing model-free techniques.
- The hierarchical accuracy of models follows the order of ANN-PCA> ANN> GEP-
- PCA> GEP> GMDH-PCA> GMDH> MLR-PCA> MLR in terms of the R<sup>2</sup>, RMSE,

- RAE and NSE values for the test stage, as given in Table 5. The percent of prediction
- improvements by utilizing the PCA as input dimension reduction in RMSE reduction
- was 57% and 3% in ANN-PCA, 4% and 4% in GEP-PCA, 9% and 45% in GMDH-
- PCA for train and testing steps, respectively. The explicit form of predictive
- equations based on the trained above eight models are as follows:
- 424 MLR:

425 
$$MLR = -3.39S + 0.02D_{gr} - 2.02\frac{U}{u_{*c}} - 5.4\frac{H}{W} - 8.3F_{r} - 0.002R_{e*} - 46.2\theta + 1.8\frac{U}{u_{*}} + 4.81$$
 11

426 MLR-PCA:

$$427 MLR - PCA = 1.95 - 2.81PC_1 + 6.35PC_2 + 3.65PC_3 12$$

428 GMDH:

429 
$$G_1 = 1.9Se^{0.27(\theta+0.64)} + 0.16S^2 + 12e^{0.27(\theta+0.64)} - 2.84S - 11.44$$

430 
$$G_2 = 411.52 * e^{0.002(G_1 + 1.73)} - 411.4$$

431 
$$G_3 = 115671.3e^{0.000009(G_2+1.35)} - 115671.3$$

432 
$$GMDH = 970.9e^{0.000956(G_3+1.36)} - 970.79$$

433 GMDH-PCA:

434 
$$GMDH - PCA = \frac{504.35}{1 + e^{4.55PC_3 - 17.7}} + 1.78e^{0.998 + 0.3008PC_3 + 0.55PC_2 - 0.225PC_1} +$$
435 
$$0.01e^{-(3.71PC_1 + 10.83)} - 0.61PC_3 + 0.44PC_3 \times PC_1 - 506.39$$

436 GEP:

437 
$$GEP = \theta \times Fr + e^{Fr-2.21} + 6.2e^{\theta} - \theta - S - 7.14$$
 15

438 GEP-PCA:

439 
$$GEP - PCA = 33.6e^{-\frac{(PC_2 - 7.81)^2}{2PC_3^2}} + 1479391.4e^{-\frac{(PC_2 - 24.13)^2}{2PC_3^2}} - 1.75e^{-0.37(PC_2 - PC_1)^2} + 440 \qquad 2^{(PC_2 - PC_1)} + 0.73$$

441 ANN:

442 
$$ANN = \frac{530.2}{1+e^{-2(T_1+T_2+T_3+T_4)-28.3}} - 265.1$$

443 
$$T_{1} = \frac{16.36}{1 + e^{0.66S - 12.94D_{gr} + 5.18\frac{U}{u_{*c}} + 0.2\frac{H}{W} + 5.44F_{r} + 7.66R_{e*} - 4.34\theta - 5.18\frac{U}{u_{*}} - 5.86} - 8.18$$

444 
$$T_{2} = \frac{-41.46}{1 + e^{31.22S + 1.74D_{gr} + 109.78\frac{U}{u_{*c}} - 165.86\frac{H}{W} + 136.76F_{r} + 2.3R_{e*} + 92.44\theta + 117.63\frac{U}{u_{*}} - 271.4}} + 20.73$$

446 
$$T_{3} = \frac{-2.98}{1 + e^{1.26S - 12.64D_{gr} + 4.8\frac{U}{u_{*}c} + 0.1\frac{H}{W} + 5.28F_{r} + 7.44R_{e*} - 2.68\theta - 4.72\frac{U}{u_{*}} + 4.2} + 1.49$$

$$T_4 = \frac{0.24}{1 + e^{26S - 4.78D_{gr} - 2.52\frac{U}{u_{*c}} + 9.24\frac{H}{W} + 6.06F_r - 0.3R_{e*} - 38.42\theta - 0.38\frac{U}{u_*} + 9.38} - 0.12$$

