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Genetic Programming for Multibiometrics

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Abstract

Biometric systems suffer from some drawbacks: a biometric system can provide in general good performances except with some individuals as its performance depends highly on the quality of the capture... One solution to solve some of these problems is to use multibiometrics where different biometric systems are combined together (multiple captures of the same biometric modality, multiple feature extraction algorithms, multiple biometric modalities...). In this paper, we are interested in score level fusion functions application (i.e., we use a multi-biometric authentication scheme which accept or deny the claimant for using an application). In the state of the art, the weighted sum of scores (which is a linear classifier) and the use of an SVM (which is a non linear classifier) provided by different biometric systems provide one of the best performances. We present a new method based on the use of genetic programming giving similar or better performances (depending on the complexity of the database). We derive a score fusion function by assembling some classical primitives functions (+, *, -, ...). We have validated the proposed method on three significant biometric benchmark datasets from the state of the art.

Keywords: Multibiometrics, Genetic Programming, Score fusion, Authentication.

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1. Introduction

1.1. Objective

Every day, new evolutions are brought in the biometric field of research. These evolutions include the proposition of new algorithms with better performances, new approaches (cancelable biometrics, soft biometrics, ...) and even new biometric modalities (like finger knuckle recognition [1], for example). There are many different biometric modalities, each classified among three main families (even if we can find a more precise topology in the literature) :

- *biological* : recognition based on the analysis of biological data linked to an individual (e.g., DNA analysis [2], the odor [3], the analysis of the blood of different physiological signals, as well as heart beat or EEG [4]);
- *behavioural* : based on the analysis of an individual behaviour while he is performing a specific task (e.g., keystroke dynamics [5], online handwritten signature [6], the way of using the mouse of the computer [7], voice recognition [8], gait dynamics (way of walking) [9] or way of driving [10]);
- *morphological* based on the recognition of different particular physical patterns, which are, for most people, permanent and unique (e.g., face recognition [11], fingerprint recognition [12], hand shape recognition [13], or blood vessel [14], ...).

Nevertheless, there will always be some users for which a biometric modality (or method applied to this modality) gives bad results, whereas, they are better in average. These low performances can be implied by different facts: the quality of the capture, the instant of acquisition and the individual itself but they have the same implication (impostors can be accepted or user need to authenticate themselves several times on the system before being accepted). Multibiometrics allow to solve this problem while obtaining better performances (i.e., better security by accepting less impostors and better user acceptance by rejecting less genuine users) and by expecting that errors of the different modalities are not

29 correlated. In this paper, we propose a generic approach for multibiometric
30 systems.

31 We can find different types of biometric multimodalities [15]. They use:

- 32 1. different sensors of the same biometric modality (i.e., capacitive or resistive
33 sensors for fingerprint acquisition);
- 34 2. several different representations for the same capture (i.e., use of points
35 of interest or texture for face or fingerprint recognition);
- 36 3. different biometric modalities (i.e., face and fingerprint recognition);
- 37 4. different instances of the same modality (i.e., left and right eye for iris
38 recognition);
- 39 5. multiple captures (i.e., 25 images per second in a video used for face recog-
40 nition);
- 41 6. an hybrid system composed of the association of the previous ones.

42 We are interested in the first four cases in this paper. Our objective is to
43 automatically generate fusion functions which combine the scores provided by
44 different biometric systems in order to obtain the most efficient multibiometrics
45 authentication scheme.

46 1.2. Background

47 1.2.1. Performance Evaluation

48 In order to compare different multibiometrics systems, we need to present
49 the how to evaluate them. Several works have already done on the evaluation of
50 biometric systems [16, 17]. Evaluation is generally realized within three aspects:

- 51 • *performance*: it has for objective to measure various statistical criteria
52 on the performance of the system (*Capacity* [18], *EER*, *Failure To En-*
53 *roll (FTE)*, *Failure To Acquire (FTA)*, computation time, *ROC* curves,
54 etc [17]);
- 55 • *acceptability*: it gives some information on the individuals' *perception*,
56 *opinions* and *acceptance* regarding the system;

57 • *security*: it quantifies how well a biometric system (algorithms and de-
58 vices) can resist to several types of logical and physical attacks such as
59 Denial of Service (DoS) attack.

60 In this paper, we are only interested in performance evaluation (because the
61 fusion approach is not modality dependant and perception and security depend
62 on the used modalities). The main performance metrics are the following ones:

- 63 • *FAR (False Acceptance Rate)* which represents the ratio of impostors ac-
64 cepted by the system;
- 65 • *FRR (False Rejection Rate)* which represents the ratio of genuine users
66 rejected by the system;
- 67 • *EER (Error Equal Rate)* which is the error rate when the system is con-
68 figured in order to obtain a *FAR* equal to the *FRR*;
- 69 • *ROC (Receiver Operating Characteristic)* curve which plots the *FRR* de-
70 pending on the *FAR* and gives an overall overview of system performance;
- 71 • *AUC (Area Under the Curve)* which gives the area under the ROC curve.
72 In our case, smaller is better. It is a way to globally compare performance
73 of different biometric systems.

74 We can also present the *HTER* (Half Total Error Rate) which is the mean
75 between the *FAR* and *FRR* for a given threshold (this error rate is interesting
76 when we cannot get the *EER*).

