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2 **Labelling and Family Resemblance in the**
3 **discrimination of polymorphous categories**
4 **by pigeons**

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20 Animal Cognition MS AC-09-0161, Accepted for publication subject to revision

21 This draft last saved 1 July 2010 by SEGL

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23

24 **Abstract**

25

26 Two experiments examined whether pigeons discriminate polymorphous categories
27 on the basis of a single highly predictive feature or overall similarity. In the first
28 experiment, pigeons were trained to discriminate between categories of photographs
29 of complex real objects. Within these pictures, single features had been manipulated
30 to produce a highly salient texture cue. Either the picture or the texture provided a
31 reliable cue for discrimination during training, but in probe tests, the picture and
32 texture cues were put into conflict. Some pigeons showed a significant tendency to
33 discriminate on the basis of the picture cue (overall similarity or family resemblance),
34 whereas others appeared to rely on the manipulated texture cue. The second
35 experiment used artificial polymorphous categories in which one dimension of the
36 stimulus provided a completely reliable cue to category membership, whereas three
37 other dimensions provided cues that were individually unreliable but collectively
38 provided a completely reliable basis for discrimination. Most pigeons came under
39 the control of the reliable cue rather than the unreliable cues. A minority, however,
40 came under the control of single dimensions from the unreliable set. We conclude
41 that cue salience can be more important than cue reliability in determining what
42 features will control behaviour when multiple cues are available.

43 **Key words:** Pigeon, Category discrimination, Feature learning, Family
44 resemblance, Labelling, Salience

45

46 **General Introduction**

47

48 The ability to categorise natural objects is expected to be widespread across the
49 animal kingdom, since it permits a reduction in the amount of information an animal
50 must acquire about its environment in order to respond adaptively (Makino and
51 Jitsumori 2007). Since the pioneering experiments of Herrnstein and Loveland (1964)
52 birds have been shown to be capable of discriminating a wide range of categories,
53 with category members varying from simple artificially constructed stimuli (e.g. Lea
54 and Harrison 1978) to photographs of complex natural-language categories such as
55 trees (Herrnstein et al. 1976; for a review see Huber 2001).

56

57 A typical categorisation experiment involves pigeons learning to associate one
58 stimulus set or category with a food reward, when there is no single feature that is
59 necessary or sufficient for category membership. However it is not clear what pigeons
60 learn to associate with the food reward. It could be that the pigeon has elaborated a
61 concept corresponding to the category. Typically however it is assumed that
62 successful category discrimination does not necessarily imply that pigeons have a
63 concept corresponding to the experimenter-defined category (despite the terminology
64 of early experiments e.g. Herrnstein et al. 1976), or in the case of natural photographs,
65 the ability to generalise from images to the corresponding real objects (Bovet and
66 Vauclair 2000). But if pigeons are not using concepts to categorize complex stimuli,
67 how might they be doing it? There are two well studied possibilities. It could be that
68 they learn how to respond to one or more specific exemplars, with other stimuli being
69 categorised by a process of generalisation. Alternatively, it may be that they learn to
70 respond to a collectivity of features that are between them good enough predictors of

71 category membership, even if none of them is individually necessary or sufficient.
72 These possibilities are usually referred to respectively as exemplar learning and
73 learning by family resemblance or overall similarity.

74

75 It is often useful to compare the cognitive abilities of non-human animals and human
76 infants, because in both cases subjects have to complete tasks without the elaborate
77 language competence that facilitates so many cognitive performances in adult
78 humans. The present experiment draws on two results that have been established in
79 the field of infant categorization, in order to pose questions about categorization in
80 pigeons.

81

82 Both phenomena involve the use of single, highly predictive features within sets of
83 multidimensional stimuli. However, they are to some extent contradictory. On the
84 one hand, under conditions where adults and older children typically categorise items
85 according to a single stimulus dimension, infants have been found to group items
86 according to overall similarity or family resemblance across numerous stimulus
87 dimensions (Smith and Kemler 1977; Smith 1981). On the other hand, the provision
88 of a verbal label which is invariantly associated with the members of one category has
89 been found to facilitate children's category acquisition (Waxman and Markow, 1995),
90 even among infants too young to have functional speech (e.g. Balaban and Waxman,
91 1997); so in contrast to the control by family resemblance seen in spontaneous
92 categorization, the presence of a single reliable feature is found to facilitate category
93 acquisition.

94

95 The first of these findings, the tendency of younger children to categorize on the basis
96 of overall similarity, has supported a general arguments that such categorization must
97 require simpler cognitive mechanisms than using a single stimulus dimension (e.g.
98 Ashby et al. 1998; Pothos 2005). Unidimensional discrimination is assumed to
99 require the ability to verbalise a rule, something that a pre-verbal infant cannot do. In
100 accordance with this assumption, Couchman et al. (2010) found that, when they used
101 stimulus sets of a kind introduced by Kemler Nelson (1984), which can be
102 discriminated either on the basis of a single dimension or on a family resemblance
103 across three other dimensions, human participants categorised them unidimensionally
104 whereas two rhesus monkey subjects categorized them mainly by family resemblance.
105

106 However, it is not the case that non-human animals always categorize complex stimuli
107 by overall similarity rather than unidimensionally, or that unidimensional
108 categorization implies more complex cognitive processing (e.g. Lea and Wills 2008;
109 Wills et al. 2009). Although it is certainly possible for pigeons to use multiple
110 stimulus dimensions in making complex discriminations (e.g. Blough 1972;
111 Herbranson et al. 1999; Kirkpatrick-Steger and Wasserman 1996; Soto and
112 Wasserman 2010; von Fersen and Lea 1990), this is not necessarily what they do most
113 easily or spontaneously. Rather than categorizing in terms of overall similarity, which
114 would involve all available features, birds in categorization experiments with complex
115 stimuli often come under the control of just one or two features (e.g. Lea et al. 1993,
116 2006); and it would be absurd to suppose that pigeons can verbalise rules (cf. Lea and
117 Wills, 2008). Furthermore, the fact that infant categorization proceeds more easily in
118 the presence of a verbal label – which is really nothing but a single highly valid

119 stimulus dimension (Plunkett et al., 2008) – suggests that discriminating a single
120 dimension may be a cognitively simpler task than discriminating overall similarity.

121

122 The aim of the present study is to explore whether, under conditions where both
123 strategies would be equally effective, pigeons would solve categorization tasks by
124 using a single highly predictive feature, or by using the overall similarity across a
125 group of features of more modest individual predictiveness. However, previous results
126 have shown that when pigeons do make use of single features, they do not always
127 select the most predictive feature: a less valid but more salient dimension of stimulus
128 variation may acquire exclusive control over behaviour (Lea et al. 2009).
129 Accordingly, the experiments were designed to enable us to dissociate the effects of
130 cue salience from those of cue validity.

131

132 Von Fersen and Lea (1990) noted that categorization in pigeons can be investigated in
133 two different ways: by using natural photographs, videos and objects, which may be
134 referred to as an analytic approach, and by using artificially designed stimuli, the
135 synthetic approach. The advantage of using artificial stimuli is that structure and
136 feature content can be carefully controlled, but such control is usually at the expense
137 of the richness and detail associated with natural exemplars. The use of photographs
138 or videos enhances the verisimilitude of categorization experiments, because it
139 presumably mimics more closely the kind of discrimination that birds have to make in
140 the wild (though it cannot do so exactly, and it cannot necessarily be assumed that the
141 birds recognize the pictures as representations of the corresponding real objects). In
142 the present paper, we used both techniques, so as to gain the advantages of both, and
143 also to provide systematic replication of our main manipulation. Experiment 1 used

144 natural photographs (though with some artificial modification), while Experiment 2
145 used wholly artificial stimuli. In both experiments, the stimuli included a single
146 feature that was a perfectly reliable predictor of category membership; by analogy
147 with the work of Waxman and others cited above, we refer to this as the “label”. In
148 addition the stimuli included several other features that were individually imperfect
149 predictors but in combination provided enough information to allow perfect
150 discrimination. Once good performance had been achieved, probe trials were
151 introduced, in which the single, perfectly reliable cue and the remaining cues were put
152 in conflict with each other, allowing us to discern which cues were controlling
153 behaviour more strongly.

