

## A study into fingermarks at activity level on pillowcases

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## 1 **1. Introduction**

2 Forensic scientists are increasingly interested in the interpretation of evidence at activity level  
3 [1]. Activity level questions focus on the activity that led to the deposition of the evidence [2].  
4 However, for fingerprint evidence, little attention has been devoted to interpretation at  
5 activity level. Most studies on fingerprint evidence focus on the interpretation at source level,  
6 while the court frequently has to address questions at activity level.

7 An example of cases in which activity level questions related to fingerprints may arise are  
8 criminal cases with a pillow as the object of interest: was the pillow used to smother a  
9 victim?<sup>1</sup> By definition, smothering is a form of suffocation caused by an obstruction of the  
10 throat and mouth [3]. In homicidal smothering cases, an item often used to obstruct the  
11 airways is a pillow [4]. In these cases, the victim usually shows very few specific marks or  
12 traces, unless the victim resisted forcefully. This is often problematic, since smothering  
13 victims usually tend to be young, old, disabled or incapacitated by illness or drugs [4].

14 Nowadays, activity level analysis of textile fibres can be used as trace evidence in smothering  
15 cases [5]. However, the transfer of the fibres depends on several factors such as the shedder  
16 capacity of the fabric and the nature of the impact. In these cases, it would be of great interest  
17 to be able to evaluate the fingerprints on the pillowcase at activity level as well.

18 For fingerprints, the area where an item is touched will potentially contain valuable  
19 information for the evaluation of propositions at activity level. In previous research [6], we  
20 identified the variable ‘location of the fingerprints’ as an important feature that may provide  
21 information about the manner of deposition of the fingerprints. The location where a surface  
22 is touched depends on the activity carried out, and therefore the location of the fingerprints  
23 may differ between activities. Until now, the location of fingerprints in relationship to  
24 activity level questions has not been addressed in any literature and it is not known whether it  
25 is possible to derive conclusions on activity level from fingerprint patterns. More importantly,  
26 an objective method to study the location of fingerprints on items is lacking.

27 The aim of this study was to create a method to analyse the location of fingerprints on two-  
28 dimensional items. For this purpose, we used pillowcases as the object of interest to study  
29 whether we could distinguish the activity ‘smothering’ from an alternative activity like  
30 ‘changing a pillowcase’ based on the location of the touch traces left by the activities. To do  
31 so, we performed an experiment on the Dutch music festival ‘Lowlands’, in which

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<sup>1</sup> A search in a database consisting of randomly selected Dutch verdicts ([www.rechtspraak.nl](http://www.rechtspraak.nl)) resulted in at least twenty cases in the last five years in which this question was relevant. Case example: Rb Rotterdam 27 November 2014, ECLI:NL:RBROT:2014:9661.

32 participants performed two activities with paint on their hands: the activity of smothering with  
33 the use of a pillow and the alternative activity of changing a pillowcase of a pillow,  
34 representing replacing the bedding. The pillowcases were photographed and a method was  
35 designed to extract the location features of the fingermarks left on the pillowcases. A binary  
36 classification model was used to classify the pillowcases into one of the two classes,  
37 smothering and changing, based on these location features. The result is a promising model  
38 for the evaluation of propositions at activity level, based on trace locations, that could be  
39 applied to two-dimensional objects in general.

## 40 **2. Materials and methods experiment**

### 41 *2.1 Participants*

42 A total of 176 visitors of the Dutch music festival Lowlands—which took place from  
43 19/08/2016-21/08/2016—voluntarily participated in the experiment. Three participants  
44 stopped during the experiment for personal reasons. Ethical approval was obtained from the  
45 Human Research Ethics Committee (HREC) of the Delft University of Technology. The  
46 fingermarks collected during the experiment were not suitable for identification by the friction  
47 ridge pattern due to the use of an excess amount of paint.

48

### 49 *2.2 Experimental design*

50 A within-subjects design was used in which every participant was assigned to the same  
51 experimental tasks, namely performing both the smothering and changing scenario once. We  
52 used across-subjects counterbalancing for the order in which the scenarios were performed by  
53 changing the order of the scenarios every hour, for a total experimental time of 24 hours.

54

### 55 *2.3 Materials*

56 The barcode stickers used were produced on 63.5 x 29.6 mm acetate silk labels. To mark the  
57 location where the pillows have been handled, UV fluorescent skin friendly paint of the brand  
58 PaintGlow Neon UV Face and UV Body Paint was applied on the hands of each participant,  
59 in the colours blue (AA1B03), pink (AA1B04) and yellow (AA1B01). Black, 100% cotton  
60 pillowcases (70 x 60cm) by the name of DVALA and pillows (70 x 60cm) by the name of  
61 AXAG, both purchased at IKEA, were used. The pillows were covered with a water-resistant  
62 pillowcase<sup>2</sup>, and the mattress was covered with plastic foil to prevent paint cross-  
63 contamination.

64 For the experiment, two separate bedrooms were created. Next to the beds, tables were  
65 situated on which a pillowcase was placed. In the smothering scenario, a life-sized dummy of  
66 ±1.80 m with a wooden head represented the victim. The dummy was positioned in the bed  
67 under a blanket, with its head on a pressure sensor such that the pressure the volunteers  
68 exercised to smother the victim was measured. A script (Matlab®) written by the TU Delft  
69 was used to measure the performed pressure over time to check whether the participants put

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<sup>2</sup> <https://www.zorgmatras.com/waterdicht-kussen.html>

70 enough effort into smothering the victim<sup>3</sup>. The carried-out scenarios were recorded with a  
71 Logitech C615 HD webcam in each bedroom.

72 The pillowcases were photographed in a light proof photography tent for optimal UV light  
73 results. A frame with the exact dimensions of the pillowcases was used to stretch the  
74 pillowcase to remove creases. The pillowcases were photographed with a Nikon D800,  
75 60mm/2.8 lens, illuminated with UV light of wavelength 320-400 nm with the use of a  
76 Lumatec.