77 1.2.2. Biometric Fusion

78 There are several studies on multibiometrics. The fusion can be operated on
79 different points of the mechanism:

- 80 • *template fusion*: the templates captured by different biometric systems
81 are merged together, then the learning process is realized on these new
82 templates [19, 20]. Figure 1(a) presents this type of fusion. The fusion

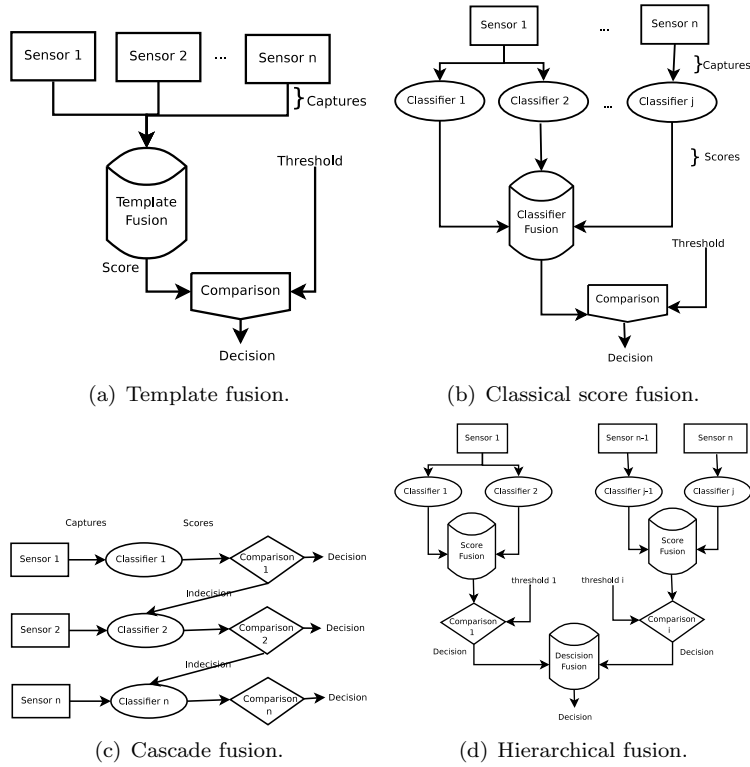


Figure 1: Illustration of different fusion mechanisms.

83 process is related to a feature selection in order to determine the most
 84 significant patterns to minimize errors.

- 85 • *decision fusion*: the decision is taken for each of the biometric authentication system, then the final decision is done by fusing the previous ones [21].
- 88 • *rank fusion*: the decision is done with the help of different ranks of biometric identification systems. The main method is the majority vote [22].
- 90 • *score fusion*: the fusion is realized considering the output of the classifiers. The Figure 1(b) presents this type of fusion.

92 BuysSENS *et al.* [23] showed the interest of biometric fusion for face recognition
 93 combining the image in visible and infrared color spaces with convolutional

94 neural networks. In [24], Mantalva and Freire have combined keystroke dynam-
95 ics with voice recognition, it seems it is the first time that multibiometrics has
96 been done with keystroke dynamics and another biometric modality. In [25],
97 Hocquet *et al.* demonstrated the interest of fusion in keystroke dynamics in
98 order to improve the recognition rates: three different keystroke dynamics func-
99 tions are used on the same capture. The sum operator (consisting in summing
100 the different scores) seems to be the most powerful approach in the literature.

101 These fusion architectures are quite simple but powerful. Results can yet be
102 improved (in term of error rate or computation time) by using different archi-
103 tectures. A cascade fusion [26] is another interesting approach. A first test is
104 done, if the user is correctly verified as the attended client or if it is detected
105 as an impostor, the algorithm stops. Otherwise, another biometric authentica-
106 tion (with another capture from another modality) is proceeded until obtaining
107 a decision of acceptance or rejection, or reaching the end of the cascade. So,
108 instead of using one decision threshold, each test (except the last one) needs
109 two thresholds: one for rejection and one for acceptance. All scores between
110 these thresholds are considered in an indecision zone. This mechanism is pre-
111 sented in Figure 1(c). Another advantage of this method is to decrease the
112 verification time by not using all the modalities, they are used only if necessary.
113 This method has been successfully applied on a multibiometric system using
114 face and fingerprint recognition in a mobile environment (where acquisition and
115 computation times are important) [26].

116 Another kind of architecture has been proposed: it is a hierarchical fusion
117 scheme [27] (called multiple layers by their authors). Shen *et al.* have pre-
118 sented this method with two different keystroke dynamics methods. The fusion
119 is done at different steps, and involves different mathematical operations on
120 scores (sum, weighted sum, product, min, max) and logical operations decision
121 (comparison to a threshold, or, and) on differents templates extracted from the
122 same capture. An extended version to any multibiometric system is presented
123 in Figure 1(d). We think our work can be seen as a generalization of this paper.

124

125 It is also possible to model the distribution of the genuine and impostor
126 matching scores, we talk about *Density-based score fusion*. In [28], scores are
127 modelled with a Gaussian Mixture Model and have been tested on three multi-
128 biometric databases involving face, fingerprint, iris and speech modalities.

129

130 Concerning non linear algorithms, Support Vector Machine (SVM) can also
131 be used in a fusion process. Each score to combine is arranged in a vector
132 and a training set is used to learn the SVM model. In [29], the SVM fusion
133 to improve face recognition gives slightly better performances than weighted
134 sum. Voice and online signature have been fused with SVM in [30]. In this
135 experiment, arithmetic mean gives best results with noise free data, while SVM
136 gives equivalent results with noisy data.

137 1.3. Discussion

138 In this paper, we are interested in biometric modality independent *transformation-*
139 *based score fusion* [28] where the matching scores are first normalized and second
140 combined. We have previously seen that in this case, arbitrary functions are
141 often used. Our work is based on these various fusion architectures based on
142 score fusion in order to produce a score fusion function automatically generated
143 with genetic programming [31].

144

145 By the way, the definition of a fusion architecture is still an open issue
146 in the multibiometrics research field [32], because the range of possible fusion
147 configurations is very large. We think that using automatically generated fusion
148 functions can bring a new solution to solve this kind of problems.

149 2. Material and Methods

150 In this section, we present all the required information in order to allow
151 other researchers to reproduce our experiment.

152 *2.1. Biometric databases*

153 As it is well known that results can be highly related to the database, for this
154 study, we have used three different multibiometric databases: the first one is the
155 BSSR1 [33] distributed by the NIST [34] (referenced as BSSR1 in the paper),
156 the second one is a database we have created for this purpose (referenced as
157 PRIVATE in the paper) and the third one is a subset of scores computed with
158 the BANCA [35] database (referenced as BANCA in the text. In fact, BANCA
159 database is composed of templates. We have used the scores available in [36]).
160 As all these databases are multi-modal, the scores are presented with tuples:
161 the i th tuple of scores is represented as $s_i = (s_i^1, s_i^2, \dots, s_i^n)$ for a database having
162 n modalities (in our case, $n \in \{4, 5\}$).