154

155 **Experiment 1**

156

157 The first experiment used photographs of everyday objects as stimuli. The two
158 categories were houses and cars, which differ along multiple stimulus dimensions. To
159 provide a single highly salient “label” cue, the roofs of the houses and the windows of
160 the cars were replaced with a standard uniform texture in a contrasting colour. If the
161 pigeons in this situation base their discrimination on the label alone, then when probe
162 stimuli are introduced in which the labels have been reversed between categories, it is
163 expected that probes containing the label previously associated with the positively
164 reinforced category (i.e. positive texture probes) will be treated as positive. If pigeons
165 respond positively instead to probes which contain the label originally associated with
166 the negatively reinforced category (positive picture probes), this would suggest they
167 are attending more to the overall similarity of exemplars. Although the stimuli were
168 based on natural photographs of objects, it was not required by the design that the

169 pigeons should recognize them as depictions of objects that they had seen;
170 photographs were used only to ensure that the stimulus sets had the kinds of variation
171 that are characteristic of functional categories in the everyday world.

172

173 **Materials and Methods**

174

175 *Subjects*

176 Twenty pigeons were used. They were acquired as discards from pigeon fanciers, and
177 so had visual experience of the world outside the laboratory. Some had previous
178 experience of the touchscreen arrangement used in the present experiment, but in
179 experiments with monochrome, geometric stimuli and a different training procedure
180 (Wills et al. 2009, Experiment 1a); the remainder were experimentally naive. The
181 pigeons were housed in an indoor aviary, measuring 2.2 m by 3.4 m by 2.4 m, and
182 given constant access to water and grit. Prior to testing, pigeons were held in
183 individual cages in which they had access to water and were released back into the
184 aviary when testing had finished for the day. All pigeons were maintained on a 12:12
185 hr light/dark cycle at 95% of their free-feeding weight.

186

187 *Apparatus*

188 Four identical operant chambers (internal size; 640 mm x 430 mm x 470 mm) were
189 used. Each consisted of a plywood box, with a 15-inch (39-cm) touch-monitor (Elo
190 Touchsystems Accutouch (resistive) or CarrollTouch (infra-red) model 1547L)
191 mounted in the front wall. The screen resolution was set to 1024 x 768 pixels. The
192 monitor was controlled by software written in Visual Basic using the Whisker control

193 system (Cardinal and Aitken 2001) running on a computer supplied by Quadvision
194 (Quadvision Ltd., Dorset, UK). Two food hoppers, positioned one on each side of the
195 main screen, could be used to deliver a 2:1 mixture of hemp seed and conditioner to
196 the pigeons, for 2.5 seconds. Each box had a webcam fitted into the side wall, 250
197 mm above the floor, allowing the pigeons' behaviour to be observed from outside the
198 test room using the imaging software ViewCommander (Internet Video and Imaging,
199 Ltd.). Each pigeon was assigned its own test chamber for all stages of the experiment:
200 six pigeons used resistive touchscreens while the remaining pigeons used infra-red
201 touchscreens.

202

203 *Stimulus Materials*

204 The images used were natural photographs comprising two stimulus sets; houses and
205 cars. There were 24 pictures of each type of object. Photographs were manipulated
206 using the GNU Image Manipulation Program (GIMP ver. 2.2). Each image was
207 isolated from the original setting, placed on a black background and then scaled to
208 ensure images were approximately the same size (160 x 120 pixels). To produce a
209 highly salient "label" feature, similar areas in the upper part of the objects were
210 selected and given a new artificial texture. For houses this was the roof, and for cars
211 the windows. Thus 12 of the houses had their roofs replaced with a leopard-skin
212 texture, and the other 12 had their roofs replaced by a blue "swimming pool" wave
213 texture. Similarly, 12 of the cars had all their windows replaced with the leopard-skin
214 texture and the other 12 had them replaced with the blue wave texture. Figure 1
215 shows examples of the stimuli. Fourteen of the pigeons were trained to discriminate
216 the houses with leopard-skin roofs from the cars with blue windows, and for these
217 pigeons probe stimuli consisted of houses with blue roofs and cars with leopard-skin

218 windows. For the remaining pigeons, the training stimuli were the houses with blue
219 roofs and the cars with leopard-skin windows, and the probe stimuli were the houses
220 with leopard-skin roofs and the cars with blue windows. Within each of these groups,
221 some of the pigeons were trained with houses as positive stimuli and the remainder
222 with cars as positive stimuli.

223

224 *General Procedure*

225 The pigeons were pre-trained using conventional methods to find food in the grain-
226 feeders, and to peck lighted discs on the touch-screen for food reinforcement. They
227 were then trained in a multi-stimulus discrimination procedure similar to that
228 described by Huber et al. (2005) and Wills et al (2009, Experiment 2b). Initially, they
229 were trained using this procedure to discriminate white filled hexagons (8mm across)
230 within a 25-mm diameter black circle, outlined in white, from a blank circle. Those
231 pigeons that mastered this discrimination proceeded to the task involving the house
232 and car stimuli.

233

234 In the multi-stimulus discrimination procedure, each session started when the house-
235 lights in the box came on, and a white disc of diameter 4cm (the observing key) was
236 displayed on the screen. When the pigeon pecked the observing key it disappeared, to
237 be replaced by an array of photographs. The array consisted of eight cells, arranged as
238 shown in Figure 2. In order to ‘open’ a cell, pigeons had to peck it twice in rapid
239 succession, causing the image to disappear. If the opened cell was positive, a side key
240 appeared to one side of the array, in the nearer of the two locations shown in Fig. 2
241 (arbitrarily, the left side key was used when the middle top cell was opened, and the
242 right side key when the middle bottom cell was). The pigeons were required to peck

243 the side key to activate the food hopper on that side for 2.5 seconds (Figure 2). If a
244 negative cell was pecked this was recorded as a miss; the image disappeared but no
245 side key appeared and the screen became unresponsive for 2.5 seconds. If a pigeon
246 pecked a blank cell this was recorded but there were no scheduled consequences.
247 When all the positive cells had been opened, the array disappeared, to be replaced
248 after an inter-trial interval that varied from 1 to 5 s by the observing key. Sessions
249 consisted of six or seven arrays depending on the stage of the experiment.

250

251 *Training*

252 Three pigeons failed to complete pre-training and were dropped from the experiment.
253 Of the remaining 17 pigeons, 11 were trained using the houses with leopard-skin roofs
254 and the cars with blue windows; of these, five were assigned houses as the positive
255 stimulus, and six were car-positive. The remaining six pigeons were trained using the
256 houses with blue roofs and the cars with leopard-skin windows, and of these three
257 were house-positive and three were car-positive. Arrays were made up of four cars
258 and four houses, pseudo-randomly arranged, with the constraints that stimuli from the
259 same set were never presented in the same place in more than three consecutive arrays
260 and that no more than three stimuli from the same set were placed next to each other
261 or reinforced on the same side. Each training session contained six arrays, so that each
262 of the 12 positive and negative stimuli was seen twice within a session.
263 Discrimination during the training phase was determined using the ρ statistic of
264 Herrnstein et al. (1976), which was used to measure the overlap between stimulus sets
265 in terms of the order in which the cells were opened. A ρ value of 0.5 indicates
266 random responding, whereas a value of 1 indicates perfect discrimination (i.e. that all
267 the positive stimuli have been opened before any of the negative stimuli). After each

268 pigeon had reached a criterion of a p value of 0.8 or more in all six arrays in a session,
269 it was exposed to probe sessions.