77

#### 78 *2.4 Experimental protocol*

79 At the start of the experiment, each participant was assigned a personal mentor who guided  
80 the participant through the experiment and tried to identify any signs of discomfort during the  
81 performance of the scenarios. In case this occurred during a scenario, the scenario was ended,  
82 and the participant went directly to the debriefing. The personal mentor started with a briefing  
83 and handed the participants four personal barcode stickers, used to mark the pillowcases used  
84 in the experiment. After providing informed consent, the participant was asked to fill in a  
85 digital questionnaire that was linked to his/her personal barcode by scanning with a hand  
86 scanner.

87 After closing the questionnaire, the participants' hands were covered with fluorescent paint  
88 using paint rollers to obtain an equal distribution of paint over the hands. Three different  
89 colours were applied to distinguish the marks of the fingers (blue), the palm (pink) and the  
90 thumb (yellow). Afterwards, the personal mentor brought the participant to the first scenario  
91 (depending on the time slot) and its corresponding bedroom. Between the scenarios, the  
92 participant washed his/her hands, and new fluorescent paint was applied.

93 In bedroom A, where pillowcases are being changed, the pillow covered in a water-resistant  
94 pillowcase was positioned on the bed. On the table next to the bed, a clean, unfolded  
95 pillowcase with its opening to the left was placed. The participant was instructed to change  
96 the pillowcase on the pillow. The instruction was to carry out this activity in the exact same  
97 way as he/she would do at home, while attempting to ignore the paint on their hands. After  
98 the scenario was carried out, the appropriate barcode stickers were placed on the pillowcase,  
99 in a corner where no paint was present. It was decided that the front side was going to be the  
100 upper side of the pillow as left on the bed. Next, the pillowcase was removed from the pillow

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<sup>3</sup> For further information on the pressure software, we would like to refer to Arjo Loeve, department Biomechanical Engineering, Delft University of Technology. Email: a.j.loeve@tudelft.nl.

101 and placed on a clothes hanger to dry. The plastic pillowcase, the foil on the mattress and the  
102 table were cleaned between experiments to prevent paint cross-contamination.

103 In bedroom B, where the smothering scenario was carried out, a pillow covered in a water-  
104 resistant pillowcase and covered in a pillowcase with its opening to the left was positioned on  
105 the table. The participant was instructed to smother the dummy using the pillow and ignoring  
106 the paint on the hands. The participant was instructed to perform enough pressure until the  
107 computer showed a blue screen, marking the end of the scenario. This occurred when a  
108 previously set pressure/time ratio was obtained. When the scenario was finished, the  
109 participant left the pillow on the bed. The pillowcases were then processed as previously  
110 described for the changing scenario. After participating in the experiment, the participants  
111 were debriefed by their personal mentor.

112 As soon as the pillowcases were dry, pictures were taken of the front side and backside of  
113 each pillowcase under UV illumination. The UV light caused the yellow paint used for the  
114 thumbs to show green, the blue paint used for the fingers to show blue and the pink paint used  
115 for the palms to show red in the resulting images.

116 **3. Image processing**

117 *3.1 Image pre-processing*

118 During the experiment, we collected four pillowcase images per donor: smothering front,  
119 smothering back, changing front and changing back. The digital images were all acquired  
120 under identical conditions. The photos were edited using Photoshop CS, following the  
121 protocol in the supplementary material. After pre-processing the images, all donors from  
122 whom four correct images were obtained were used for further analysis. A method to measure  
123 the location of the fingermarks left on the pillowcases had to be designed. We chose to  
124 transform each image into a grid in which the cells that contain fingermarks were marked.

125

126 *3.2 Image processing*

127 A software tool was developed to segment the fingermarks from the images. This  
128 segmentation process was performed in separate steps, which can be found in the  
129 supplementary material. The whole segmentation process resulted in two grid representations  
130 per pillowcase, one of the front and one of the back, in which the presence of fingermarks is  
131 marked.

## 132 4. Analysis

133 All analyses were conducted using R, version 0.99.896 [7].

134

### 135 4.1 Classification task

136 Formally, the purpose of classification is to assign the objects to a class  $C$  based on  
137 measurements on the objects [8]. The objects in our study are the pillowcases with the two  
138 classes, smothering and changing. The image classification task can then be defined as: to  
139 which class does a pillowcase belong given the position of the fingermarks? To perform this  
140 classification task, a supervised learning algorithm is used. A part of the pillowcase data set is  
141 used as a training set to train the algorithm. For all the pillowcases in this training set, we  
142 know to which class they belong. The trained algorithm is used to predict the class of  
143 pillowcases in an unseen test set. These class predictions are compared to the known classes  
144 of the pillowcases in the test set to determine the accuracy of the model.

145

### 146 4.2 Data pre-processing

147 For the data pre-processing, the design shown in Figure 1  
148 was used. Since the front and the back of one pillowcase  
149 are dependent, we decided to concatenate each two sides  
150 of a pillowcase. As a result, we obtained a 20 x 46 grid  
151 for one pillowcase, in which the right side represents the  
152 front and the left side represents the back. The final  
153 dataset consisted of two concatenated grids for each  
154 scenario per donor.

155 All donors were randomly split into three subsets: a  
156 training set, a test set and a validation set. Of the total  
157 dataset, 70% is used as training set 1 and 30% is used as a  
158 test set. Training set 1 was again divided into a training  
159 set 2 (70% of training set 1) and a validation set (30% of  
160 training set 1). Training set 2 and the validation set were used to find the right data  
161 construction and the best algorithm. Herein functioned the validation set as a test set to test  
162 each algorithm we tried during this phase. After the final algorithm was found and the results  
163 were optimized, the model was trained on training set 2, and the obtained model was used to  
164 make predictions about the unseen test set.