163 The three databases are presented in detail in the following subsections while
164 Table 1 presents a summary of their description.

165 *2.1.1. BSSR1 database*

166 The BSSR1 [33] database consists of an ensemble of scores sets from different
167 biometric systems. In this study, we are interested in the subset containing
168 the scores of two facial recognition systems and the two scores of a fingerprint
169 recognition system applied to two different fingers for 512 users. We have 512
170 tuples of intra-scores (comparison of the capture of an individual with its model)
171 and $512 * 511 = 261, 632$ tuples of inter-scores (comparison of the capture of an
172 individual with the model of another individual). Each tuple is composed of 4
173 scores: $s = (s_{bssr1}^1, s_{bssr1}^2, s_{bssr1}^3, s_{bssr1}^4)$, they respectively represent the score of
174 the algorithm A of face recognition, the score of algorithm B of face recognition
175 (the same face image is used for the two algorithms), the score of the fingerprint
176 recognition with left index, the score of fingerprint recognition with right index.
177 This database has been used several times in the literature [28, 37].

178 *2.1.2. PRIVATE database*

179 The second database is a chimeric one we have created by combining two
180 public biometric template databases: the AR [38] for the facial recognition and

181 the GREYC keystroke [39] for keystroke dynamics.

182

183 The AR database is composed of frontal facial images of 126 individuals
184 under different facial expression, illumination conditions or occlusions. This is
185 a quite difficult database in reason of these specificities. These images have
186 been taken during two different sessions with 13 captures per session. The
187 GREYC keystroke contains the captures on several session during a two months
188 period involving 133 individuals. Users were asked to type the password "greyc
189 laboratory" 6 times on a laptop and 6 times on an USB keyboard by interleaving
190 the typings.

191 We have selected the first 100 individual of the AR database and we have
192 associated each of these individuals to another one in a subset of the GREYC
193 keystroke database having 5 sessions of captures. We then used the 10 first
194 captures to create the model of each user and the 16 remaining ones to compute
195 the intra and inter scores.

196 These scores have been computed by using two different methods for the
197 face recognition (the scores $s_{private}^1$ and $s_{private}^2$ and three different ones for the
198 keystroke dynamics ($s_{private}^3$, $s_{private}^4$ and $s_{private}^5$ scores). The face recognition
199 algorithms are based on eigenfaces [11] and SIFT keypoints [40] comparisons
200 between images from the model and the capture [41]. Keystroke dynamics scores
201 have been computed by using different methods [42] based on SVM, statistical
202 information and rhythm measures.

203 2.1.3. BANCA database

204 The latest used benchmark is a subset of scores produced by the help of
205 the BANCA database [36]. The selected scores correspond to the following
206 one labelled: IDIAP_voice_gmm_auto_scale_25_100_pca.scores for s_{banca}^1 , SUR-
207 REY_face_nc_man_scale_100.scores for s_{banca}^2 , SURREY_face_svm_man_scale_0.13.scores
208 for s_{banca}^3 and
209 UC3M_voice_gmm_auto_scale_10_100.scores for s_{banca}^4 .

210 We have empirically chosen this subset. G1 set is used as the learning set,

Table 1: Summary of the different databases used to validate the proposed method

Nb of	BSSR1	PRIVATE	BANCA
users	512	100	208
intra tuple	512	1600	467
inter tuple	261632	158400	624
items/tuples	4	5	4

211 while G2 set is used as the validation set. Users from G1 are different than users
 212 from G2.

213 *2.1.4. Discussion*

214 The main differences between these three benchmarks are:

- 215 • the biometric modalities used in BSSR1 and BANCA have better perfor-
 216 mances than the ones in PRIVATE;
- 217 • the quantity of intra-scores is more important in PRIVATE (only one tuple
 218 of intra-score per user in BSSR1 instead of several in PRIVATE);
- 219 • BSSR1 and BANCA are databases of scores (by the way, we do not know
 220 the biometric systems having generated them) whereas PRIVATE is a
 221 database of templates (we had to compute the scores);
- 222 • BSSR1 and BANCA are more adapted to physical access control appli-
 223 cations (i.e., a building is protected by a multi-modal biometric system),
 224 while PRIVATE is more adapted to logical access control (i.e., the au-
 225 thentication to a Web service is protected by a multi-modal biometric
 226 system).

227 In the following subsections, we describe the proposed methodology to auto-
 228 matically generate a score fusion function with genetic programming. We adopt
 229 the classical score fusion context described in Figure 1(b). Before using the
 230 scores provided by different biometric systems, we need to normalize them.

231 *2.2. Score Normalization*

232 It is necessary to normalize the various scores before operating the fusion pro-
 233 cess: indeed, these scores come from different classifiers and their values do not

234 necessarily evolve within the same interval. We have chosen to use the *tanh* [43]
 235 operator to normalize the scores of each modality. Equation (1) presents the
 236 normalization method, where μ_{gen}^m and σ_{gen}^m respectively represents the average
 237 and standard deviation of the genuine scores of the modality m . The genuine
 238 scores are obtained by comparing the model and the capture of the same user:
 239 they are also called the *intra scores*. In opposition, the *inter scores* are obtained
 240 by comparing the model of a user with the capture of other users. $score'$ and
 241 $score$ respectively represents the scores after and before normalisation.

$$score' = \frac{1}{2} \left\{ \tanh \left(\frac{1}{100} \left(\frac{score - \mu_{gen}^m}{\sigma_{gen}^m} \right) + 1 \right) \right\} \quad (1)$$

242 We have selected this normalization procedure from the state of the art
 243 because it is known to be stable [44] and does not use impostors patterns which
 244 can be hard or impossible to obtain in a real application. The aim of this
 245 paper is not to analyse the performance of biometric systems depending on the
 246 normalization procedure, but to present a new multibiometrics fusion procedure.
 247 The scores of each modality have been normalized using this procedure.