448 ANN-PCA:

449 
$$ANN - PCA = \frac{530.2}{1 + e^{-2(T_1 + T_2 + T_3) - 68.43}} - 265.1$$

450 
$$T_1 = \frac{-5.065}{1 + e^{-37PC_1 + 34.63PC_2 - 2PC_3 - 30.2}} + \frac{0.09}{1 + e^{20.2PC_1 - 16.9PC_2 + 2.75PC_3 + 9.68}} + 2.48$$

451 
$$T_2 = \frac{68.9}{1 + e^{1025.7PC_1 + 34.7PC_2 + 195.9PC_3 + 347.4}} + \frac{5.64}{1 + e^{-36.4PC_1 + 18.4PC_2 - 8.6PC_3 - 18.6}}$$
452 
$$-37.3$$

453 
$$T_3 = \frac{-0.03}{1 + e^{-9.75PC_1 + 17.1PC_2 + 0.3PC_3 + 3.6}} + 0.013$$

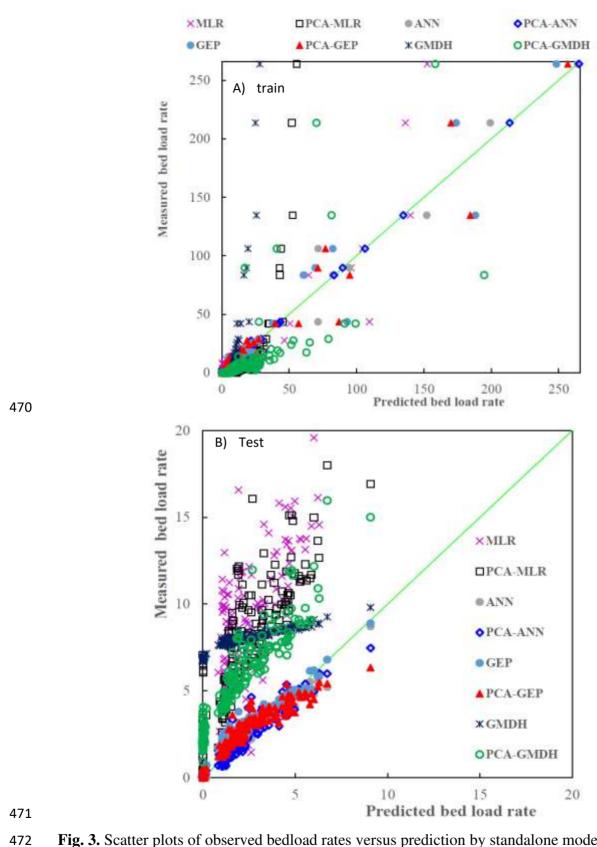
The scatter plots of the measured bedload rate against predicted by the models are presented in Fig.3. This figure shows that MLR, MLR-PCA, GMDH and GMDH-PCA model have underestimations for the bedload rate. As the results in this figure confirmed, the GEP-PCA, ANN-PCA models are most consistent with the 1:1 line and provide superior predictions for bedload transport rate in rivers compared to the standalone models of ANN, GEP, MLR, and GMDH.

As the first main motivation and contribution of the current study was to introduce the feasibility of utilizing the pre-processing model-free techniques of SSMD, GT and PCA and their ensemble ability with standalone models for bedload transport rate prediction in rivers, these techniques show an improved generalization capacity than non-preprocessed predictions and is confirmed with high estimation accuracy obtained.