165

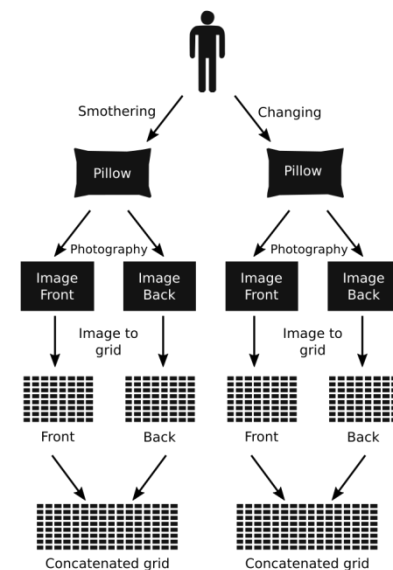


Figure 1: Data construction. The process results in two concatenated rasters per donor.



166 *4.3 Feature extraction*

167 The location of the fingermarks had to be extracted from the grids to perform the  
 168 classification task. Since it was expected that there is a higher similarity between two grids of  
 169 the same class than between two grids of a different class, we decided to use a similarity  
 170 measure between the grids. Each grid can be represented by a large vector in which every grid  
 171 cell is translated to a vector element. The similarity between two binary vectors can be  
 172 represented by a so-called similarity index, *SI* [9]. The value for *SI* ranges from 0 to 1; two  
 173 completely similar vectors have a similarity index of 1 and two completely different vectors  
 174 have a similarity index of 0. The similarity index is based on the 2 x 2 contingency table in  
 175 Table 1, in which: *a* represents the number of cells for which both vectors contain a 1  
 176 (fingermark); *b* represents the number of cells for which vector one contains a 1 (fingermark)  
 177 and vector two contains a 0 (no fingermark); *c* represents the number of cells for which vector  
 178 one contains a 0 (no fingermark) and vector two contains a 1 (fingermark); and *d* represents  
 179 the number of cells for which both vectors contain a 0 (no fingermark).

		Vector of pillowcase 2		
		1	0	
Vector of pillowcase 1	1	<i>a</i>	<i>b</i>	<i>a + b</i>
	0	<i>c</i>	<i>d</i>	<i>c + d</i>
		<i>a + c</i>	<i>b + d</i>	<i>n</i>

181 *Table 1: Contingency table. Values in this table are used to calculate the similarity between two pillowcases.*

182 A similarity coefficient between two vectors can be calculated in several ways. Since we  
 183 observed that the absence of fingermarks on a pillowcase also provides information on the  
 184 class to which the pillowcase belongs, we chose for the ‘simple matching coefficient’ of  
 185 Sokal and Michener [10], which also takes the matching ‘empty’ cells into account:

$$SI = \frac{a + d}{n} \tag{1}$$

186 Using the *SI*, the Euclidean distance (*d*) between two vectors can be expressed as:

$$d = \sqrt{1 - SI} \tag{2}$$

187 This method was used to obtain a distance measure between two grids of pillowcases. For  
 188 each grid, the distances to each of the grids in the training set smothering and to each of the  
 189 grids in the training set changing were calculated. As a result, each grid can be represented as  
 190 a feature vector  $\begin{pmatrix} x_1 \\ x_2 \end{pmatrix}$  where  $x_1$  represents its mean distance to the training set smothering and

191  $x_2$  represents its mean distance to the training set changing. A grid of a smothering pillowcase  
192 will be more similar to the grids of other smothering pillowcases than to the grids of changing  
193 pillowcases, resulting in a lower distance to the smothering training set and a higher distance  
194 to the changing training set. For the grid of a changing pillowcase, the reverse reasoning  
195 holds. Based on these distance measures, we expect that the grids of the pillowcases of both  
196 scenarios can be quite well separated.

197 The feature vectors of all pillowcases together form a so-called feature space and a  
198 classification rule partitions the feature space into regions [11]. In our study, we were looking  
199 for a classification rule that partitioned the feature space into the two regions smothering and  
200 changing. To determine the decision boundary between these two regions, the approach of  
201 Quadratic Discriminant Analysis (QDA) was used.

202

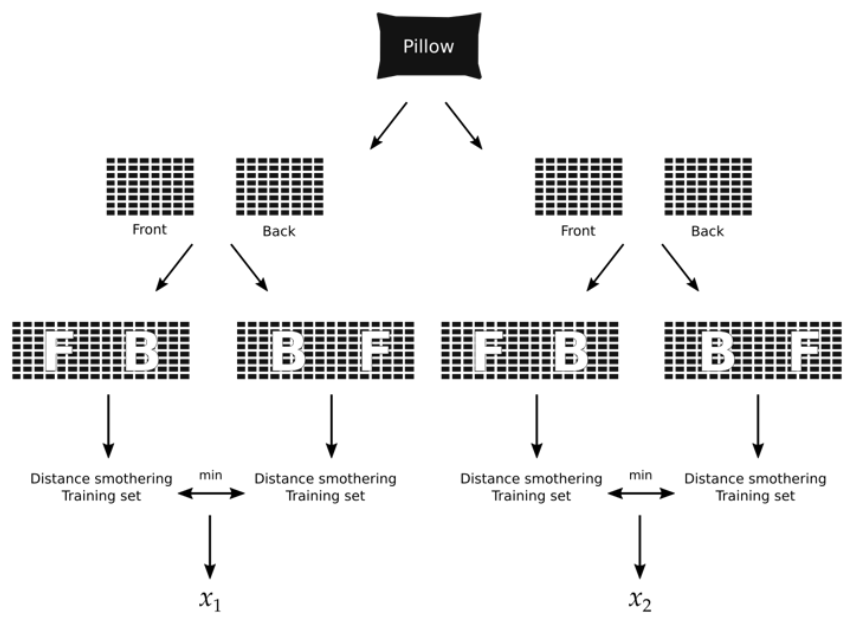
#### 203 *4.4 Classification*

204 To construct the classification system, a quadratic discriminant analysis (QDA) classifier was  
205 used to classify each feature vector of a pillowcase into one of the classes smothering or  
206 changing. For further explanation of quadratic discriminant analysis, see James, Witten,  
207 Hastie and Tibshirani [12].