248 2.3. Fusion Procedure

249 In this study, we have chosen to use genetic programming [31] in order to
 250 generate score fusion functions. Genetic programming belongs to the family of
 251 evolutionary algorithms and its scheme is quite similar to the one of genetic
 252 algorithms [45]: a population of computer programs (possibly represented by a
 253 tree) evolves during several generations; different genetic operators are used to
 254 create the new population. Programs are evaluated by using a fitness function
 255 which produces a value that is used for their comparisons and gives a probability
 256 of selection during the tournaments. In a system where the computer programs
 257 are represented by trees, their leaves mainly represent the entries of the problem,
 258 the root gives the solution to the problem and the other nodes are the various
 259 functions taking into arguments the values of their children nodes.

260 The leaves are called terminals and can be of several kinds: (a) pseudo-
 261 variables containing the real entries of the problem (in our case, the list of

262 scores of each modality), (b) some constants possibly randomly generated, (c)
263 functions without any arguments having any side effect, or (d) some ordinary
264 variables.

265 The different genetic operators usually used during the evolution are (a)
266 the crossover, where randomly choose sub-trees have two different trees are
267 exchanged, (b) the mutation, where a sub-tree is destroyed and replaced by
268 another one randomly generated, or (c) the copy, where the tree is conserved in
269 the next generation. The different steps of a genetic programming engine are
270 presented as following:

- 271 1. An initial population is randomly generated. This population is composed
272 of computer programs using the available functions and terminals. The
273 trees are built using a recursive procedure.
- 274 2. The following steps are repeated until the termination criterion is satis-
275 fied (the fitness function has reached the right value, or we reached the
276 maximum number of generations).
 - 277 (a) Computation of the fitness measure of each program (the program-
278 ming is evaluated according to its input data).
 - 279 (b) Selection of programs with a probability based on their fitness to
280 apply them the genetic operations.
 - 281 (c) Creation of the new generation of programs by applying the follow-
282 ing genetic operations (depending on their probabilities) to the pre-
283 viously selected programs:
 - 284 • Reproduction: the individual is copied to the new population.
 - 285 • Crossover: A new offspring program is created by recombining
286 randomly chosen parts from two select programs. An example is
287 provided in Figure 2.
 - 288 • Mutation: A new offspring program is created by mutating one
289 node of the selected program at a randomly chosen place. An
290 example is provided in Figure 3.
- 291 3. the single best program of the whole population is designated as the win-
292 ner. This can be the solution or an approximate solution to the problem.

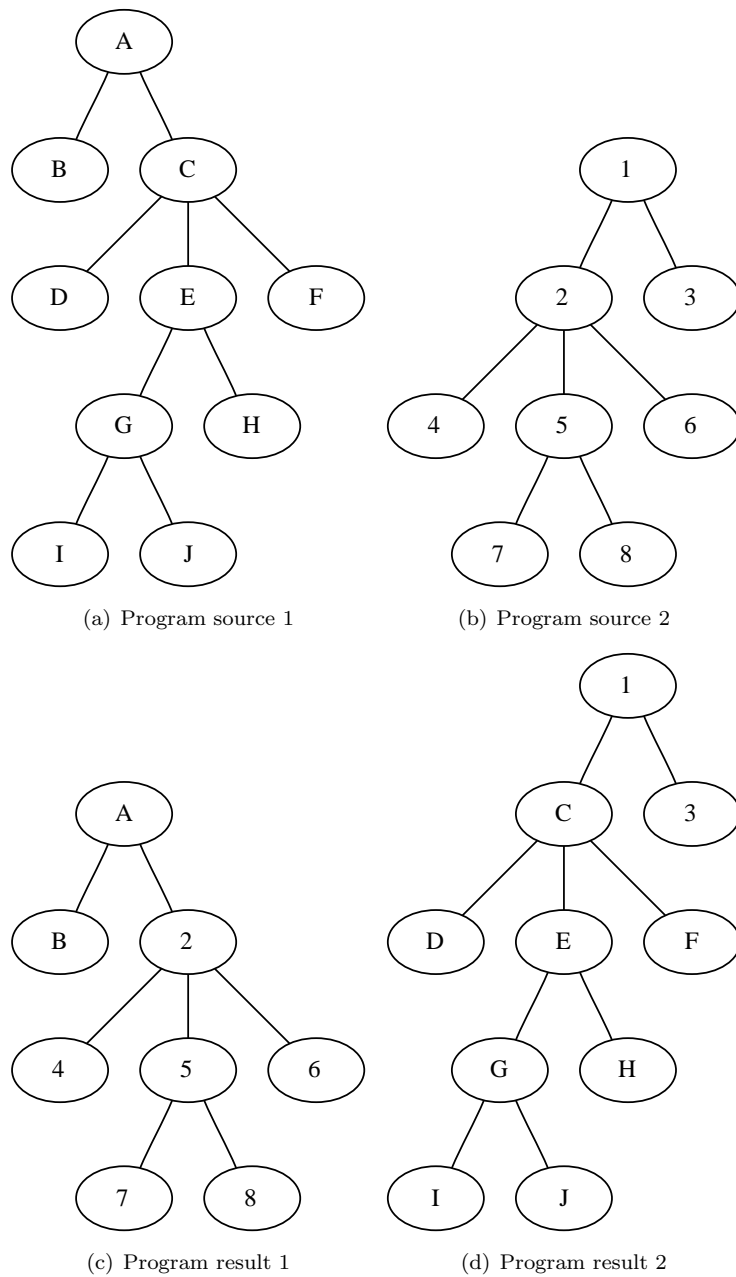


Figure 2: Crossover in genetic programming: node C from tree 1 is exchanged with node 2 from tree 2. Program result 1 is the new individual to add to the new generation.