**Table 5.** The statistical measures of standalone models in the train and testing steps

	Train								
		MLR-	ANN- GEP-			GEP-	GMDH-		
	MLR	PCA	ANN	PCA	GEP	PCA	GMDH	PCA	MME
$\mathbb{R}^2$	0.76	0.37	0.98	0.996	0.94	0.95	0.80	0.55	0.997
<b>RMSE</b>	6.27	10.27	1.66	0.71	3.06	2.94	12.77	9.39	0.6
RAE	0.90	1.39	0.10	0.08	0.24	0.20	2.28	1.26	0.06
NSE	0.76	0.37	0.98	0.996	0.94	0.95	0.02	0.47	0.997

						Test			
		MLR-		ANN-		GEP-		GMDH-	
	MLR	PCA	ANN	PCA	GEP	PCA	GMDH	PCA	MME
$\mathbb{R}^2$	0.78	0.76	0.96	0.96	0.93	0.92	0.88	0.89	0.98
<b>RMSE</b>	5.66	4.75	0.37	0.36	0.56	0.54	6.16	3.37	0.24
RAE	3.06	2.83	0.18	0.16	0.34	0.25	4.21	2.16	0.1
NSE	-9.03	-6.07	0.96	0.95	0.90	0.91	-10.91	-2.57	0.98



**Fig. 3.** Scatter plots of observed bedload rates versus prediction by standalone models in A) tarin and B) test data sets

492

3-3- Level 2: Performance of EMM approach: Ensembles-POMGGP 475 In the developed new strategy of EMM approach for bedload transport predictions, 476 the Pareto optimality in conjunction with the multi-gene genetic programming is 477 used to predict bedload transport by considering the output of standalone models. In 478 this strategy the MLR, MLR-PCA, ANN, ANN-PCA, GEP, GEP-PCA, GMDH and 479 GMDH-PCA predictions are used as the input vector to the POMGGP model and 480 481 the feasible inputs are selected automatically by the geniting programming. The Multi-Model input variable importance is shown in Fig. 4. As this figure shows, 482 483 the most important sub-model is the ANN with an importance probability of 0.301, followed by the ANN-PCA sub-model with an importance probability of 0.286, and 484 the MLR model with an importance probability of 0.225. Less important sub-models 485 in the ensembles multi-model for predicting the dimensionless bedload transport rate 486 follows the order of GMDH (probability=0.075)> GEP (probability=0.071)> 487 GMDH-PCA (probability=0.029)> GEP-PCA (probability=0.014)> and MLR-PCA 488 489 (probability=0.0). The importance probability graph of sub-models in Fig. 4 shows that using the 490 results of ANN, ANN-PCA, MLR, GMDH and GEP models, we are able to derive 491 a predictive equation with an importance probability of 95.8%. So, in order to reduce

the complexity of the final multi-model, and increase the application feasibility of the results, the Pareto optimality is used to derive the equation of final multi-model.



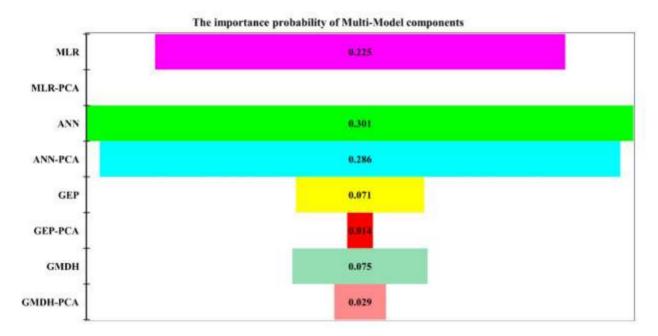


Fig. 4. The Multi-Model input variable (standalone models) importance

The parameters in Table 6 are determined by trial and error and using those suggested in the literature. The multi-gene genetic programming is trained and optimized by the least square error, the RMSE as the fitness function, basic math operators in function set, and Pareto optimality as the selection criteria. The Pareto graph of the evolved multi-models for bedload predictions using all sub-models as inputs, i.e: MLR, MLR-PCA, ANN, ANN-PCA, GEP, GEP-PCA, GMDH, and GMDH-PCA are shown in Fig. 5.

The Pareto-optimal solution of different multi-models on the Pareto front are chosen not more than 10% decrease occurred in model accuracy neither in the train nor at testing step. In this figure, the Pareto front is demonstrated with green circles, and the best final multi-model as the optimal solution is displayed by a circle with red perimeter and green color filled. The structural properties of the final multi-model include the overall complexity of 367, with 89 nodes in the selected symbolic expression, 4 individual genes, depth value 6 and -7.77 as the bias term, with MLR, ANN, ANN-PCA, GEP-PCA, GMDH as selected optimum input sub-models in agreement with probability importance graph in Fig. 4.