208

#### 209 *4.5 Side of the pillowcase*

210 The proposed model was built under the assumption that it was known which side of the  
211 pillowcase was used for smothering. Because it is highly unlikely that this information is  
212 available in forensic casework, we classified the test set without using this information. For  
213 each donor in the test set, we concatenated the two grids of a pillowcase in two ways: one of  
214 which the front side was on the left and one of which the front side was on the right, as shown  
215 in Figure 2. For both these concatenated grids, the distance to the set smothering and to the set  
216 changing were determined. The concatenated grid for which the distance to the training set  
217 smothering was minimal was taken to be the most likely concatenation for a smothering  
218 pillowcase; this distance is used for the value of  $x_1$ . The concatenated grid for which the  
219 distance to the set changing was minimal was taken to be the most likely concatenation for a  
220 changing pillowcase; this distance is used for the value of  $x_2$ . By comparing the concatenation  
221 order chosen by the model with the known concatenation order for the test set, we can study  
222 the ability of the model to predict the front and the back of a pillowcase.



223

224 *Figure 2: Data construction. Process of testing the test set without using the side of the pillowcase.*

225 *4.5 Programming in R*

226 For the implementation of the analysis in R, the following packages were used:

- 227 - *Raster* for all grid computations [13];
- 228 - *Ade4* to compute distance measures [14];
- 229 - *MASS* to perform QDA [15]; and
- 230 - *MVN* to test assumptions for QDA [16].

231

232 **5. Results**

233 *5.1 Participants*

234 We obtained two pillowcases each from 173 volunteers, resulting in 704 images.  
235 Unfortunately, not every image was suitable for analysis due to photography issues such as  
236 movement, incorrect lightning or creases. For these images, the quality of the image was too  
237 poor or the location of the fingermarks was shifted due to creases, and therefore these images  
238 could not be used for further analysis. For the final analysis, we selected all donors for whom  
239 all four images were determined correct according to the protocol described in the  
240 supplementary material, resulting in 132 donors and 528 images. Table 2 shows the  
241 characteristics of these 132 participants. The group consisted of 59 men and 68 women, with  
242 an age ranging from 18 to 60 years old ( $M = 28.0$ ,  $SD = 8.3$ ).

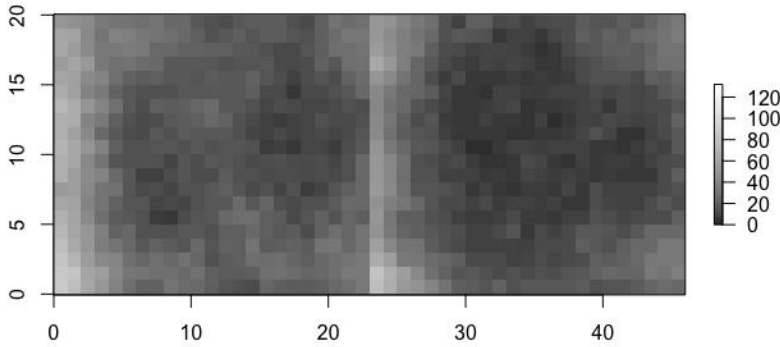
243

Characteristics of participants		<i>n</i>	Percentage
<b>Sex</b>	Men	59	45%
	Women	68	51%
	Unknown	5	4%
<b>Age</b>	<30	82	62%
	31-50	43	33%
	>50	4	3%
	Unknown	3	2%

244 *Table 2: Characteristics of the volunteers who participated in the experiment.*

245 *5.2 Heat map*

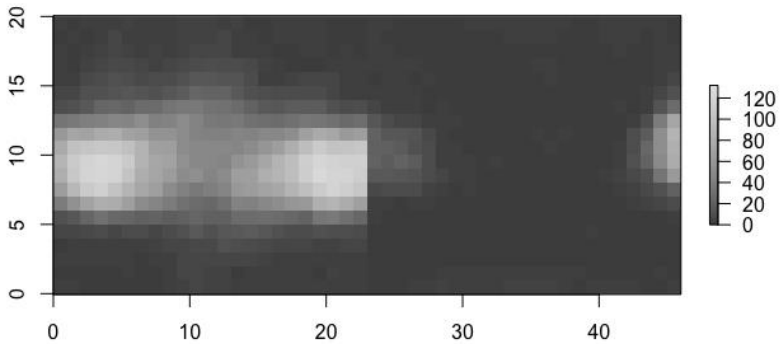
246 Figure 3 and Figure 4 show heat maps of the grids for the changing scenario and the  
247 smothering scenario, respectively. These heat maps show the concatenated grids of the front  
248 side and back side of the pillowcase, with the opening on the left-hand side. The heat maps  
249 show meaningful differences with regard to the location of the fingermarks between the two  
250 scenarios. The traces caused by changing a pillowcase show a random distribution over the  
251 pillowcase for both the front and the backside of the pillowcase, with a higher distribution of  
252 fingermarks around the opening of the pillowcase. The traces caused by smothering with the  
253 pillow show a high density of traces in the middle lane of the front side of the pillowcase. On  
254 the back side of the smothering pillowcases, almost no fingermarks are found, and the  
255 fingermarks that are found are mostly around the opening of the pillowcase.



256

257

Figure 3: Heat map changing. Shows the heat map of the concatenated pillowcases used under the scenario changing.



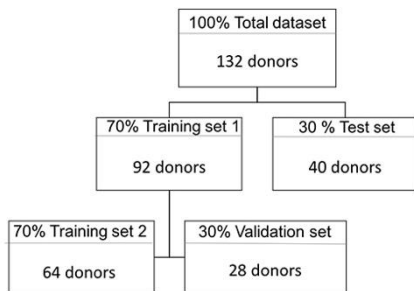
258

259

Figure 4: Heat map smothering. Shows the heat map of the concatenated pillowcases used under the scenario smothering.