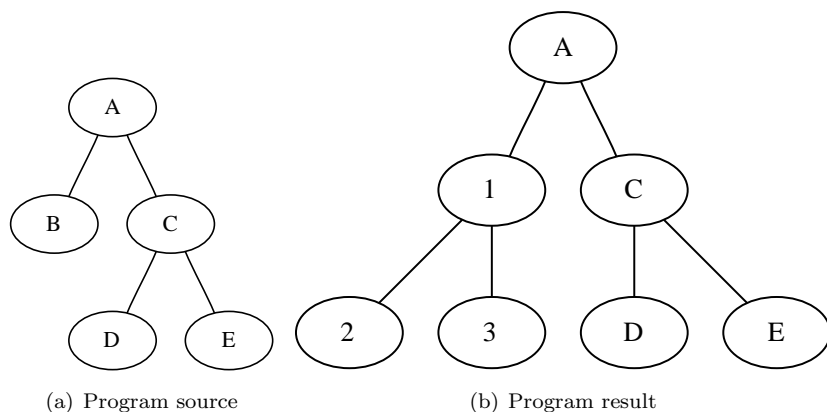


Figure 3: Mutation in genetic programming: node B is replaced by another sub-tree.

294 Different applications to genetic programming are presented in [46] as well
 295 as their bibliographic references. The fields of these applications can be listed
 296 in curve fitting, data modelling, symbolic regression, image and signal process-
 297 ing, economics, industrial process control, medicine, biology, bioinformatics,
 298 compression... but, it seems, so far of our knowledge, that it has not been
 299 yet applied to multibiometrics. We only found one reference on genetic pro-
 300 gramming in the biometrics field. In this paper [47], authors have used genetic
 301 programming to learn speaker recognition programs. They have used an island
 302 model where different islands operate their genetic programming evolution, and,
 303 after each generation some individuals are able to leave to another island. The
 304 obtained performance was similar to the state of the art in speaker recognition
 305 in normal conditions, but, the generated systems performed better in degraded
 306 conditions.

307 More information about the configuration of the genetic programming sys-
 308 tem is presented in the next section.

309 2.4. Parameters of the Genetic Programming

310 We want to use a score fusion function that returns a score related to the
 311 performance of a multibiometric system. This score has to be compared with a
 312 threshold in order to make the decision of acceptance or rejection of the user.

313 In this case, none logical operation is required in the generated programs and
314 different information can be extracted from the result of the fusion function (we
315 can compute the ROC curve, the EER, ...).

316 2.4.1. Fitness Function

317 The EER (Error Equal Rate) is usually used to compare the performance
318 of different biometric systems together. A low EER means that FAR and FRR
319 are both low and the system has a good performance if its threshold is config-
320 ured accordingly to obtain this value. For this reason, we have chosen to use
321 this running point to evaluate the performance of the generated score fusion
322 functions.

323 To compute the EER, we consider the highest and lowest values in the final
324 scores generated by the genetic programming. Then, we set a threshold at the
325 lowest score and linearly increment it until obtaining the highest score value in
326 1000 steps. For each of these steps, we compute the FAR (comparison between
327 the threshold and the inter scores) and FRR (comparison between the threshold
328 and the intra scores). The ROC curve can be obtained by plotting all these
329 couples of (FAR, FRR), while the EER is the mean of FAR and FRR for the
330 couple having the lowest absolute difference. So, the fitness function is $fitness =$
331 $(FAR_i + FRR_i)/2$, where i is the threshold for which $abs(FAR_i - FRR_i)$ is
332 minimal.

333 2.4.2. Genetic Programming Parameters

334 In this section, we present the various parameters used in the genetic pro-
335 gramming algorithm. Table 2 presents the various parameters of the evolution-
336 ary algorithm.

337 To achieve this experiment, we used the PySTEP [48] library. The generated
338 programs contain basic functions (+, -, *, /, *min*, *max*, *avg*). The terminals
339 are the scores of the biometric systems and random constants between 0 and 1.

340 The whole fitness cases are completed with a single tree evaluation, thanks to
341 the numpy [49] library. Each fitness case is a tuple of scores (where each score

Table 2: Summary of the configuration of the genetic programming iterations. Numbers used in function set can be scores or constants.

Configuration	Values						
Objective	Generates a function producing a multibiometrics score.						
Functions set	<ul style="list-style-type: none"> • +: addition of two numbers, two numbers, • -: subtraction of two numbers, • *: multiplication of two numbers, • <i>max</i>: returns the maximum of two numbers, • /: division of two numbers, • <i>avg</i>: returns the mean of two numbers • <i>min</i>: returns the minimum of 						
Fitness function	Computes the EER of the multibiometric system						
Terminal set	<table style="width: 100%; border-collapse: collapse;"> <thead> <tr> <th style="width: 33%; text-align: left;">BSSR1</th> <th style="width: 33%; text-align: left;">PRIVATE</th> <th style="width: 33%; text-align: left;">BANCA</th> </tr> </thead> <tbody> <tr> <td> <ul style="list-style-type: none"> • <i>a</i>: scores from s_{bssr1}^1, • <i>b</i>: scores from s_{bssr1}^2, • <i>c</i>: scores from s_{bssr1}^3, • <i>d</i>: scores from s_{bssr1}^4, • 50 constants linearly distributed between 0 and 1. </td> <td> <ul style="list-style-type: none"> • <i>a, b, c</i>: keystroke dynamics scores ($s_{private}^3, s_{private}^4, s_{private}^5$), • <i>d, e</i>: face recognition scores ($s_{private}^1, s_{private}^2$), • 50 constants linearly distributed between 0 and 1. </td> <td> <ul style="list-style-type: none"> • <i>a</i>: scores from s_{banca}^1, • <i>b</i>: scores from s_{banca}^2, • <i>c</i>: scores from s_{banca}^3, • <i>d</i>: scores from s_{banca}^4, • 50 constants linearly distributed between 0 and 1. </td> </tr> </tbody> </table>	BSSR1	PRIVATE	BANCA	<ul style="list-style-type: none"> • <i>a</i>: scores from s_{bssr1}^1, • <i>b</i>: scores from s_{bssr1}^2, • <i>c</i>: scores from s_{bssr1}^3, • <i>d</i>: scores from s_{bssr1}^4, • 50 constants linearly distributed between 0 and 1. 	<ul style="list-style-type: none"> • <i>a, b, c</i>: keystroke dynamics scores ($s_{private}^3, s_{private}^4, s_{private}^5$), • <i>d, e</i>: face recognition scores ($s_{private}^1, s_{private}^2$), • 50 constants linearly distributed between 0 and 1. 	<ul style="list-style-type: none"> • <i>a</i>: scores from s_{banca}^1, • <i>b</i>: scores from s_{banca}^2, • <i>c</i>: scores from s_{banca}^3, • <i>d</i>: scores from s_{banca}^4, • 50 constants linearly distributed between 0 and 1.
BSSR1	PRIVATE	BANCA					
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Initial population	500 random trees with a depth between 2 and 8 built with the ramped half and half method.						
Evolution parameters	<ul style="list-style-type: none"> • Number of individuals: 500, • Maximal number of generations: 50, • Depth limited to: 8, • Probability of crossover: 45%, • Probability of mutation: 50% • Probability of reproduction: 5% (with elitism), • Selection: tournament of size 10 with a selection probability of 80%. 						
Termination criterion	Best individual has a fitness inferior at 0.001 (by the way, this value would never be met ...) or maximal number of generations reached.						
Learning set	First half of the intra-scores tuples and first half of the inter-scores tuples.						
Validating set	Second half of the intra-scores tuples and second half of the inter-scores tuples.						