270

271 *Probe Sessions*

272 Probe sessions consisted of four training arrays, alternated with three probe arrays.

273 Each session began and ended with a training array. A probe array contained four

274 training stimuli, two from each stimulus set, and four probe stimuli in which the

275 artificial manipulations were reversed. Two of the probes in each array, one from each

276 stimulus set, were assigned positive reinforcement contingencies and the other two

277 were assigned negative contingencies. Probes that were designated positive the first

278 time they were displayed were not reinforced when next shown. Pigeons were

279 required to open the two positive training cells and all four probe cells in order to

280 complete a probe array. Each pigeon received two probe sessions, which were

281 alternated with training sessions, to ensure pigeons recovered their original training

282 performance. Following the procedure of Wills et al. (2009, Experiment 2b),

283 responses to probe stimuli were categorised according to the order in which the

284 pigeons opened them within the array. A probe that was one of the first four stimuli

285 opened in an array was considered to have been treated as positive, and a probe that

286 was one of the last four stimuli opened was considered to have been treated as

287 negative.

288

289 **Results**

290

291 The pigeons reached criterion in a median of 5 training sessions (range 2-8 sessions).
292 This corresponds to the presentation of a median of 30 arrays, i.e. 240 stimuli. There
293 were no significant differences in the rate of acquisition between the house-positive
294 and the car-positive groups, or the leopard-skin positive and blue-wave positive
295 groups, nor was the interaction between the positive picture and the positive texture
296 significant.

297

298 In probe sessions, responses to training stimuli continued at the same high level of
299 accuracy. Different pigeons behaved differently towards the probe stimuli. Probes
300 included either a positive picture (house or car, depending on which was positive
301 during training) and a negative texture (blue wave or leopard-skin), or a negative
302 picture and a positive texture. Figure 3 shows, for each pigeon, the proportion of
303 probe stimuli that were treated in accordance with the texture cue present (that is,
304 responded to as positive if they included the positive texture or as negative if they
305 included the negative texture). In all, 7 of the 17 pigeons (Ro, Ma, Fr, Sn, Su, Ba and
306 Ze) responded to more than half the probes in accordance with the texture cue, 9
307 pigeons (Io, Le, Ri, Ru, Rs, Ti, Ot, Ry, and Se) responded to more than half the
308 probes in accordance with the picture cue, and the remaining pigeon (At) treated both
309 kinds of probes equally. Chi-square tests were used to determine whether the
310 proportions of probe stimuli treated in accordance with the texture or picture cue
311 differed significantly from 0.5 for individual pigeons. Four pigeons responded to the
312 probes in accordance with the texture cue on significantly more than half the trials
313 (Ro, Fr, Sn and Su; χ^2 values of 15.04, 14.09, 8.52 and 4.34, $df=1$, $p<0.0001$, 0.0001,
314 0.01 and 0.05 respectively). Three pigeons responded to the probes in accordance
315 with the picture cue on significantly more than half the trials (Io, Ri, and Ry; χ^2 values

316 of 15.04, 9.38 and 7.04, $df=1$, $p<0.0001$, 0.01 and 0.01 respectively). The sum of the
317 individual χ^2 values across the pigeons was 83.05; by virtue of the additive property of
318 the chi-square distribution (Weatherburn, 1957, p. 177), this can be tested as a χ^2 value
319 against the sum of their degrees of freedom. The resulting significance level is <
320 0.0001, showing that despite the fact that not all results were in the same direction, we
321 can reject the null hypothesis that all pigeons had a 50% chance of treating any probe
322 in accordance with the picture cue.

323

324

325 Discussion

326

327 The rapid learning shown by all the pigeons confirms the conclusion of Huber et al.
328 (2005) and Wills et al (2009) that multi-stimulus training methods are an efficient
329 means of establishing complex pattern discriminations. It is a property of such
330 methods, as they have been implemented previously and in the present experiment
331 that, within an array, a correct response makes the remainder of the task more difficult
332 (because it reduces the ratio of positive to negative stimuli remaining) and an
333 incorrect response makes it easier. This makes the task of predicting the probability
334 of a correct response mathematically more difficult, but is not a concern when as here,
335 the intention is to compare the probabilities of choosing different probe stimuli that
336 are presented at the same frequency.

337

338 Responses to probe stimuli were reinforced non-differentially. This was because
339 substantial numbers of probe trials were required, and the alternative, non-

340 reinforcement, could have led to the pigeons learning that all probe stimuli were
341 essentially negative. Non-differential reinforcement carries the alternative risk that
342 subjects will come to behave at random towards probes. However in the present
343 experiment, sustained non-random behaviour towards probes was observed in most
344 pigeons, showing that non-differential reinforcement had no major effect.

345

346 The pigeons were almost equally split between showing greater control by the picture
347 and showing greater control by the texture (the “label” cue). This was not a result of
348 random behaviour, but of systematic behaviour that differed between individuals.
349 Seven of the 17 pigeons showed significant differences in their responses to probes
350 containing the positive picture rather than the positive texture, but in three of these it
351 was the picture cue that controlled probe responding, and in the other four it was the
352 texture cue. As regards the remaining pigeons, given that performance on training
353 stimuli remained highly accurate throughout the probe sessions, the most plausible
354 conclusion is that they were under the control of both the picture and the texture. The
355 distribution of control by the picture or texture was not even between the groups, and
356 in particular all four of the pigeons that showed a significant trend to texture control
357 had the leopard-skin texture as positive. If the leopard-skin was more salient than the
358 blue-wave texture for the pigeons, this could be accounted for as a feature-positive
359 effect (Jenkins and Sainsbury 1970), but in the absence of independent evidence about
360 relative salience, this can only be a speculation. Furthermore one of the pigeons that
361 showed a significant trend towards control by the picture cue (Io) also had the
362 leopard-skin texture as positive, weakening the argument that the overall pattern of
363 results could be due to high salience of the leopard-skin cue.

364

365 Because the pictures were natural photographs, we cannot easily specify which
366 features were supporting discrimination between them, but inevitably the features
367 available will have varied somewhat between instances. We therefore interpret
368 discrimination based on the picture cue as categorization by overall similarity or
369 family resemblance. The texture cue, by contrast, was (to the human eye at least)
370 highly salient, and it was consistent across the training and probe stimulus sets.
371 Discrimination of the texture cue can therefore be regarded as unidimensional
372 categorization. Nevertheless, it only exerted dominant control over behaviour in
373 probe trials for four pigeons.

374

375 It is possible that the pigeons learned to discriminate the categories by learning each
376 exemplar of a house or car separately. Previous results make this unlikely given the
377 number of exemplars used: an experiment by Bhatt, cited by Wasserman and Bhatt
378 1992, showed that pigeons switch from exemplar control to featural control in
379 category discriminations where the number of exemplars rises above about six. In
380 any case it would not matter to the present experiment, whose point was to investigate
381 whether the pigeons would come under the control of the single feature provided by
382 the texture cue or the multiple features provided by the picture cue: if the pigeons
383 discriminated houses from cars on the basis of exemplars, that would involve the
384 learning of even more different features than doing so by extracting a few general
385 features that were positively but not perfectly correlated with reinforcement.