### 260 5.3 The classification model

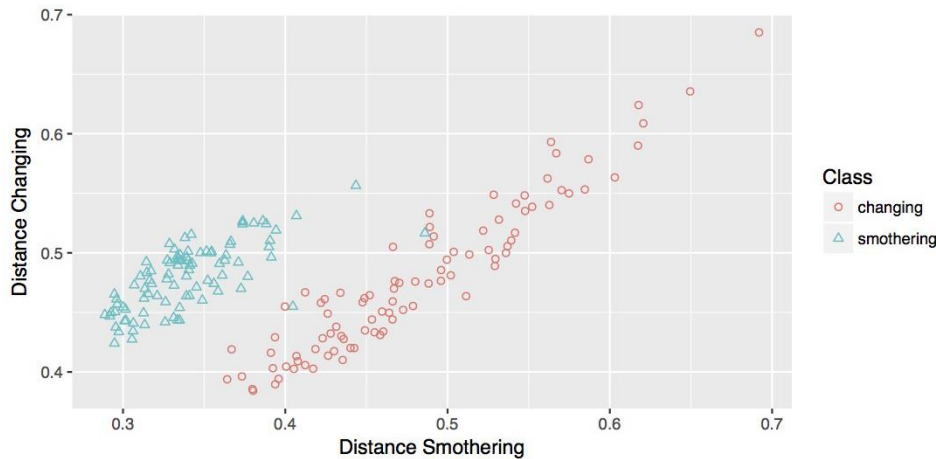
261 The 132 donors were randomly split into three subsets, a training set, test set and a validation  
 262 set, as shown in Figure 5. Training set 2 and the validation set were used to optimally fit the  
 263 model. For each pillowcase in training set 2, the distances to the training set smothering and  
 264 to the training set changing are calculated. The resulting feature space is shown in Figure 6.  
 265 The red dots represent the changing pillowcases, and the blue dots represent the smothering  
 266 pillowcases. Figure 6 shows that the two classes smothering and changing are distributed into  
 267 two reasonably separate regions.



268

269

Figure 5: Subsets of total dataset. Division of donors into three separate subsets.



270

271 *Figure 6: Feature space. Shows the distribution of the pillowcases based on the distance measures.*

272 A QDA classifier assumes the classes to be multivariate normally distributed. We have tested  
 273 this assumption using the Mardia test and QQ plots (see supplementary material). From the  
 274 Mardia test, it appeared that the data were not multivariate normal within the classes. Because  
 275 multivariate outliers are a reason for violation of the multivariate Gaussian assumption [16],  
 276 we studied the QQ plot of each class. It appeared that there are a few outliers that distort the  
 277 normality assumption. Besides these outliers, the data follow a normal distribution, and we  
 278 assume that with a bigger dataset, the assumption of a multivariate Gaussian distribution for  
 279 each class is met and QDA can be applied. A summary of the resulting QDA model is  
 280 available as supplementary material.

281

#### 282 *5.4 Evaluation of the model*

283 Table 3 summarizes the results of classifying the observations in the test set with the QDA  
 284 classifier. The model classified 39 of the 40 pillowcases correctly, representing a model  
 285 accuracy of 98.8%. Of particular interest are the errors obtained when applying the model.  
 286 Table 3 shows that the error is a smothering pillowcase that is classified as a changing  
 287 pillowcase. Within the forensic science community, these false-negative errors are determined  
 288 to be less problematic than false-positive errors, which are highly undesirable since they  
 289 involve a higher possibility of an unfair decision-making [17]. When looking more closely at  
 290 the pictures and video footings of this false negative, we found that the donor rotated the  
 291 pillow 45 degrees before starting smothering, resulting in a trace pattern exactly 45 degrees  
 292 rotated from the pattern observed in the heat map for smothering.

293

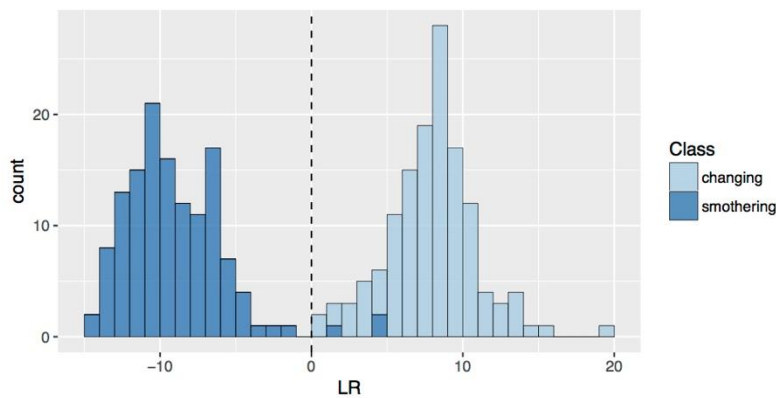
Test set	Changing	Smothering
Changing predicted	40	1
Smothering predicted	0	39

294 *Table 3: Confusion matrix for the Test set using the QDA classifier.*

### 295 5.5 Likelihood ratio

296 Since classification using QDA is based on the posterior probability  $Pr(Y = k|X = \mathbf{x})$  for  $k=$   
 297 (smothering, changing) and  $\mathbf{x}$  a feature vector of the corresponding pillowcase, a likelihood  
 298 ratio can be determined for each pillowcase. With the use of a prior probability of 0.5 for each  
 299 class, the posterior probability is equal to the likelihood ratio. Therefore, the model directly  
 300 provides a likelihood ratio for each pillowcase in the classes smothering and changing. The  
 301 distribution of the likelihood ratios obtained from the total set can be observed in Figure 7, in  
 302 which the range of the  $\log_{10}(\text{LR})$  values can be seen on the x-axis. This figure shows that the  
 303 likelihood ratios for the classes changing and smothering are almost perfectly separated.  
 304 However, there are smothering pillowcases that obtain a likelihood ratio in favour for the  
 305 scenario changing, resulting in misleading evidence in these cases [18]. These are the three  
 306 misclassified smothering pillowcases discussed previously.

307



308

309 *Figure 7: Likelihood ratio distribution. Shows the calculated LR for each pillowcase.*

### 310 5.6 Side of the pillowcase

311 Table 4 represents the results of predicting the order of concatenation of the grids in the test  
 312 set. The results show that the front and back side of the smothering pillowcases were all  
 313 predicted correctly. The front and back side of the changing pillowcases are wrongly  
 314 predicted in 37.5% of the cases. This can be explained by the fact that the front and the back  
 315 side of the changing pillowcases show similar distributions of fingermarks, whereas the front  
 316 and the back side of smothering pillowcases show very different distributions of fingermarks.