342 comes from a different biometric modality) and its result value is the score
343 returned by the generated multimodal system. The global fitness value of a tree
344 is the EER value computed with the previously generated scores (computation
345 of the ROC curve, then reading of the EER value from it).

346 PySTEP is a strongly typed genetic programming engine, but, in our case,
347 we do not use any particular constraints: the root node can only have a function
348 as child (no terminal in order to avoid an unimodal system, and any function of
349 the set), while the other function nodes can have any of the functions as children
350 as well as any of the terminals.

351 The maximal depth of the generated trees is set to 8. In order to avoid
352 to stay in a local minimal solution, the mutation probability is set to 50%.
353 500 individuals evolve during 50 generations. We have set this few quantities,
354 because during our investigations, using a population of 5000 individuals on
355 100 generations did not give so much better results (gain not interesting in
356 comparison to the computation time). Each database has been splitted in two
357 sets of equal size: the first half is the learning set and the second half is the
358 validation set.

359 The mutation rate is set to 50%, the cross-over rate to 45% and the repro-
360 duction rate to 5%. For mutation and cross-over the individuals are selected
361 with a tournament of size 10 with a probability of 80% to select the best individ-
362 ual. The same individual can be selected several times. For the reproduction,
363 the individuals are selected with an elitism scheme: the 5% best individuals are
364 copied from generation $n - 1$ to generation n . During a crossover, only the first
365 offspring (of the two generated ones) is kept.

366 **3. Results**

367 In this section, we present the results of the generated fusion programs on
368 the three benchmark data sets.

369 The results are compared to other functions from the state of the art: (a)
370 the *min* rule which returns the minimum score value, (b) the *mul* rule which

371 returns the product of all the scores, (c) the *sum* rule which returns the sum
 372 of the scores, (c) the *weight* rule which returns a weighted sum, and (d) an
 373 SVM implementation. The weights of the weighted sum have been configured by
 374 using genetic algorithm on the training sets [50, 51] (in order to give the best
 375 results as possible). The fitness function is the value of the EER and the genetic
 376 algorithm engine must lower this value. Table 3 presents the configuration of
 377 the genetic algorithm.

Table 3: Configuration of the genetic algorithm to set the weights of the weighted sum

Parameter	Value
Population	5000
Generations	500
Chromosome signification	weights of the fusion functions
Chromosome values interval	$[-10; 10]$
Fitness	EER on the generated function
Selection	normalized genetic selection (probability of 0.9)
Elitism	True

378 For the SVM, we have computed the best parameters (i.e., search the C
 379 and γ parameter giving the lowest error rate) using the learning database on
 380 a 5-fold cross validation scheme. We have used the *easy.py* script provided
 381 with libSVM [52] for this purpose. We have then tested the performance on the
 382 validation set. We only obtain on functional point (and not a curve) when using
 383 an SVM. That’s why we have used the HTER instead of the EER.

384 Table 4 presents the performances, for the three databases, of each biometric
 385 systems, fusion mechanisms from the state of the art, and our contribution.

386 Concerning the state of the art performances, can see that the simple fusion
 387 functions *sum* and *mul* tend to give better performances compared to the best
 388 biometric method of each database, but they are outperformed by the *weight* rule.
 389 The *min* operator gives quite bad results (it does not improve the best biometric

390 system). The *SVM* method gives good results but is outperform by the *weight*
391 method.

392 Table 5 presents the gain of performance against the *weight* operator (which
393 gives the best results in Table 4) in term of EER and AUC.

This gain is computed as following:

$$gain = 100 \frac{(EER_{weight} - EER_{gpfunc})}{EER_{weight}} \quad (2)$$

394 where EER_{weight} and EER_{gpfunc} are respectively the EER values of the weighted
395 fusion and the generated score fusion function (the same procedure is used for
396 the AUC). Better values than the weighted sum are represented in bold. The
397 EER gives a local performance for one running point (system configured in or-
398 der to obtain an FAR equal to the FRR), while the AUC gives a gives a global
399 performance of the whole system. These two information are really interesting
400 to use when comparing biometric systems. Figure 4 presents the ROC curves
401 of the generated programs against the weighted sum. Performance of the initial
402 biometric systems are not represented, because we have already seen that they
403 are worst than the weighted sum (same remark for the other fusion functions).
404 Logarithmic scales are used, because error rates are quite small.