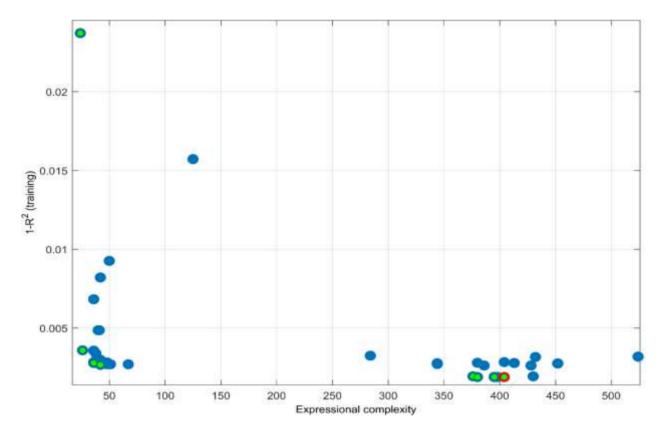


Fig. 5. Pareto graph of the best evolved multi-models

The final parse tree of the Pareto selected multi-model is presented in Fig. 6. This figure presents the symbolic expression of each gene in the multi-gene model. The corresponding equation and simplified expression of each gene, the individual gene weights, the number of nodes, individual complexities and depths are presented in Table 7.

As shown in Table 7, The bias term with weight=-7.77, Gene 2 that includes ANN and ANN-PCA with weight=7.6, Gene 4 with weight= -3.86 added the MLR, GMDH followed by Gene 1, has the highest weight and importance in the multigene model solution. To evaluate the statistical significance of individual genes the p-value of each gene calculated and the p-values in all of the genes were smaller than 0.00001, confirms and indicates the statistical importance of individual genes in the multigene model. Finally, by applying the coefficients of individual genes and simplifying the final Pareto solution of multi-gene expression, the final explicit predictive equation for dimensionless bedload rate based on the developed multi-model strategy with its effective sub-models is derived as

533 
$$\emptyset = 1.13\text{ANN}_{PCA} - 0.079\text{ANN} - 0.073\text{GEP}_{PCA} + 0.027\text{MLR} +$$
534 
$$7.6e^{\left(Exp\left(-3.34Exp\left(ANN_{PCA}^{2}\right)\right)\right)} + \frac{3.93\text{ANN}}{\text{GMDH}} - \frac{4\text{ANN}_{PCA}}{\text{GMDH}} + \frac{0.073\text{MLR}}{\text{GMDH}} -$$
535 
$$\frac{ANN_{PCA}}{13.74\text{GMDH} - 3.82\text{ANN}_{PCA} + \frac{13.74\text{MLR}}{\text{ANN}}} - 7.77$$
19