317

	<b>Correct predicted order</b>	<b>Incorrect predicted order</b>
<b>Smothering</b>	40	0
<b>Changing</b>	25	15

318 *Table 4: Results of predicting the order of concatenation.*

319



## 320 **6. Discussion and Conclusion**

321 The purpose of this study was to create a method to analyse the location of fingerprints on  
322 two-dimensional items. For this purpose, we used pillowcases as the object of interest to study  
323 whether the activity of smothering with a pillow can be distinguished from the alternative  
324 activity of changing a pillowcase, based on the fingerprints left by the activity. The results of  
325 our classification model show that the fingerprint patterns caused by smothering with a  
326 pillow can be well distinguished from the fingerprint patterns caused by changing a  
327 pillowcase based on the location of the traces, with a model accuracy of 98.8%. The results  
328 support the expectation that the location of the fingerprints on a pillowcase provides valuable  
329 information about the activity that is performed with it.

330 The proposed model misclassified one pillowcase for belonging to the changing class when it  
331 actually belonged to the smothering class. When studying this pillowcase, we learned that the  
332 resulting trace pattern showed a rotation of 45 degrees compared with the trace pattern on the  
333 other smothering pillowcases. This was the only pillowcase in the test set for which this  
334 pattern is observed, and the model directed us to this 'exception'. After examining the training  
335 set and the validation set, we found two other pillowcases showing this trace pattern. We  
336 expect that with a larger sample size, these rotated pillowcases will be observed more often,  
337 resulting in a larger number of rotated pillowcases in the training set. Consequently, the  
338 learning algorithm based on the training set will probably learn that the rotated variant also  
339 belongs to the class smothering, resulting in a model that might predict the right class for the  
340 rotated variant. Another possibility might be to assign a third class representing the rotated  
341 variants. This might result in a classification model in which the pillowcases are classified  
342 into three separate classes: changing, smothering and rotated smothering.

343 In this experiment, the side of the pillowcase that was used for smothering is known. In  
344 forensic casework, this information will not be available. Therefore, we tested the pillowcases  
345 in the test set without using this information. The results show that the front and the back of  
346 the pillowcases used for smothering are determined correctly in 100% of the cases. For  
347 changing pillowcases, 62.5% of the pillowcases were correctly determined. It is not of much  
348 interest to determine the front and back of a pillowcase that is used for changing; however, it  
349 can be highly valuable to be able to determine the front and back of a pillowcase that is used  
350 for smothering, since it makes a targeted sampling for DNA possible. This information,  
351 together with the location information of the fingerprints, may provide valuable information  
352 in smothering cases, especially on the activity level interpretation of the fingerprints.

353 Performing the experiment at a music festival such as Lowlands allowed us to obtain many  
354 participants in only one weekend. Normally in forensic casework, it is often challenging to  
355 obtain a dataset of the size we obtained. For cases in which this might be challenging, citizen  
356 science projects such as the one we performed on Lowlands may offer a solution, as also  
357 shown by Zuidberg, Bettman, Aarts, Sjerps and Kokshoorn [19]. The results show a large  
358 variety of donors, and the results of the experiment can be based on a relatively large sample.  
359 Although the results of our experiment are promising, there are some important limitations  
360 that make direct implementation in casework difficult. One drawback of practical experiments  
361 in forensic science is that it is difficult to reconstruct a realistic murder scenario. In real life,  
362 the person who is smothered will very likely resist. This could not be simulated in our  
363 experiment. Additionally, the time it takes to smother a person will be up to a few minutes  
364 [20]. Due to the fact that the experiment had to be suitable for a festival and we did not want  
365 to emotionally and physically burden participants excessively, we used a smothering time of  
366 around 45 seconds, depending on the pressure performed. Another point to mention is that we  
367 used paint for the detection of the fingermarks. The resulting paint traces are not directly  
368 comparable to the results when visualizing fingermarks with the use of visualization methods.  
369 Further research should reveal whether the model is also applicable to visualized fingermarks.  
370 An additional limitation is that we only considered the two activities smothering and  
371 changing, both independent of each other. In real life, a pillowcase that is used for smothering  
372 may contain other fingermarks caused by changing the pillowcase and other activities. It  
373 would be of interest to study these combined activities to see whether it is possible to select  
374 the fingermarks that resulted from smothering to make targeted DNA sampling possible.  
375 It must be noted that the likelihood ratio values for the pillowcases obtained with our model  
376 are very high. These are not the likelihood ratio values we expect to obtain in real cases.  
377 However, this research shows a first proof of concept of the possibility to differentiate  
378 between two separate activities based on the location of the fingermarks. Further research  
379 should demonstrate whether these results are also applicable to casework situations in which  
380 pillows are the object of interest.

381 A limitation of the proposed classification model is that the training set must consist of data  
382 that has exactly the same dimensions as the data in the test set. For example, the resulting  
383 model based on a training set consisting of pillowcases with dimensions 60 x 70 may not  
384 directly be applicable to pillowcases with a different ratio because the size of the fingermarks  
385 does not change in the same ratio as the size of the pillows. Further research is necessary to  
386 overcome this problem.

387 Of great importance is that the resulting model is not only limited to pillowcases; we propose  
388 a promising model for studying trace locations at activity level that could be applied to two-  
389 dimensional objects in general. This means that the model can be applied to all two-  
390 dimensional items for which we expect that different activities will lead to different locations  
391 of fingerprints. As long as the traces can be visualized, the proposed method can be trained to  
392 classify the items into separate classes based on the location of the traces. The only difference  
393 is that the learning algorithm of the model must be trained with a new training set consisting  
394 of grids representing these new two-dimensional objects. In the future, the method may even  
395 be adjusted to account for studying fingerprint locations on three-dimensional objects. This is  
396 a recommendation for further research.