405 We can see from Table 5 and Figure 4 that most of the time, the automati-
406 cally generated functions with genetic programming give slightly better results
407 than the weighted sum. These improvements can be local and global and vary
408 between 16% and 59% for the EER and 0.05% and 76% for the area under
409 the curve. When there is no improvement, the results are equal or (in one
410 case) slightly inferior. Even if there is some difference between training (not
411 represented in this paper) and validating sets, we cannot observe overfitting
412 problem. The BSSR1 dataset presents the largest difference of performance
413 between training and validation sets, but, the results are still better than the
414 ones from the state of the art (and the same problem can be observe with the
415 weighted sum). By the way, the fitness criterion has never been met, we did
416 not achieve to obtain fusion functions doing no error. So, the evolution always

Table 4: Performance (HTER in %) of the initial methods ($s_*^1, s_*^2, s_*^3, s_*^4, s_*^5$), the state of the art fusion functions ($sum, min, mul, weight$) and our proposal on the three databases. Bold values represent better performance than the initial biometric systems, and * represents fusion results better than state of the art.

(a) BSSR1

Method	HTER	
BSSR1		
Biometric systems	s_{bssr1}^1	04.30%
	s_{bssr1}^2	06.19%
	s_{bssr1}^3	08.41%
	s_{bssr1}^4	04.54%
	s_{bssr1}^5	04.54%
Fusion functions	sum	00.70%
	min	05.04%
	mul	00.70%
	$weight$	00.38%
	SVM	0.77% (FAR=1.16%, FRR=0.39%)
Proposal	gpI	0.40%

(b) PRIVATE

Method	HTER	
PRIVATE		
Biometric systems	$s_{private}^1$	8.92%
	$s_{private}^2$	11.53%
	$s_{private}^3$	15.69%
	$s_{private}^4$	06.21%
	$s_{private}^5$	31.43%
	$s_{private}^6$	31.43%
Fusion functions	sum	02.70%
	min	13.72%
	mul	02.67%
	$weight$	02.26%
	SVM	05.47% (FAR=10.87, FRR= 0.07%)
Proposal	gpA	01.57%*

(c) BANCA

Method	HTER	
BANCA		
Biometric systems	s_{banca}^1	04.38%
	s_{banca}^2	11.54%
	s_{banca}^3	08.97%
	s_{banca}^4	07.32%
	s_{banca}^5	07.32%
Fusion functions	sum	01.28%
	min	04.38%
	mul	01.28%
	$weight$	00.91%
	SVM	01.01% (FAR= 1.71 %, FRR=0.32%)
Proposal	gpΦ	00.75%*

Table 5: Performance gain between our proposal and the weighted sum (which gives the best results in the methods of the state of the art).

Database	EER	AUC
BSSR1	-5.26%	0.05%
PRIVATE	34.85%	23.85%
BANCA	17.58%	76.74%

417 ended when reaching the 50th generation.

418 Figure 5 represents the fitness evolution during all the generations of one
 419 genetic programming run on the BSSR1 database. A logarithmic scale has been
 420 used to give more importance to the low values and track easier the fitness
 421 evolution of the best individual of each generation. We can observe the same
 422 kind of results with the other databases. The fitness convergence appears several
 423 generations before the end of the computation. The worst program of each
 424 generation is always very bad which implies that the standard deviation of the
 425 fitness is also always quite huge. This can be explained by the high quantity of
 426 mutation probability and the low quantity of good programs kept for the next
 427 generation. When running the experiment several times, we obtain the same
 428 convergence value. We can say that we reach the maximum performance of the
 429 system.

430 4. Discussion

431 The score fusion functions generated by the proposed approach give a slightly
 432 better performance than the fusion functions used in the state of the art in multi-
 433 biometrics. We can argue that genetic programming is adapted to automatically
 434 define score fusion functions returning a score. The tradeoff of this performance
 435 gain is the need of training patterns which are not necessary for *sum*, *mul* or
 436 *min* (but this requirement is already present for the weighted sum or the use
 437 of an SVM). By the way, this is not really a problem, because we already need
 438 training patterns to configure the threshold of decision (if we do not want to do
 439 it empirically) or if we need to normalize the scores before doing the fusion.

440 Another problem inherent to genetic programming is the complexity of the

441 generated programs. It is probable that some subtrees could be pruned or sim-
442 plified without losing performance. Another trail would be to add regulariza-
443 tion parameter to the fitness function (for example, the number of nodes or the
444 depth of the tree). Generated programs would be more readable by a human
445 and quicker to interpret. Figure 6 presents a simple generated tree (depend-
446 ing on the database, they can be more or less complex). Even if the program
447 is quite short (comparing to the other generated functions), it includes useless
448 code (e.g., the subtree $avg(a, a - 1/12)$ could be simplified by $a - 1/24$). Some
449 generated trees include preprocessing steps by not using all the modalities in
450 the terminal set.

451 Genetic programming generated score fusion functions give performance
452 slightly equal or better than genetic algorithm configured weighted sum. Even
453 if computation time is more important than for genetic algorithm, we can think
454 that the gain is not really important between the two methods, but, to obtain
455 these results, genetic programming needed a population ten times smaller and
456 ten times less of generations.

457 5. Conclusion

458 We propose in this paper a new approach for multibiometrics based on the
459 automatic generation of score fusion functions. We have seen interesting ap-
460 proaches in the state of the art and decided to improve them by automatically
461 generated score fusion programs by the help of genetic programming.

462 Our contribution concerns the designing of multibiometric systems while
463 using a generic approach based on genetic programming (and is inspired from the
464 state of the art architectures). The proposed method returns a multibiometrics
465 score to be compared with a defined threshold. The proposed multibiometric
466 system has been heavily tested on three different multibiometric databases. We
467 obtained great improvements compared to classical fusion functions used in the
468 state of the art. We hope to have opened a new path in the fusion of biometric
469 systems thanks to genetic programming.

470 Results could surely be improved by using different parameters in the genetic
471 programming engine (i.e., more individuals and generations, different range of
472 constants, different functions, . . .). It could be interesting to test other perfor-
473 mance metrics could be improved by adding quality measures of the capture,
474 and if genetic programming could produce template fusion programs.