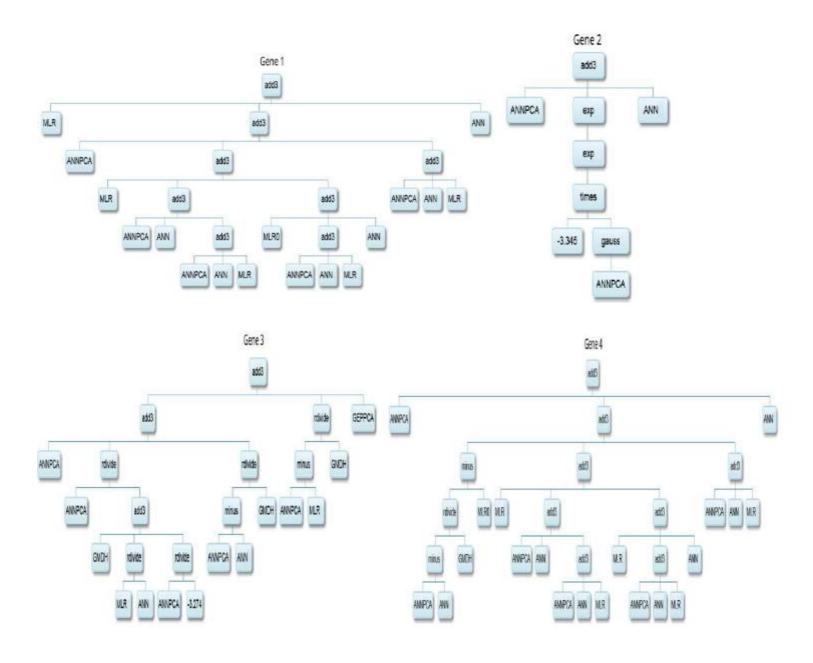


Fig. 6. The final parse tree of the Pareto selected multi-model

Performance indices of the final MME predictions for bedload are compared in Table 5 in train and test stages. Graphically, the results of Eq. 34 as the final multimodel solution are presented in Figs. 7 and 8 for the test stage and in Figs. 9 and 10

for the train stage. The scatter plots and series plots show the multi-model is 542 accurately capable of capturing low and high values of bedload with different 543 conditions in input observations. This is one remarkable aspect of our multi-model 544 in mimicking low and high flows. 545 These results revealed that the multi-model approach improved the generalization 546 capacity of single standalone single models, as confirmed with better estimation 547 accuracy obtained in this extensive dataset (Train: R<sup>2</sup>=0.997, RMSE=0.6, 548 RAE=0.06, NSE=0.997, and in test Train: R<sup>2</sup>=0.98, RMSE=0.1, RAE=0.24, 549 NSE=0.98. Comparing the results of the training period of multi-model with the 550 greatest improvement, about 16% in RMSE, and 25% in RAE was obtained 551 compared to the best standalone model, ANN-PCA. 552 Based on the results in training step, the Multi-model had a decrease of 90% (in 553 RMSE, RAE) and an increase of 31% (in NSE, R<sup>2</sup>) compared to MLR. Multi-Model 554 also showed a decrease of 95% (in RMSE, RAE) and an increase of 169% (in NSE, 555 556 R<sup>2</sup>) compared to MLR-PCA; a decrease of 63% and 40%% (in RMSE, RAE) and an increase of 2% (in NSE, R<sup>2</sup>) compared to ANN, a decrease of 80% and 75%% (in 557 RMSE, RAE) and an increase of 6% (in NSE, R<sup>2</sup>) compared to GEP; a decrease of 558 79% and 70% (in RMSE, RAE) and an increase of 5% (in NSE, R<sup>2</sup>) compared to 559 GEP-PCA in train stages.

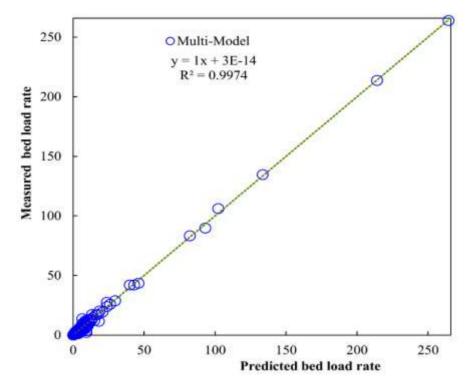


Fig. 7. Scatter plot of multi-model in training step

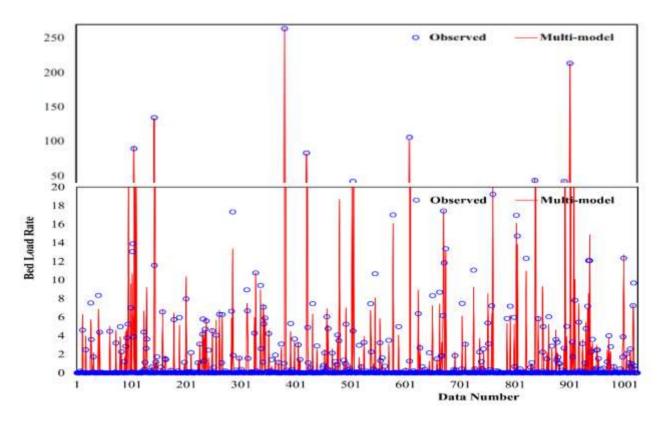


Fig. 8. Comparison of observed versus predicted bedload transport in training step