397 For the analysis of fingerprints at activity level, this study provides an important step  
398 forward. Until now, many of the variables that provide information for fingerprint evaluation  
399 at activity level have not been studied yet, and their probabilities can only be based on expert  
400 experience. We showed an example of how the variable location can be studied with the use  
401 of an experiment. This information can be implemented in a Bayesian network to study the  
402 evaluation of fingerprints at activity level in casework [6].

403 **Supplementary material**

404 *1. Image processing protocol*

- 405 1. Duplicate image.
- 406 2. Rotate the image such that the opening of the pillowcase points to the left.
- 407 3. Adjust the brightness such that the corners of the pillowcase can be observed.
- 408 4. Crop the pillowcase with a 60 x 70 cm frame.
- 409 5. In case the pillowcase is smaller than the 60 x 70 cm frame due to incorrect stretching of
- 410 the pillowcase during the photography, use the option `transform > distort` based on bicubic
- 411 interpolation. Stretch the picture such that the pillowcase matches the 60 x 70 cm frame.
- 412 6. Mask the barcode label on the pillowcase.
- 413 - If there is no paint near the barcode label, we assume the barcode label was placed
  - 414 on a non-paint area as instructed in the protocol. Place a grey rectangle with an
  - 415 RGB value of (20,20,20) and of size equal to the barcode label over the barcode
  - 416 sticker.
  - 417 - During the experiment, we observed that on some pillowcases, it was difficult to
  - 418 place the label in a non-paint area. If there is an indication for the presence of paint
  - 419 beneath the label, place a transparent rectangle of 0% of size equal to the barcode
  - 420 label over the barcode sticker. Transparent pixels will later in the process be
  - 421 translated to missing values.
- 422 7. In case part of the pillowcase is not photographed due to movement of the camera or
- 423 skewing of the pillow, mask the area within the 60 x 70cm frame that contains missing
- 424 data with a transparent layer of 0%.
- 425 8. Save the picture as a JPEG file if there are no transparent areas in the image. Save the
- 426 picture as a PNG file if there are transparent areas in the image.
- 427 9. In case one of the following problems occurs, remove the donor from the dataset.
- 428 - Borders of the pillowcase could not be determined due to movement of the camera
  - 429 or wrong lightning conditions during the image-acquisition process.
  - 430 - Wrong stretching of pillowcase caused a substantial distortion in the pillowcase.

431 2. Segmentation software Lexie

432 A software tool called Lexie was developed to segment the fingerprints from the images. This  
433 segmentation process was performed in separate steps.

434

435 2.1 Colour extraction

436 Different areas of the hand left different coloured marks on the pillow. These marks were  
437 extracted to three separate images based on the colour vectors and the hue of the pixel values,  
438 resulting into three grey scale images. The image intensity ranges were then normalized to the  
439 same intensity range to allow the same segmentation settings for each image.

440 To extract a colour from an image, all pixel values were compared to three predefined colours  
441 that defined the fingerprints for the fingers, palm and thumb of the hand. A colour vector  $\vec{c}$  is  
442 equivalent to the triple red, green and blue value of a pixel. The more the colour vectors of the  
443 pixel and of the predefined colour point in the same direction, taking the length of the vector  
444 into account, the more a pixel is considered to match the predefined colour. To strengthen the  
445 colour extraction, the hue of the pixel and the predefined colours were also compared. The  
446 hue value of a pixel ranges between 0 and 360 and it is circular, meaning that a hue of 360 is  
447 equal to the hue of 0. If the hue of the pixel compared to the hue of the predefined colour  
448 differed more than 120, the colours were considered not equal, resulting in an intensity of 0  
449 for that pixel in the resulting image. If the difference was less than 120, the linear ratio of this  
450 difference was defined as the hue-factor.

451 This extraction process, which extracts an intensity  $I$  for each pixel  $p$  can be formally defined  
452 as:

$$I_{i,p} = 255 \cdot \frac{\vec{c}_i \cdot \vec{c}_p}{|\vec{c}_i|} \cdot H_{i,p} \quad (1)$$

453

454 where  $i$  represents fingers, palm or thumb,  $\vec{c}_i$  its corresponding predefined colour and  $\vec{c}_p$  the  
455 color of the pixel  $p$ . The hue-factor  $H_{i,p}$  is defined as:

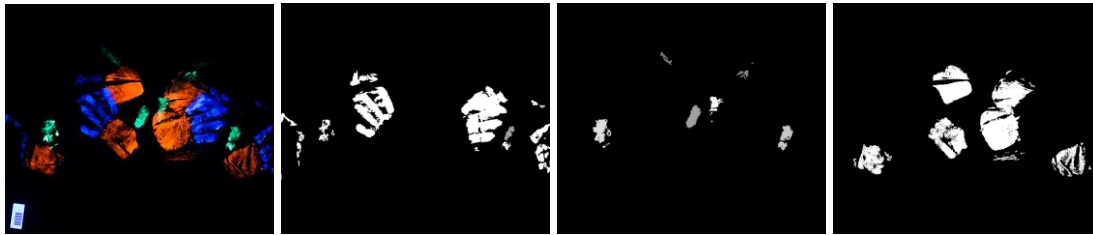
456

$$H_{i,p} = \max\left(\frac{|h_i - h_p| \bmod 360 - 180}{120}, 0\right) \quad (2)$$

457 where  $h_i$  is the hue value of  $\vec{c}_i$  and  $h_p$  the hue value of  $\vec{c}_p$ . Applying this for the three  
458 predefined colours resulted into three grey scale images with intensity ranging between 0 and

459 255. Figure S1a shows an example of a pre-processed image, before analysis in Lexie. Lexie  
460 extracts the colours as denoted in Figures S1b-S1d.