386

387 The roughly equal distribution of subjects between control by the texture cue and
388 control by the picture cue is an unexpected result, given that pigeons have a tendency
389 to be dominated by single features (Lea et al. 1993, 2006). However, while it is true

390 that birds rarely use all the features available, they can certainly be trained to use
391 more than one (e.g. von Fersen and Lea 1990; Lea et al. 2006). Indeed, some of the
392 pigeons used in the present experiment had been trained in Experiment 1a of Wills et
393 al. (2009), in which they had to learn to discriminate several different features
394 presented on separate trials (though the stimuli and the procedure were very different
395 from those of the present experiment). In the generalisation tests of that experiment,
396 when the features were combined, the behaviour of one of the pigeons that was re-
397 used here (Io) was shown to be under the control of multiple features. However,
398 bringing behaviour under the control of multiple stimulus dimensions often requires
399 special training procedures (as in von Fersen and Lea 1990) or extended training,
400 whereas the pigeons in the present experiment learned very quickly.

401

402 Because the picture cues were based on natural photographs, it remains an assumption
403 that discrimination of the pictures was on the basis of overall similarity. It is possible
404 that within the pictures there were other highly predictive features apart from the one
405 manipulated, which were consistent within each category and which were salient for
406 the pigeons. All the car stimuli, for example, contain wheels and so pigeons might
407 have learnt the discrimination on the basis of the presence or absence of silver
408 ellipses. We did our best to ensure there was no such single predictive feature for
409 each category, for example the angle at which the photographs of the stimuli were
410 taken was varied, as was the colour of the cars (so that although all the cars had
411 wheels, those wheels themselves formed a polymorphous set), but there can be no
412 certainty that such attempts will succeed. Conversely, although the label features
413 were much more consistent, they did have some variability: for example, the shapes of
414 the house roofs and car windows to which the textures were applied varied between

415 instances. To avoid these problems while investigating the question of whether birds
416 are more likely to rely on single features or family resemblance, in Experiment 2 we
417 took the alternative approach of using artificial compound stimuli, and so bringing
418 feature content under control.

419

420 **Experiment 2**

421

422 Experiment 2 was formally very similar to Experiment 1, but used artificial
423 multidimensional stimuli. The stimuli in the present experiment were made up of four
424 spatially separated elements, each of which constituted a stimulus dimension; they
425 were based on those used by Wills et al. (2009, Experiment 1a) and Lea et al. (2009).
426 The elements are referred to as the Annulus, the Bar, the Checks and the Diamond,
427 and examples are shown in Figure 4. Each element could be used in four graded
428 versions, two of them positively and two negatively correlated with the occurrence of
429 reinforcement. The stimulus sets had exactly the same formal structure as those used
430 by Kemler Nelson (1984) with children and Couchman et al (2010) with rhesus
431 monkeys, but the stimulus dimensions used were different.

432

433 Corresponding to the reliable, salient, texture cue used as a label in Experiment 1, one
434 of the dimensions was designated as a label cue for each pigeon. This dimension was
435 used only in its extreme versions, and one of these always occurred in the presence of
436 reinforcement while the other one never did. The remaining three dimensions were
437 used to construct a two-out-of-three artificial polymorphous stimulus of the sort
438 introduced by Dennis, Hampton and Lea (1973) and used with pigeons by Lea and

439 Harrison (1978): that is, whenever at least two of the three dimensions took its
440 positive value, the stimulus as a whole was designated as positive. For these
441 dimensions, all four versions were used. These three dimensions collectively
442 corresponded to the picture feature in Experiment 1: if a pigeon was to discriminate
443 on the basis of these cues, it would have to come under the control of the overall
444 similarity of the stimuli to an ideal positive form if it was to achieve 100% correct
445 performance, since each individual dimension within this set was imperfectly
446 correlated with reinforcement. These artificial stimulus sets had several additional
447 advantages. First, the spatial separation of the stimulus elements ensured that the
448 dimensions of the stimuli could be manipulated entirely independently. Second, the
449 four dimensions could be used in a balanced way, with each dimension being used to
450 provide the label (reliable) feature for some pigeons. Thirdly, within the set of
451 features used to create the polymorphous concept, it was possible to assess
452 empirically whether all three features were controlling behaviour, and therefore
453 whether it is appropriate to describe the pigeons as coming under the control of
454 overall similarity. Finally, on the basis of results with similar stimuli (Lea et al 2009,
455 Wills et al 2009) we had reason to think that the saliences of the feature differences
456 used on the four dimensions of the stimuli were comparable.

457

458 **Materials and Methods**

459 *Subjects*

460 Sixteen pigeons were used in this experiment. None of them had previous experience
461 of this kind of discrimination task. They were maintained under the same conditions
462 as the pigeons used in Experiment 1.

463

464 *Apparatus*

465 The same apparatus was used as in Experiment 1, except that only infra-red
466 touchscreens were used.

467

468 *Stimuli and Experimental design*

469 The stimuli were modified from those used in Wills et al. (2009, Experiment 1a) and
470 Lea et al. (2009). Examples are shown in Figure 4. All stimuli consisted of a square
471 array of four elements. Elements of the same type were always placed in the same
472 location. The element types were an annulus (A), a horizontal bar (B), a square
473 chequerboard (C), and a diamond shape made up of equal-width stripes (D). There
474 were four versions of each element, designated as the X, x, y, and Y forms. All
475 versions of all elements were placed on a black background, and contained the same
476 number of white pixels (within 2%). The specifications of the four forms of each
477 element are listed in Table 1, and they are illustrated in Figure 4. Note that because of
478 the constraint that all elements should have the same area, some elements varied on
479 two dimensions simultaneously: for example, when the Annulus was made smaller, it
480 was also made wider. With four versions of each of four elements, there were 256
481 possible stimuli, but not all pigeons experienced all stimuli. All versions of all
482 elements fitted within a 60 x 60 pixel square with some black border, so that the entire
483 stimulus including borders fitted within a 120 x 120 pixel square; at a pigeon's typical
484 pre-peck viewing distance, 120 pixels subtended about 25° of arc, and would thus fit
485 within the extent of the pigeon's frontal, binocular viewing field (Martin and Young,
486 1983).

487

488 For each pigeon, one of the four dimensions was designated as Reliable, and either the
489 X or the Y value of it was designated as positive; the less extreme (x and y) values of
490 the Reliable dimension were not used either in training or in probe stimuli. The
491 remaining three dimensions were designated as Unreliable, and either all their X and x
492 values, or all their Y and y values, were designated as positive. Positive training
493 stimuli always included the extreme positive value of the Reliable dimension, and
494 positive values (either extreme or less extreme) of at least two of the Unreliable
495 dimensions. Negative training stimuli always included the extreme negative value of
496 the Reliable dimension, and negative values (either extreme or less extreme) of at
497 least two of the Unreliable dimensions. As an example, consider Pigeon Mo, for
498 which the Reliable dimension was the Annulus and the X values of both the Reliable
499 and Unreliable dimensions were designated as positive (see Table 2). For this pigeon,
500 positive stimuli always contained the X value of the Annulus, and at least two of the
501 other three dimensions (Bar, Chequerboard and Diamond) in either their X or their x
502 forms. The negative stimuli always contained the Y form of the Annulus, and either
503 the y or the Y form of at least two of the other three dimensions. There were 32
504 stimuli in each of the positive and negative categories. These categories could be
505 discriminated perfectly in either of two ways (or by a mixture of them). The pigeon
506 could use the Reliable dimension alone, and ignore the three Unreliable dimensions.
507 Alternatively, it could ignore the Reliable dimension, and respond on the basis of the
508 preponderance of values of the three Unreliable dimensions. Each Unreliable
509 dimension took one of its negative values in a quarter of the positive training stimuli,
510 and a positive value in a quarter of the negative training stimuli, so each Unreliable
511 dimension considered on its own can be described as being 75% valid during training;
512 their collectivity, however, was 100% valid. The Reliable cue thus served as a label,

513 while the Unreliable cues defined a 2-out-of-3 polymorphous stimulus set. Stimuli
514 that included the positive value of the Reliable dimension with negative values of at
515 least two of the Unreliable dimensions, or the negative value of the Reliable
516 dimensions with positive values of at least two of the Unreliable dimensions, were
517 ambiguous, and were not used in training. There were 64 such ambiguous stimuli for
518 each pigeon, and a selection of these were used as probes, in tests conducted after
519 training was complete.

