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2 Contingency in Learning

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6 Synonyms

7 Assignment of credit; Association; Contingency; Correla-
8 tion; Dependence; Prediction; Retrodiction

9 Definitions

10 *Contingency*: the extent to which knowledge of one event
11 reduces uncertainty about another. *Prediction*: the extent
12 to which knowledge of one event's occurrence enables
13 one to anticipate whether and/or when another event
14 will occur. *Assignment of credit*: determining to which
15 past event an outcome event should be attributed (retro-
16 diction). *Association*: perceived contingency. *Instrumental*
17 *conditioning*: a learning protocol in which a desired or
18 undesired is contingent on an action of the subject
19 (or agent). *Pavlovian conditioning*: a learning protocol in
20 which the contingency between two events is varied.
21 *Entropy*: the measure of amount of uncertainty, aka the
22 amount of information available in a probability distribu-
23 tion. *Mutual information*: the sum of the entropies of the
24 marginal distributions minus the entropy of their joint
25 distribution. *Uncertainty coefficient*: the percent reduction
26 in uncertainty about whether and/or when a predicted
27 event will occur that is produced by the occurrence
28 of a predictor event. **[Aut]** A broadly useable measure of
29 contingency or association.

30 Theoretical Background

31 The concept of contingency plays a central role in the
32 analysis of commonly studied learning paradigms and
33 also in research on human judgments of dependence,
34 contingency, and causality. Despite its conceptual impor-
35 tance, there is surprisingly small psychological literature
36 focusing on the following question: What is the proper
37 definition or measure of contingency?

In Instrumental/Operant Conditioning 38

39 The concept of contingency is important in the study of
40 instrumental conditioning, because the reinforcing
41 event only reinforces the instrumental response if it is
42 contingent on that response. In the operant conditioning
43 literature, the concept has often been treated as
44 unproblematic, perhaps because the experimenter
45 specified the contingencies that were taken to be of
46 interest. However, implicit in many treatments of rein-
47 forcement – and explicit in discussions of the role of delay
48 of reinforcement – is the assumption that what really
49 matters is not contingency per se but rather the close
50 *temporal pairing* of response and reinforcement.
51 This makes the question of the role of contingency in
52 instrumental conditioning the same as the question of its
53 role in Pavlovian conditioning. The challenge in both cases
54 is to specify what constitutes “close.”

55 As the study of reinforcement learning from
56 a computational perspective has become a significant
57 focus of research in computer science and cognitive neu-
58 roscience, there has been a greater realization that it was
59 not obvious which aspects of a sequence of actions should
60 be regarded as the aspect on which the feedback-providing
61 outcome was contingent. How to determine this is
62 the assignment of credit problem. It is the contingency
63 problem seen from the other end. It can be reformulated
64 as: What aspect or aspects of an action sequence is
65 an outcome contingent on? One wants a measure of
66 contingency or dependency that is mathematically well
67 grounded and lends itself to the apportionment of
68 contingency or dependency among possible predictors.

In Pavlovian/Classical Conditioning 69

70 The concept of contingency became important in the
71 study of Pavlovian conditioning in the late 1960s when
72 a series of experiments from different laboratories called
73 into question the assumption that the temporal pairing
74 was what drove the formation of an association between
75 two stimuli or events (hereafter called the CS and US, with
76 the CS being the predictor and the US the predicted event
77 or stimulus). Rescorla (1968) posed the question whether
78 it was the temporal pairing of CS and US or the CS–US
79 contingency that led to the emergence of a conditioned

80 response (a response to the CS that anticipates the US). He
 81 fixed the number of co-occurrences (temporal pairings) of
 82 the CS and US and varied the contingency by varying the
 83 frequency of US during intervals when the CS was absent.
 84 When there were no US in the absence of the CS, a strong
 85 conditioned response was seen on the post-conditioning
 86 test trials, even when $p(US|CS)$, the probability of the US
 87 given the CS was as low as 0.1. Regardless of the value
 88 of $p(US|CS)$, as the frequency of the US in the absence
 89 of the CS increased, the strength of the conditioned
 90 response on test trials diminished (see Fig. 1). When
 91 $p(US|\sim CS) = p(US|CS)$, that is, when the contingency
 92 was eliminated, there was no conditioned response.
 93 Thus, it is predictive (and retrodictive) power or contin-
 94 gency rather than temporal pairing that drives condition-
 95 ing. That is also the implication of the phenomena of
 96 *blocking*, *overshadowing*, and *relative validity*, which were
 97 discovered at about the same time. All of these phenomena
 98 imply that the critical aspect of a conditioning protocol
 99 is the predictive power of CS (or of the response in instru-
 100 mental conditioning), the extent to which it improves the
 101 subject's ability to anticipate when the US will occur.