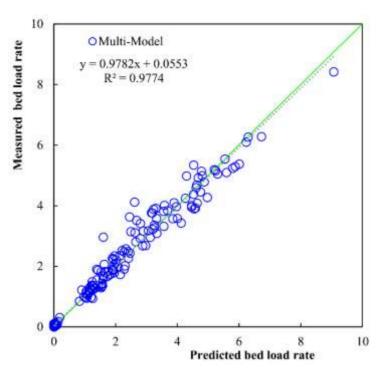


Fig. 9. Scatter plot of multi-model in testing step

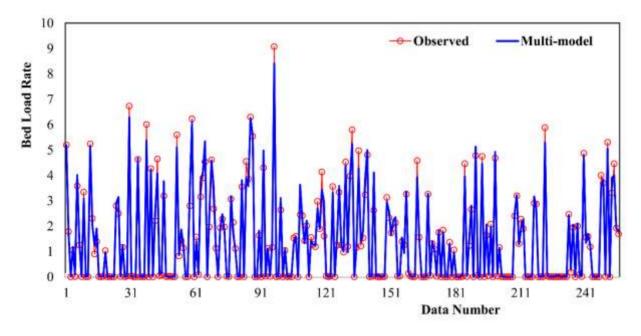


Fig. 10. Comparison of observed versus predicted bedload transport in testing step

The percentage of improvements in the test stage of multi-Model when these results are compared with other standalone predictions, are presented in Table 8. The improvement percentages in this table indicates that utilizing the multi-model strategy the RMSE and RAE values, as major error indices are decreased from 33% in ANN-PCA up to 96% in MLR and GMDH. In the R<sup>2</sup> measure the improvement varies from 2% up to 29%, and in the NSE, the gain varies from 2 up to 138%. These values confirm the superiority of the developed strategy in the generalization of bedload prediction.

models, in Figs. 11 and 12 the standardized error distribution of prediction in terms of RDR versus probability and the Taylor diagram of all models in train and test stages are shown. As these figures show the MLR, MLR-PCA, ANN and GEP models have underestimated and the GMDH-PCA, GMDH, ANN-PCA and GEP-PCA models overestimated for the bedload, while the multi-Model have reasonable estimates in training step. In the testing step, the RDR graph in Fig. 10, declares that the multi-Model strategy provides more generalities in the predictions and the RDR distribution is accurately around the 0, while the ANN, GEP, MLR and MLR-PCA have considerable underestimates and ANN-PCA, GEP-PCA, GMDH-PCA and GMDH have high overestimate in bedload. Reasonable accuracy and generality and parsimonious structure, endorse the developed multi-model approach for bedload

transport estimation in practice. The leading cause behind the improvement in MME originates from the inherent multi-process nature and different patterns of bedload transport in the extremely low flows up to high flows is that the sediment transport is a mixture of a laminar, turbulent, linear and nonlinear phenomenon in rivers that would be taken into account by integration of linear and nonlinear models.

**Table 6.** Parameter setting for the MME development.

Run parameter	Value	Run parameter	Value			
Population size	100	Gaussian perturbation of constant	0.05			
Max. generations	500	Max. genes	4			
Generations elapsed	500	Max. tree depth	6			
Input variables	8	Max. total nodes	Inf			
Training instances	1024	ERC probability	0.3			
Tournament size	50	Crossover probability	0.84			
High level Crossover	0.2	Low level Crossover	0.8			
Elite fraction	0.75	Mutation probabilities	0.14			
Sub-tree mutation	0.9	Input Mutation probabilities	0.05			
Lexicographic selection pressure	On	Complexity measure	Expressional			
Function set $*, -, +, /, ^{\wedge}, \sqrt{}$ , exp, ln, multi3, cub, gauss, add3, square,						