520

521 The way the categories were used was varied between pigeons so as to balance the use
522 of the different dimensions, as shown in Table 2. Each dimension was assigned as
523 Reliable for four of the pigeons. For two of these, the X value of the Reliable
524 dimension was assigned as positive, and for the other two its Y value was assigned as
525 positive. For one of the pigeons for which each value of the Reliable dimension was
526 assigned as positive, the X and x values of the Unreliable dimensions were assigned
527 as positive, and for the other one the Y and y values of the Unreliable dimensions
528 were assigned as positive.

529

530 To reduce the risk that the pigeons would learn the contingencies associated with
531 probe stimuli, only 36 probe trials were given to each pigeon, so not all the 64
532 possible probe stimuli were used. The stimuli used as probes are summarised in Table
533 3. Four of these stimuli were used 6 times each, so as to provide a strong test of the
534 basic question of whether the pigeons' behaviour was governed by the Reliable or the
535 Unreliable dimensions. These stimuli involved:
536 The positive value of the Reliable dimension, combined with the extreme negative
537 values of all three Unreliable dimensions

538 The negative value of the Reliable dimension, combined with the extreme positive
539 values of all three Unreliable dimensions

540 The positive value of the Reliable dimension, combined with the less extreme
541 negative values of all three Unreliable dimensions

542 The negative value of the Reliable dimension, combined with the less extreme
543 positive values of all three Unreliable dimensions.

544

545 In addition six other probe stimuli, as indicated in Table 3, were used twice each.

546 These stimuli involved the positive value of the Reliable dimension and the extreme

547 positive value of one of the Unreliable dimensions, and the extreme negative values of

548 the other two Unreliable dimensions; or the negative value of the Reliable dimension

549 and the extreme negative value of one of the Unreliable dimensions, and the extreme

550 positive values of the other two Unreliable dimensions. Each of the Unreliable

551 dimensions was paired with the Reliable dimension in an equal number of probe

552 stimuli. These stimuli allowed a test of which of the Unreliable dimensions was

553 controlling behaviour. Each probe stimulus was associated an equal number of times

554 with the reinforcement contingencies appropriate to positive and negative stimuli.

555

556 *Procedure*

557 The pigeons were pretrained as in Experiment 1. They were then trained, using the

558 same procedure as in Experiment 1, to discriminate between two positive and negative

559 training categories. Training was continued for a maximum of 20 sessions, but was

560 stopped earlier if a pigeon reached a criterion of a p value of 0.8 or more in all six

561 arrays in a session. Three probe sessions were then given. As in Experiment 1,

562 probe sessions consisted of seven arrays: four training arrays, alternated with three

563 probe arrays. Each session began and ended with a training array. A probe array
564 contained four training stimuli and four probe stimuli, of which two were associated
565 with the reinforcement contingencies appropriate to positive stimuli and two with
566 those appropriate to negative stimuli. Responses to probe stimuli were classified as
567 positive if they occurred within the first four stimulus cells pecked in an array, and as
568 negative otherwise.

569

570 Results

571 Ten of the 16 pigeons reached criterion before their 20th training session; the number
572 of sessions required ranged from 9 to 18 (median, including the pigeons that did not
573 reach criterion, 17). The other six pigeons were showing no obvious further
574 improvement in performance after 20 sessions, though all but one of them was
575 showing p values consistently above 0.5; the performance of the remaining pigeon
576 was erratic.

577

578 For comparison with Fig. 3, Fig. 5 shows the proportions of probe trials in which the
579 response was correctly predicted by the Reliable stimulus dimension rather than the
580 preponderance of the Unreliable dimensions. For 10 of the 16 pigeons, this
581 proportion was greater than 0.5, and over the group as a whole the deviation from 0.5
582 was significant (1-sample, 2-tailed Wilcoxon test, $T = 23.5$, $N=16$, $p<0.05$). For each
583 of these pigeons individually, the proportion deviated from 0.5 significantly (2-tailed
584 binomial test). For one of the six pigeons where the proportion was below 0.5 (Ba),
585 the deviation was significant (2-tailed binomial test, $p < 0.001$). The pigeons that did
586 not respond according to the Reliable stimulus on the probe trials had taken

587 significantly longer to reach criterion than those that did (2-tailed Mann-Whitney test,
588 $U = 10.5, N_1 = 6, N_2 = 10, p = 0.03$).

589

590 Figure 6 shows, for each pigeon, the proportions of probe trials in which the response
591 was correctly predicted by each dimension of the stimulus, whether it was reliable or
592 unreliable for that pigeon. In this figure, any proportion above 0.67 or below 0.33
593 would be significantly different from 0.5 on an individual test, though the number of
594 data points involved and the correlations between dimension values mean that simple
595 significance tests cannot be interpreted directly. It can be seen that for each of the
596 pigeons where the unreliable dimensions predicted the response to probe trials
597 markedly better than the reliable dimension (Mo, Bn and Jk), there was one of the
598 unreliable dimensions that predicted response particularly well (Checks for Mo and
599 Jk, and Annulus for Bn). The same trend can be seen more weakly in two of the
600 pigeons where the dominance of the unreliable dimensions was more marginal, Mr
601 and Sa, where Checks and Annulus respectively seemed to have majority control over
602 behaviour. The remaining pigeon, Cr, showed apparently random behaviour towards
603 probe stimuli. For the pigeons where behaviour towards the probe stimuli was
604 dominated but not 100% predicted by the reliable dimension, there was no evidence
605 that individual unreliable dimensions contributed disproportionately to controlling
606 behaviour.

607

608 Because the values of the unreliable dimensions used in the training stimuli varied, it
609 was possible to examine the relative control over behaviour of these dimensions under
610 training as well as probe conditions. Table 4 shows the results of such an analysis,
611 carried out on the training stimuli that were used within the probe sessions so as to

612 maximise comparability with the probe stimulus results shown in Figures 5 and 6.
613 There were 96 such trials, so any proportion greater than 0.58 or less than 0.42 would
614 be significantly different from 0.5 in a single analysis, though the same cautions about
615 the number of tests and the non-independence of the dimensions must be applied as
616 with Figure 6. These data confirm the dominance of the Annulus dimension for
617 pigeons Sa and Ba, and weakly support the dominance of Checks for Mr, but do not
618 support the dominance of Checks for Mo or Jk. On the other hand they suggest that
619 Cr may have been somewhat under the control of the Bar dimension. It is notable
620 (and logically inevitable) that in those pigeons where one of the unreliable dimensions
621 exerted disproportionate control over behaviour, overall discrimination performance
622 was poorer than in the other pigeons.