461



462

463 (a) Original

(b) Fingers

(c) Thumbs

(d) Palms

464 *Figure S1: Image segmentation with Lexie. Visualization of the segmentation steps.*

## 465 2.2 Segmentation

466 Contours of the fingermarks on pillows were identified using a four-neighbour based region  
467 growing segmentation using seed and thresholding [21]. This pixel based segmentation  
468 method uses a threshold for contour definition and a seed for region selection and could be  
469 easily applied to the three grey scale images. Pixels with an intensity equal to the seed value  
470 or higher are called the seeds. Neighbouring pixels of the seeds were evaluated. If its intensity  
471 was above the threshold level, then its neighbouring pixels were also evaluated. This process  
472 continued until it reached a pixel that was below the threshold level. This resulted in regions  
473 around the seeds, which defined clusters of pixels identified as fingermarks.

474

## 475 2.3 Filtering

476 After segmentation, an additional filter was applied based on the surface of the fingermarks to  
477 remove noise elements from the segmentation. Noise elements are small regions that can be  
478 caused by drops of paint or dust reflection of the pillow. The surface-threshold allows  
479 removing these regions that are not considered fingermarks. Regions with a surface smaller  
480 than the surface-threshold were removed from the segmentation.

481

## 482 2.4 Partitioning

483 For the final analysis, the three images are partitioned by a grid, which represents the location  
484 areas. For each partition, the number of pixels that are part of a fingermark were counted,  
485 which allowed for an analysis of fingermark occurrences per cell. If a fingermark was present  
486 that contained more than 5% of the surface of the cell, then the cell was marked as containing  
487 a fingermark.

488 Some pillowcase images contained hidden fingermarks due to skewing of the pillow during  
489 photography or when the personal barcode stickers were placed on paint. These areas were  
490 marked by changing the transparency of these pixels to 0% during the image pre-processing  
491 step. If in a grid cell 5% of the surface of the cell was transparent, then the whole cell was  
492 marked with NA.

493

#### 494 2.5 Settings Lexie

495 To find the optimal settings of the segmentation software, manually prepared grids were  
496 compared to the results of the software for different settings of the threshold, seed and the  
497 250 surface-threshold. Four pillowcase pictures of one donor were manually transformed into  
498 a grid by two independent researchers. The manual results were compared, and in  
499 consultation, one grid for each pillowcase was found. These final manual grids were  
500 compared to the results obtained by Lexie for different settings. The optimal settings were  
501 used for the analysis of all images, in which each image is transformed to a 20 x 23 grid with  
502 cell size of 3 x 3cm.

503 3. Multivariate Normality testing

504 The assumption of multivariate normally distributed data within each class is tested using the  
505 Mardia test and QQ plots. The results are shown in Figures S2, S3, S4, S5 and S6.

506

```
Mardia's Multivariate Normality Test
-----
data : data_smoren_training[c(1, 2)]

g1p      : 3.159239
chi.skew : 48.44167
p.value.skew : 7.634565e-10

g2p      : 12.60609
z.kurtosis : 5.522511
p.value.kurt : 3.341886e-08

chi.small.skew : 51.12066
p.value.small : 2.106315e-10

Result    : Data are not multivariate normal.
-----
```

507

508 Figure S2: Output R for the Mardia test to assess multivariate normality for the class smothering.

509

```
Mardia's Multivariate Normality Test
-----
data : data_opmaken_training[c(1, 2)]

g1p      : 0.9620601
chi.skew : 14.75159
p.value.skew : 0.005245167

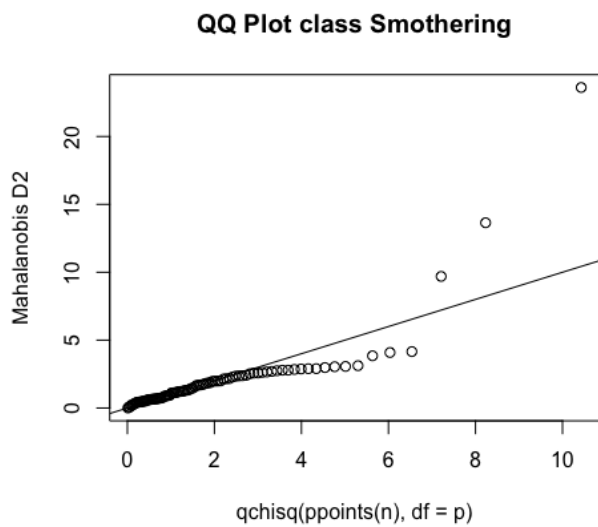
g2p      : 7.382909
z.kurtosis : -0.7398666
p.value.kurt : 0.459381

chi.small.skew : 15.5674
p.value.small : 0.003658131

Result    : Data are not multivariate normal.
-----
```

510

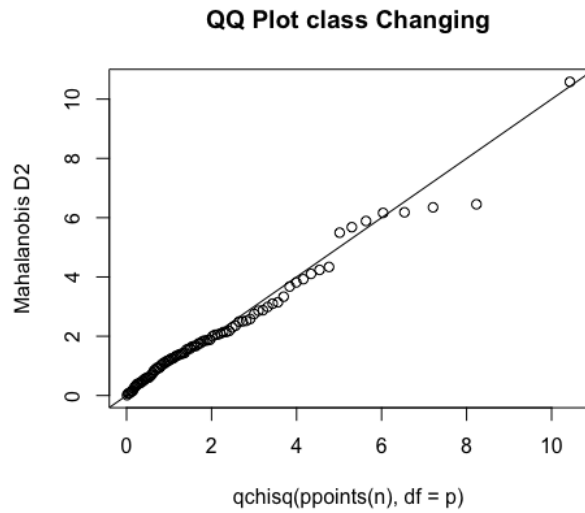
511 Figure S3: Output R for the the Mardia test to assess multivariate normality for the class changing.



512

513 Figure S4: QQ plot smothering. Used to assess multivariate normality for the class smothering.





514

515 *Figure S5: QQ plot changing. Used to assess multivariate normality for the class changing.*

516

```
Call:
qda(Klasse ~ ., data = alledata[, c(1, 2, 3)])
```

Prior probabilities of groups:

changing	smothering
0.5	0.5

Group means:

	Dist_Smoren	Dist_Opmaken
changing	0.4746263	0.4690370
smothering	0.3351587	0.4746263

517

518 *Figure S6: Fitted QDA model.*

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