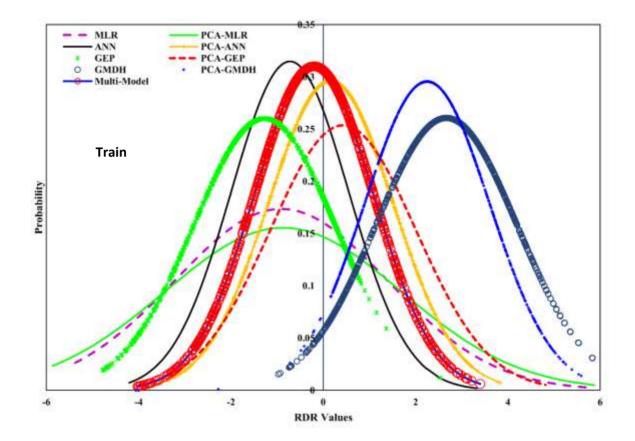
**Table 7.** The Multi-gene results of Pareto solution in MME.

Term	Value	Gene weights	Nodes	Depth	Complexity	
Bias	-7.77	-7.71	-	-	-	
Gene 1	15.4 ANN + 12.9 ANN <sub>PCA</sub> + 15.4 MLR	2.57	34	6	151	
Gene 2	$7.6 \text{ ANN} + 7.6 \text{ ANN}_{PCA} + 7.6 \text{ Exp}(\text{Exp}(-3.34 \text{ gauss}(\text{ANN}_{PCA})))$	7.6	30	6	125	
Gene 3	(0.0728 ANN)/GMDH - 0.0728 GEP <sub>PCA</sub> - ( ANN <sub>PCA</sub> )/(13.74 GMDH - 4.2 ANN <sub>PCA</sub> + (13.74 MLR)/ANN) - 0.0728 ANN <sub>PCA</sub> - (0.146 ANN <sub>PCA</sub> )/GMDH + (0.0728 MLR)/GMDH	-0.0728	9	6	31	

Term	Value	Gene weights	Nodes	Depth	Complexity
Gene 4	-(3.86 (ANN <sub>PCA</sub> -ANN + 6.0 ANN $\times$ GMDH + 5.0 ANN <sub>PCA</sub> $\times$ GMDH + 4.0GMDH $\times$ MLR))/GMDH	-3.86	23	6	97

**Overall Structure of Multi-Model:** Genes:4; Nodes:89; Complexity: 367; Depth:6; Inputs selected: MLR, ANN, ANN-PCA, GEP-PCA, GMDH





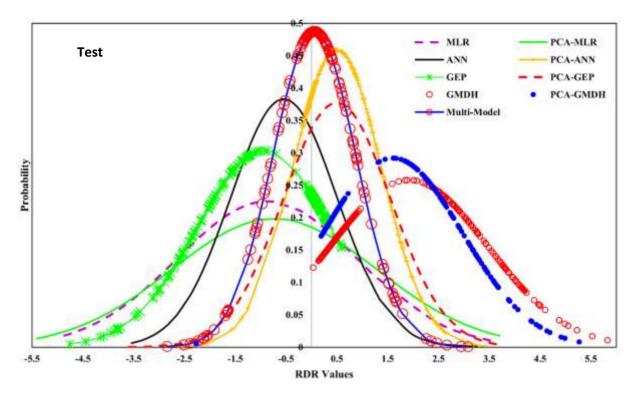
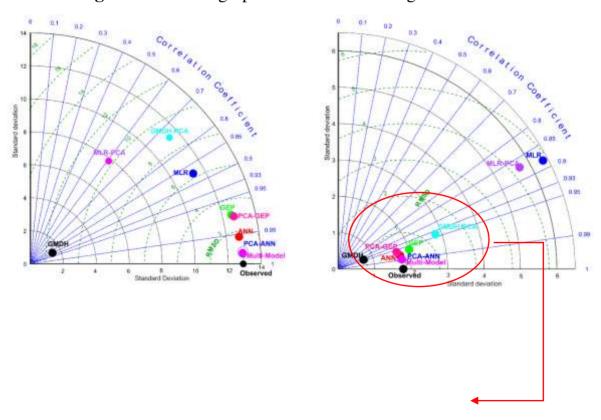