623

624 Discussion

625 In Experiment 1, we found that the texture and picture cues (which we interpret
626 respectively as a single reliable dimension, and a collectivity of presumably unreliable
627 cues) were about equally likely to control behaviour, and in many cases individual
628 pigeons probably came under the control of both types of cue. However the two kinds
629 of cues differed in nature. In Experiment 2, where the same cues were used as
630 reliable and unreliable dimensions in a fully balanced way, we found a clear trend for
631 reliable dimensions rather than a collectivity of unreliable dimensions to control
632 behaviour, even though either of them could have enabled the pigeons to perform with
633 perfect accuracy. There were some individuals in which the unreliable dimensions
634 exerted dominant control over behaviour, but in at least some cases it appears that this
635 was not because the collectivity of those dimensions was enabling perfect
636 discrimination, but rather because one of those dimensions was controlling behaviour

637 to the exclusion of the others and of the reliable dimension, with the result that the
638 pigeon's discrimination accuracy was limited. The pigeons that did not come under
639 the control of the Reliable dimension were slower to reach criterion (indeed, 4 out of
640 these 6 pigeons did not reach it within 20 sessions, as against only 2 of the 10 pigeons
641 that did show Reliable dimension control), and if they were coming under the
642 exclusive control of a less valid dimension, this is a more or less inevitable result.
643 Our results thus differ from those obtained by Couchman et al (2010), using stimulus
644 sets that were structurally identical to those used here, though with different elements:
645 Compared with humans exposed to the same task, Couchman et al's rhesus monkeys
646 always showed a greater tendency to categorise novel stimuli in terms of overall
647 similarity rather than in terms of the label dimension, and the authors concluded that
648 this was because the monkeys could not use verbal rules to categorize complex
649 stimuli. Since we do not believe that pigeons use verbal rules, yet in our experiment
650 they showed a clear tendency to use the label dimension rather than family
651 resemblance, we conclude that categorization by family resemblance is not an
652 inevitable consequence of failing to use verbal rules. We therefore also conclude that
653 unidimensional categorization is not a reliable sign that a verbal rule is being used, in
654 accordance with the conclusions of Lea and Wills (2008) and Wills et al (2009). The
655 difference between our results and those of Couchman et al may be due to the subject
656 species, or it may be due to differences in the details of the stimuli involved, a matter
657 to which we return in the General Discussion.

658

659 Part of the reason why clear dominance of the reliable dimension was found in the
660 present experiment is that with the values on them used in the present experiment, the
661 salience of the four dimensions seems to have been, if not equal, at least comparable,

662 as we predicted on the basis of our previous results with similar stimuli (Wills et al
663 2009, Lea et al 2009). As Table 4 shows, when used as the reliable dimension, all
664 four dimensions achieved dominant control over behaviour in at least one of the four
665 pigeons for which they were Reliable; assuming that salience is reflected in the
666 number of pigeons for which this was so, the order for salience was Checks >
667 Annulus> Diamond > Bar. The data on dominance by Unreliable dimensions (also
668 shown in Table 4) confirm this pattern, with Checks and Annulus being the only
669 dimensions to achieve dominant control when unreliable. Acquisition data also show
670 that the pigeons for which these dimensions were Reliable tended to learn more
671 quickly than the others.

672

673 Learning in Experiment 2 was substantially slower than in Experiment 1. In
674 Experiment 1, pigeons took a median of 5 training sessions (range 2-8 sessions) to
675 reach criterion; in Experiment 2, the median number of sessions to the same criterion
676 was 12, with the number required ranging from 3 to over 20. This difference is
677 consistent with the fact that in several studies where polymorphous concept
678 acquisition has been slow or incomplete artificial stimuli have been used (e.g. Lea et
679 al., 1993, 2006), whereas otherwise quite similar studies using natural photographic
680 stimuli have found more rapid acquisition (e.g. von Fersen and Lea, 1990). It may be
681 that there is something about abstract geometric stimuli that makes it hard for pigeons
682 to learn about them. The present results do however rule out one explanation for the
683 ease of learning discriminations based on natural photographs, which is the possibility
684 that natural stimuli contain artefacts that enable the subjects to make an easy single-
685 dimensional discrimination, whereas the more fully controlled artificial stimuli do not.

686 In Experiment 2, all the discriminations could have been solved by the use of a single
687 dimension of demonstrated salience, and yet they were not easy for the pigeons.

688

689 **General Discussion**

690 Both of the present experiments examined the effect on pigeons' category
691 discrimination of providing a single, salient dimension, in effect a category label,
692 alongside more variable information that was nonetheless sufficient when taken as a
693 whole to support perfect discrimination. In Experiment 1, the label was a distinctive
694 texture, applied to parts of natural photographs. In Experiment 2, it was one of four
695 artificial geometric elements, distinguished by the fact that it had 100% validity as a
696 cue to reward, whereas the other elements only had 75% validity individually, though
697 collectively they were fully valid. In Experiment 1, some pigeons clearly based their
698 discrimination on the label, but slightly more of the pigeons used the pictorial
699 information instead. In Experiment 2, on the other hand, almost all the pigeons used
700 the label (the Reliable feature); of the small number that did not, most came under the
701 control of just one of the Unreliable features, and were thereby prevented from
702 achieving accurate discrimination.

703

704 The results of Experiment 1 demonstrate a tendency that is evident in the literature on
705 complex visual discriminations in pigeons; that detailed photographic material can
706 serve as highly effective stimuli in such discriminations, competing in effectiveness
707 with simple unidimensional cues on highly salient dimensions such as colour. On the
708 other hand, taking the results of the two experiments together, they do not support the
709 idea that pigeons discriminate photographs readily because the birds have a strong
710 tendency to integrate the wealth of different and possibly unreliable cues that pictures

711 contain. Rather, the results support the conclusions of Lea and Wills (2008), Wills et
712 al. (2009) and Lea et al. (2009), that pigeons have a strong tendency to come under
713 the control of single salient cues when exposed to complex discrimination tasks.
714 What photographs offer is a wide range of strong cues, so there is a good chance that
715 any pigeon will find one that is salient for it – salience being presumably, in part, a
716 function of the individual’s previous experience. The carefully balanced design used
717 in Experiment 2 enables us to conclude that, other things being equal, if there is a
718 fully reliable cue available it is likely that it will achieve dominant control over
719 behaviour. Nonetheless, it is not certain that this will happen; it is possible for more
720 salient but less reliable cues to dominate, resulting in imperfect discrimination. The
721 extreme case of this is the occurrence of position habits in experiments where animals
722 have to make spatial choices in a discrimination task; by the design of the experiment,
723 position is normally a cue that carries no information at all, but it can still achieve
724 dominant control over behaviour that continues despite evidence that an animal has
725 detected the truly predictive contingencies (e.g. Mahut 1954).

726

727 Despite the strong tendency for pigeons to come under the control of single
728 dimensions, it is clearly not the case that they cannot be controlled by more than one
729 dimension. Nor is it the case that control by multiple dimensions only occurs when it
730 is essential for perfect discrimination, as in a compound discrimination. These
731 generalizations were again confirmed in the present experiments: In Experiment 1, it
732 appeared that a majority of the pigeons came under the control of both the picture and
733 the texture, and in Experiment 2 at least a few of the pigeons showed evidence of
734 control by more than one of the unreliable dimensions.

735

736 While control by multiple dimensions is not impossible, it seems from the present
737 results, and previous data, that it is not the natural tendency for pigeons. It may be
738 that it is easier for primates, and this would be one account of the difference between
739 our results and those of Couchman et al (2010) with rhesus monkeys. If
740 multidimensional control does not come naturally to pigeons, tasks that require it,
741 such as polymorphous discriminations, are likely to be difficult for pigeons or other
742 birds to learn, and previous data support this position (e.g. Lea and Harrison 1978;
743 Lea et al. 2006). Lea et al. concluded that the reason was a limitation of attention: any
744 task that requires the processing of more than one stimulus dimension is inherently
745 difficult for a bird. The present data are consistent with that position.