102 Measures of Contingency

103 Most measures of contingency in the psychological litera-
 104 ture derive from the numbers in a 2×2 contingency table
 105 (Table 1). Several have been used, but only two have
 106 suitable mathematical properties, such as ranging from
 107 0 to 1 and not depending on N . Both of these are
 108 properties of the correlation coefficient, but that measure
 109 cannot be computed for dichotomous variables. For
 110 dichotomous variables in psychological experiments,
 111 Pearson's mean square coefficient of contingency

$$\phi = \sqrt{\chi^2/N} = \sqrt{\frac{(ad - bc)^2}{(a + b)(c + d)(a + c)(b + d)}}$$

112 is recommended by Gibbon et al. (1974), while the
 113 difference in the conditional probabilities of the US,

$$\Delta P = p(US|CS) - p(US|\sim CS) = \frac{a}{a + b} - \frac{c}{c + d}$$

114 has been used extensively in studies of human contingency
 115 and causality judgment (see, e.g., Allan et al. 2008).

116 Table-based measures are, however, problematic when
 117 applied to instrumental and Pavlovian conditioning
 118 experiments, which do not reliably have a definable trial
 119 structure (Gallistel and Gibbon 2000). This is apparent
 120 when one considers how to construct the contingency
 121 table for Rescorla's experiment. In that experiment, the
 122 CS always lasted 2 min. The interval between CS varied

around an average of 10 min. There is no doubt about how
 many CS and US there were, so the first cell (a in Table 1) is
 readily determined. All the other cells are problematic,
 because there is no objectively justifiable answer to the
 question: How many not-US and how many not-CS
 were there? The values of contingency underlying Fig. 1
 were obtained by following the common practice of
 assuming that the intervals between CS presentations are
 composed of "trials" of 2-min durations each, during
 which a US either occurs or does not. The number of
 $\sim CS$ is taken to be the number of such arbitrary
 subdivisions. The number of $\sim US$ is the total number of
 2-min intervals, including those when the CS was present,
 minus the number in which a US occurred. However, the
 2-min "trials" during the intervals between CS are
 a fiction, as is the number of not-US. Absent objectively
 defined trials, not-US and not-CS have no objectively
 definable relative frequency, so one cannot construct
 a contingency table. This problem is acute in the instru-
 mental conditioning case, because there are no trials in
 those protocols.

A second problem with measures based on
 a contingency table, and with the correlation coefficient
 as well, is that they take no account of time. The contin-
 gencies of ordinary experience are defined over time,
 and the temporal intervals between the events are
 centrally relevant to the psychological perception of
 contingency and causality. The importance of "close"
 temporal pairing – of response and reinforcer, or of CS
 and US – has always been stressed in the conditioning
 literature. However, attempts to specify what constitutes
 "close" have never succeeded. Clearly, a psychologically
 useful measure of contingency must take time into
 account.

A measure that does this is the uncertainty coefficient,
 also known as the entropy coefficient. It is the percent
 reduction in uncertainty about when (or whether)
 a predicted event (US) will occur gained from knowledge
 of the times at which (or trials on which) the predictor
 event (CS) occurred:

$$UC = I(CS; US)/H(US). \quad (1)$$

$I(CS; US)$ is the mutual information between CS and US.
 $H(US)$ is the US entropy of the US distribution. It is also
 called the amount of "available" or "source" information.
 It is the information-theoretic measure of the uncertainty
 regarding when and/or whether a US will occur. In the case
 of atemporal dichotomous variables, where there are
 objectively definable trials, hence objective probabilities
 for the failure of a US to occur,

$$H(US) = \sum p_i \log_2(1/p_i) = p(US) \ln(1/p(US)) + p(\sim US) \ln(1/p(\sim US)). \quad (2)$$