Fig. 11. The RDR graph in train and test stages of the MME



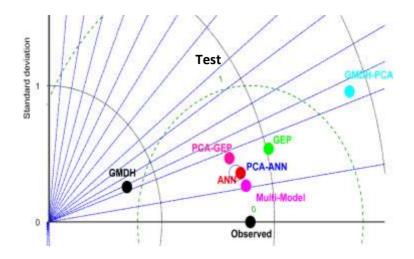


Fig. 12. The Taylor diagram in train and test stages

**Table 8.** The percentage of improvements in bedload rate prediction with multi-Model strategy in testing step

	MLR	MLR- PCA	ANN	ANN- PCA	GEP	GEP- PCA	GMDH	GMDH- PCA
R <sup>2</sup>	26	29	2	2	5	7	11	10
RMSE	-96	-95	-35	-33	-57	-56	-96	-93
RAE	-97	-96	-44	-38	-71	-60	-98	-95
NSE	111	116	2	3	9	8	109	138

## 4- Conclusions

In this study, a new multi-Model strategy integrated with pre-processing techniques of SSMD, GT, and PCA is developed to derive an explicit predictive equation for the bedload transport in rivers with extensive dataset. The framework of enhanced multi-modelling improves the accuracy and heuristic capability to learn tendencies within residuals of individual model results and gain an insight into the nature of bedload transport in three-level strategy.

- At level 0, the pre-processing, input selection and dimension reduction are carried out by SSMD, GT, PCA. At level 1, the standalone models of MLR, MLR-PCA, ANN, ANN-PCA, GEP, GEP-PCA, GMDH, GMDH-PCA are compared to derive explicit predictive equations. At level 2, the EMM is developed by utilizing the output of individual models as an external input to the multigene genetic programming with Pareto optimality. The main conclusions of this ensemble modeling are as follows:
- 1- The hierarchical accuracy of models follows the order of ANN-PCA> ANN>

  GEP-PCA> GEP> GMDH-PCA> GMDH> MLR-PCA> MLR in terms of the

  R<sup>2</sup>, RMSE, RAE and NSE values for the test stage.

- 2- The percent of prediction improvements by utilizing the PCA as input dimension reduction in terms of RMSE reduction was 57% and 3% in ANN-PCA, 4% and 4% in GEP-PCA, 9% and 45% in GMDH-PCA for training and testing steps respectively.
- 3- The MME had a decrease of 90% (in RMSE, RAE) and an increase of 31% (in NSE, R<sup>2</sup>) compared to MLR, a reduction of 95% (in RMSE, RAE) and an increase of 169% (in NSE, R<sup>2</sup>) compared to MLR-PCA; a decrease of 63% and 40%% (in RMSE, RAE) and an rise of 2% (in NSE, R<sup>2</sup>) compared to ANN, a decrease of 80% and 75%% (in RMSE, RAE) and an increase of 6%

(in NSE, R<sup>2</sup>) compared to GEP; a reduction of 79% and 70% (in RMSE, RAE) and an increase of 5% (in NSE, R<sup>2</sup>) compared to GEP-PCA.

4- The explicit predictive equation based on EMM approach has resulted in the gaining of a robust system with significant predictive accuracy improvement, (i.e., 33–96% in terms of RMSE; 2-29% in terms of R<sup>2</sup>, 2-138% in terms of NSE and 38-98% in terms of RAE in testing step).

Finally, the authors would like to acknowledge the not always subtle differences in the previous studies' data measurement/collection methods. These differences constitute a limitation of the current research and a potential source of error when compiling the data set for machine learning. However, most of the sources used for compiling the comprehensive data set needed for the training and testing of the machine learning models have followed similar data measurement methods and standard data analysis, and reporting protocols to serve a truly global international community of researchers in this field.

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776