746

747 It is possible that the difficulty of attending to multiple dimensions is exacerbated
748 when the dimensions are spatially separated, as in the present experiments. It is
749 notable that in the stimuli used by Couchman et al (2010), who obtained
750 categorization by overall similarity from rhesus monkeys, the four elements were
751 spatially contiguous, and this could be an alternative account of the difference
752 between their results and ours. Spatial grouping does affect pigeons: for example,
753 Sainsbury (1971) found that the feature positive effect in pigeons, which depends on
754 the elements of a stimulus being processed separately, was attenuated when the
755 elements were grouped more closely together. It is also possible that some kinds of
756 stimulus dimension are more separable than others for cognitive rather than perceptual
757 reasons (Soto and Wasserman 2010). However it is not a foregone conclusion that
758 spatial separation or cognitive compatibility will lead to a greater tendency towards
759 categorization by overall similarity: they could instead act to mitigate confusion

760 between the dimensions. This is a matter that would repay experimental
761 investigation.

762

763 The task used in the present Experiment 2 had many points in common with a
764 standard *m-out-of-n* artificial polymorphous discrimination task. The essential
765 difference was that there was an additional, completely reliable stimulus dimension (a
766 label in the terms used by Waxman and Markow, 1995), whereas in a standard
767 polymorphous task, all dimensions are equally unreliable. It is therefore not
768 surprising that the task was learned relatively quickly, especially when the complex
769 and abstract nature of the stimuli is taken into account. The speed of learning may
770 have been partly due to the multiple simultaneous presentation procedure, which is
771 known to produce faster learning than a simple go/no-go task (Huber et al. 2005,
772 Wills et al. 2009), and it would be interesting to try a standard polymorphous
773 discrimination using this procedure. However the present design does raise an
774 intriguing possibility. We normally think of the different dimensions of a stimulus as
775 competing for a subject's limited capacity for attention, and this is the basis of most
776 attention-based theories of discrimination learning, e.g. Sutherland and Mackintosh
777 (1971). Indeed, earlier attention-based theories assumed that animals could only
778 attend to a single stimulus dimension at once (e.g. Krechevsky, 1932). Our results are
779 certainly consistent with the idea that animals' learning of complex stimuli is limited
780 by their attentional capacity. It is conceivable, however, that a highly reliable
781 dimension could act to inform an animal that a task can be learned, and this might
782 heighten attention to other dimensions rather than diminishing it, especially if the
783 reliable dimension was removed once learning had taken place; something of this sort
784 might underlie the "labeling" phenomenon as it occurs in young children (Waxman

785 and Markow, 1995; Balaban and Waxman, 1997; Plunkett et al., 2008). This
786 possibility gives more potential empirical bite to limited-attention theories, which can
787 seem to add little to the empirical generalization that multiple-dimension
788 discriminations are difficult. It could perhaps be investigated by exploring the
789 mechanisms of attention in more detail by comparing the amount learned about
790 unreliable stimulus dimensions in the presence or absence of more reliable cues.

791

792 **Acknowledgements**

793 Experiment 1 was carried out by Beth Nicholls in partial fulfilment of the
794 requirements for the MSc degree in Animal Behaviour of the University of Exeter.
795 The experimental work described here was supported by the New and Emerging
796 Science and Technologies activity of the European Community Framework
797 Programme 6, under grant no. 516542, “From Associations to Rules in the
798 Development of Concepts” (FAR). We are grateful to Lisa Leaver, Louise Millar,
799 Andy Wills and John Endler for discussion, and to Denis Mareschal and Caspar
800 Addyman (Birkbeck, University of London) for supplying the photographs used as
801 stimuli in Experiment 1. The experimental work was carried out within the terms of
802 the United Kingdom Animals (Scientific Procedures) Act (1986).

803

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890

891 Table 1. Experiment 2: The four forms of each dimension of the stimuli

Element	dimension of variation	X value	x value	y value	Y value
Annulus	width:radius ratio	5:1	4:1	2:1	1.5:1
Bar	length:width ratio	7.5:1	6:1	3.3:1	2:1
Chequerboard	number of elements	2 x 2	4 x 4	6 x 6	8 x 8
Diamond	orientation of stripes	vertical	60°	30°	horizontal

892

893

894 Table 2

895 Experiment 2: Use of stimulus dimensions in training stimuli for each pigeon

Pigeon	Reliable dimension (label)	Positive value	Unreliable dimensions	Positive values
Mo	Annulus	X	Bar, Chequerboard, Diamond	X, x
Ct	Annulus	X	Bar, Chequerboard, Diamond	Y, y
Ch	Annulus	Y	Bar, Chequerboard, Diamond	X, x
Kc	Annulus	Y	Bar, Chequerboard, Diamond	Y, y
Bn	Bar	X	Annulus, Chequerboard, Diamond	X, x
Mr	Bar	X	Annulus, Chequerboard, Diamond	Y, y
Sa	Bar	Y	Annulus, Chequerboard, Diamond	X, x
Cu	Bar	Y	Annulus, Chequerboard, Diamond	Y, y
Rg	Chequerboard	X	Annulus, Bar, Diamond	X, x
Hy	Chequerboard	X	Annulus, Bar, Diamond	Y, y
Yw	Chequerboard	Y	Annulus, Bar, Diamond	X, x
Bu	Chequerboard	Y	Annulus, Bar, Diamond	Y, y
Fl	Diamond	X	Annulus, Bar, Chequerboard	X, x
Cr	Diamond	X	Annulus, Bar, Chequerboard	Y, y
Dd	Diamond	Y	Annulus, Bar, Chequerboard	X, x
Jk	Diamond	Y	Annulus, Bar, Chequerboard	Y, y

896

897

898 Table 3. Example showing the Probe stimuli used in Experiment 2. In this example,
 899 A was the Reliable dimension, with its X value as positive, and B, C and D were the
 900 Unreliable dimensions, with their X and x values as positive. The dimensions used as
 901 reliable and unreliable, and the values used as positive and negative, varied between
 902 birds in a balanced fashion.

Stimulus dimension				
Annulus (Reliable)	Bar	Chequerboard (Unreliable)	Diamond	Number of times used
X	Y	Y	Y	6
Y	X	X	X	6
X	y	y	y	6
Y	x	x	x	6
X	X	Y	Y	2
Y	Y	X	X	2
X	Y	X	Y	2
Y	X	Y	X	2
X	Y	Y	X	2
Y	X	X	Y	2

903

904

905 Table 4. Experiment 2: Proportions of training trials within test sessions in which the
 906 response was correctly predicted by each dimension of the stimulus

		Proportions of trials correctly predicted by value of				
Reliable dimension	Pigeon	Positive stimulus (Reliable dimension and majority of unreliable dimensions)	Individual unreliable dimensions			
			Annulus	Bar	Checks	Diamond
Annulus	Mo	0.64		0.63	0.58	0.58
Annulus	Ct	0.87		0.76	0.67	0.70
Annulus	Ch	0.98		0.77	0.77	0.78
Annulus	Kc	0.98		0.72	0.79	0.79
Bar	Bn	0.70	0.85		0.55	0.50
Bar	Mr	0.64	0.53		0.60	0.58
Bar	Sa	0.66	0.69		0.58	0.53
Bar	Cu	0.95	0.80		0.68	0.73
Checks	Rg	0.88	0.77	0.61		0.73
Checks	Hy	0.98	0.79	0.73		0.77
Checks	Yw	0.89	0.80	0.70		0.66
Checks	Bu	1.00	0.81	0.76		0.77
Diamond	Fl	0.86	0.72	0.68	0.68	
Diamond	Cr	0.46	0.41	0.61	0.38	
Diamond	Dd	0.86	0.74	0.70	0.68	
Diamond	Jk	0.71	0.77	0.64	0.51	

907

908 **Figure Legends**

909

910 Fig. 1. Experiment 1: Examples of stimuli showing each of the two picture types
911 (house and car) associated with each of the two artificially introduced textures
912 (leopard-skin and blue wave). Each stimulus measured 160 x 120 pixels. (From
913 original photographs by Casper Addyman, used with permission)

914

915 Fig. 2. Diagram of touch screen display, showing size and position of array and two
916 side keys. Cells were numbered 1-8, clockwise from top left. Cells 3-6 were
917 reinforced on the right of the screen: cells 7, 8, 1 and 2 reinforced on the left.