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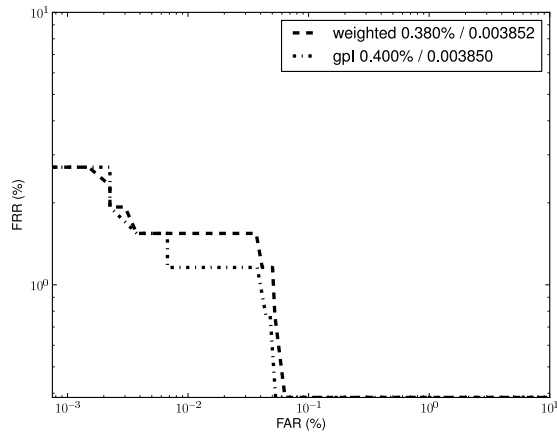
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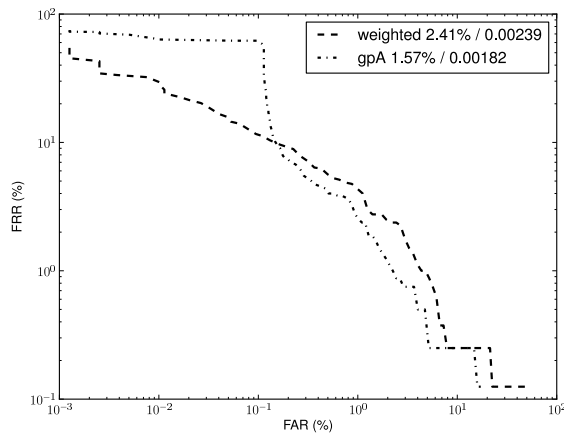
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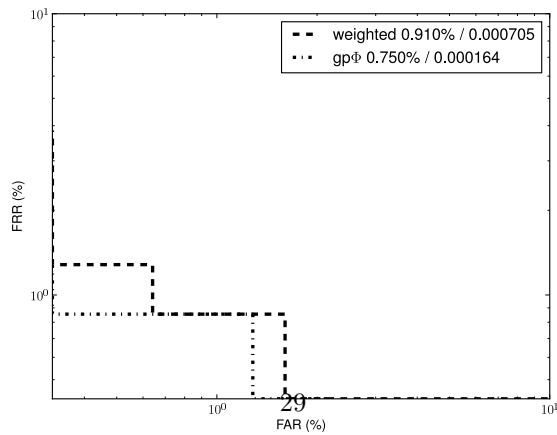
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(a) Validation with BSSR1



(b) Validation with PRIVATE



(c) Validation with BANCA

Figure 4: ROC curves of the fusion systems from the state of the art and with genetic programming. The EER of each fusion function is presented in the legend. Note the use of a logarithmic scale.

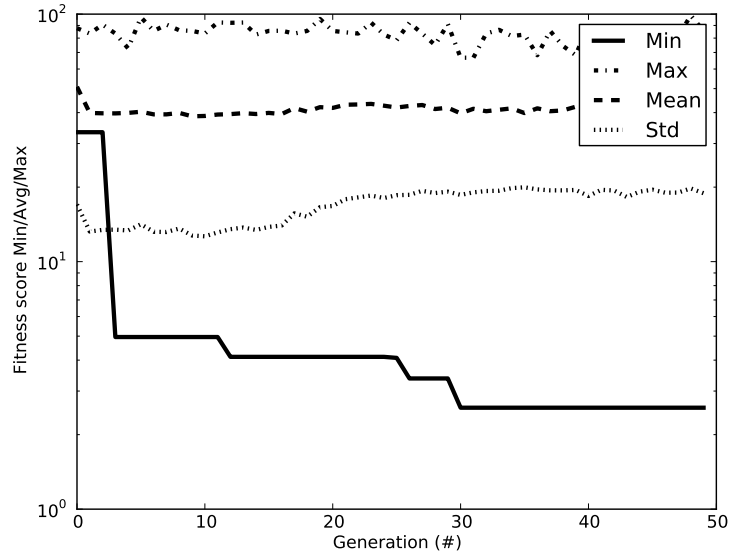


Figure 5: Fitness evolution of one run of the genetic programming evolution. The max, min, mean and std values of the fitness are represented. We want to minimize the fitness value, so lower is better.

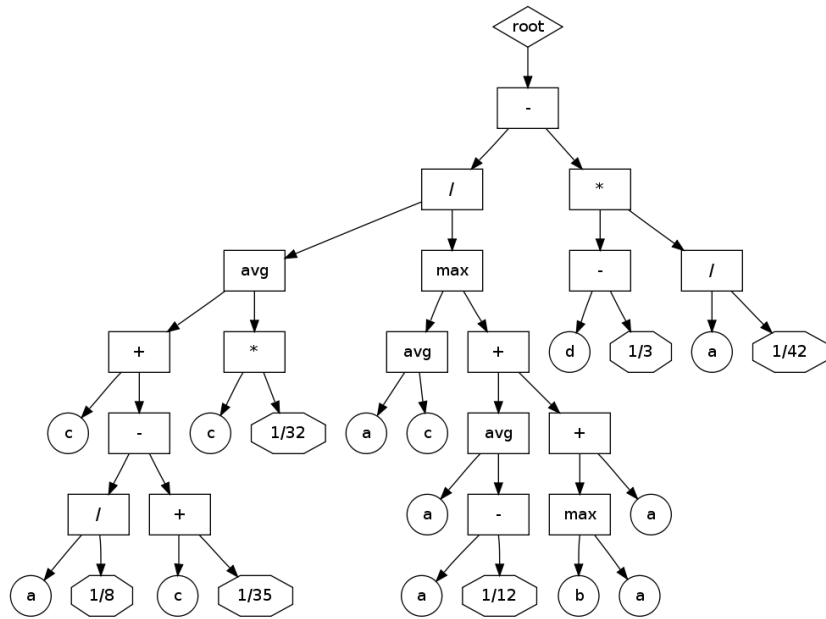


Figure 6: Sample of a "simple" generated program. We can observe the complexity of the generated fusion function.