918

919 Fig. 3. Proportions of probe stimuli responded to in accordance with the texture cue.
920 Data are shown separately for each pigeon, separated by the stimulus type that was
921 positive in training (*=Difference from 0.5 significant at 0.05 level).

922

923 Fig. 4. Examples of the stimulus sets used in Experiment 2. The upper two panels
924 show the A and a versions of each dimension, the lower two panels the b and B
925 versions.

926

927 Fig. 5. Experiment 2: Proportions of probe trials in which the stimulus was responded
928 to in accordance with the Reliable stimulus dimension. Data are shown separately for
929 each pigeon, grouped by the dimension that was designated as Reliable (*=Difference
930 from 0.5 significant at 0.05 level).

931

932 Fig. 6. Experiment 2: Proportions of probe stimulus trials on which each of the
933 Unreliable dimensions correctly predicted response. Data are shown separately for
934 each pigeon, grouped by the dimension that was designated as Reliable. Gaps within
935 the cluster of bars for each pigeon correspond to the Reliable dimension
936

Figure 1.

Fig. 1 Experiment 1: Examples of stimuli showing each of the two picture types (house and car) associated with each of the two artificially introduced textures (leopard-skin and blue wave). Each stimulus measured 160 x 120 pixels. (From original photographs by Casper Addyman, used with permission)

a



b



Figure 2

Fig. 2 Diagram of touch screen display, showing size and position of array and two side keys. Cells were numbered 1-8, clockwise from top left. Cells 3-6 were reinforced on the right of the screen: cells 7, 8, 1 and 2 reinforced on the left.

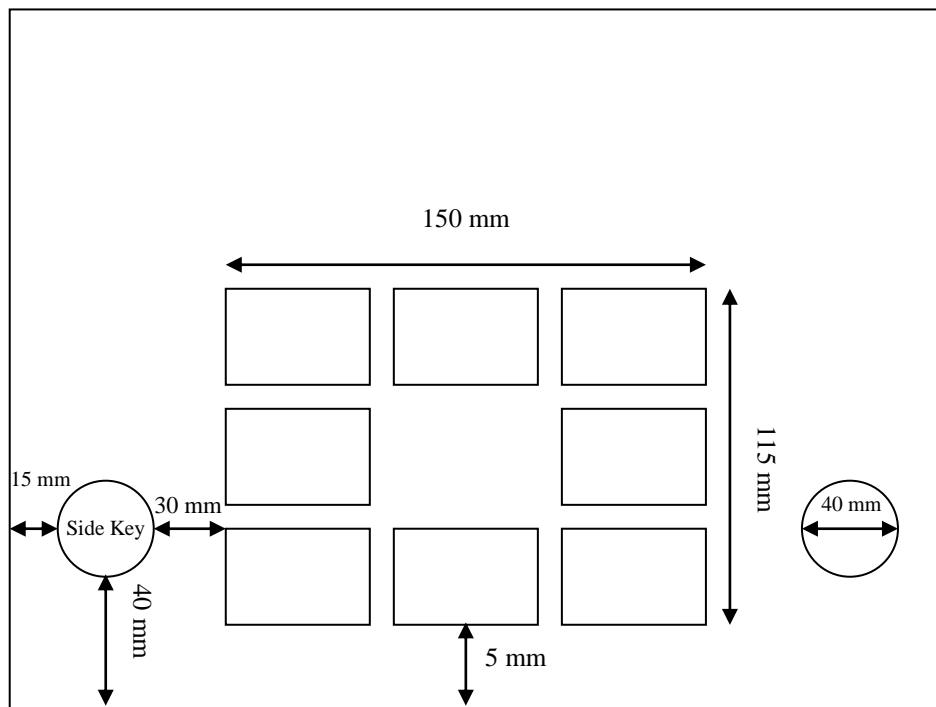


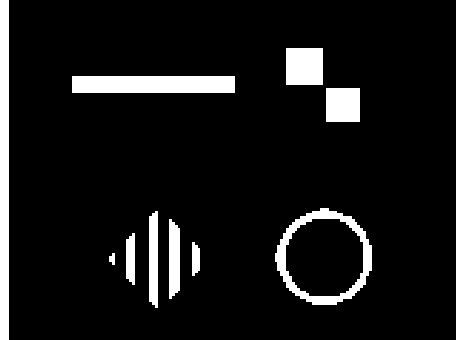
Figure 3

Fig. 3 Experiment 1: Proportions of probe stimuli responded to in accordance with the texture cue by each pigeon, separated by the stimulus type that was positive in training (*=Difference from 0.5 significant at 0.05 level).

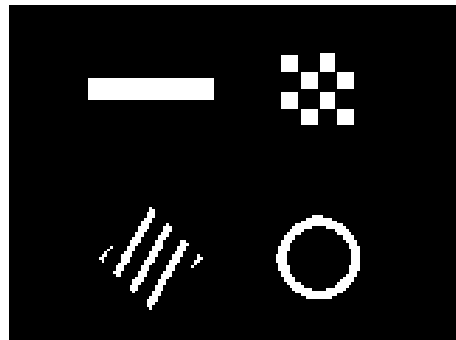
[see separate file]

Fig. 4. Examples of the stimulus sets used in Experiment 2. The upper two panels show the X and x versions of each dimension, the lower two panels the y and Y versions.

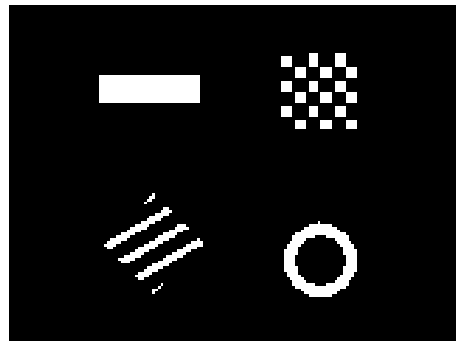
X form of all dimensions



x form of all dimensions



y form of all dimensions



Y form of all dimensions

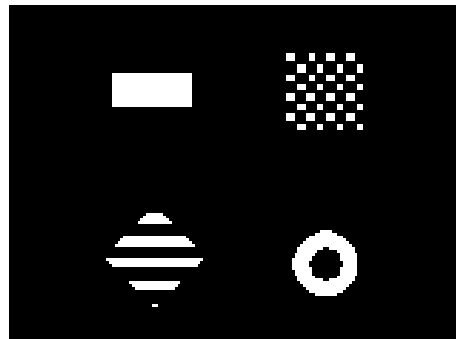


Fig. 5 Experiment 2: Proportions of probe trials in which the stimulus was responded to in accordance with the Reliable stimulus dimension. Data are shown separately for each pigeon, grouped by the dimension that was designated as Reliable (*=Difference from 0.5 significant at 0.05 level). Gaps within the cluster of bars for each pigeon correspond to the Reliable dimension

[see separate file]

Fig. 6. Experiment 2: Proportions of probe stimulus trials on which each of the Unreliable dimensions correctly predicted response. Data are shown separately for each pigeon, grouped by the dimension that was designated as Reliable.