171 The $\ln(1/p_i)$ is the amount of information provided
172 by the occurrence of the i th event in the set of possible
173 events over which a probability distribution is defined
174 (e.g., the US and \sim US events). It is also called the surprisal.
175 Intuitively, the less probable the event, the more
176 unexpected or surprising it is, the more we are informed
177 by its occurrence – but, by the same token, the less often
178 we are so informed. As may be seen from Eq. 2, the
179 entropy of a distribution, ~~H , of a distribution~~ is simply
180 the average surprisal, that is, the amount of information
181 provided by each of the possible events weighted by its
182 relative frequency.

183 Entropy is the technical term for the amount of uncertainty
184 in a probability distribution, which is the same as
185 the amount of information available from that distribution,
186 because information reduces uncertainty. The
187 mutual information between two events with observed
188 or experimenter-defined probability distributions is:

$$I(CS; US) = H(CS) + H(US) - H(CS, US),$$

189 where $H(US, CS)$ is the entropy of the joint CS–US distribution.
190 In the case where a contingency table can be
191 constructed, the US distribution is given by the normalized
192 column totals, that is, the column totals in Table 1
193 divided by N ; the CS distribution is given by the normalized
194 row totals; and the joint distribution is given by the
195 normalized cell values (a/N , b/N , c/N , d/N). For each
196 distribution, the entropy is: $H = \sum p_i \ln(1/p_i)$.

197 The UC measure applies to temporal uncertainty as
198 well (Balsam and Gallistel 2009). If US_i (or reinforcers)
199 occur at random times, then the uncertainty regarding
200 when the next US will occur is the entropy of an exponential
201 distribution, which depends only on the average
202 US–US interval (the reciprocal of the base rate). This
203 entropy is the basal uncertainty about when the next US
204 will occur. It is the amount of available information. If
205 a CS always precedes a US and always tells us exactly when
206 to expect the US, then there is no residual *objective* uncertainty
207 about when the next US will occur once the CS has
208 occurred. In that case, the UC is 1, that is, the CS reduces
209 the uncertainty about when the next US will occur by
210 100%. However, humans and other common laboratory
211 animals can only estimate the duration of an elapsing
212 interval with about $\pm 15\%$ accuracy. To be useful, the
213 CS must precede the US by some interval. Our residual
214 uncertainty about when exactly to expect the US is then
215 determined by our imprecision in estimating when the

remembered CS–US interval has elapsed. Thus, the effective
216 percent reduction in our uncertainty depends on the
217 ratio between the basal interevent interval (the average
218 US–US interval) and the CS–US interval (the delay of
219 reinforcement). The greater this ratio is, the greater the
220 percent reduction in our uncertainty. Thus, this way of
221 measuring contingency explains why “close” temporal
222 pairing is important. However, “close” is relative (to the
223 basal interevent interval), not absolute; there is no critical
224 interval that defines whether two events are or are not
225 temporally paired. 226

227 Important Scientific Research and Open Questions 228

229 The UC measure of contingency provides a rationale for
230 the two ideas in the famous Rescorla–Wagner model of
231 association formation: $\Delta V = \alpha(\lambda - \sum V)$, where V is
232 associative strength. This formula rests on two assumptions:
233 (1) The sum across all the associations from different
234 CS to one US cannot exceed some limit, which is
235 represented by the asymptote parameter, λ . (2) Associative
236 strengths are additive; their sum is subtracted from λ in
237 determining the amount by which any associative strength
238 is to be incremented, ΔV . The entropy of the US distribution,
239 which determines the amount of available information,
240 puts an upper limit on the amount of information that all
241 predictors combined can provide. Moreover, the entropies of
242 independent events (and independent conditional entropies)
243 are additive. An open question is how far this can take us
244 in understanding the objective basis for the phenomena of
245 cue competition (blocking, overshadowing, relative validity) –
246 see Balsam and Gallistel (2009). 247

248 Another open question is whether and how the brain
249 can compute the uncertainties on which the UC measure
250 of contingency depends. 250

251 Cross-References 251

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262 **References**

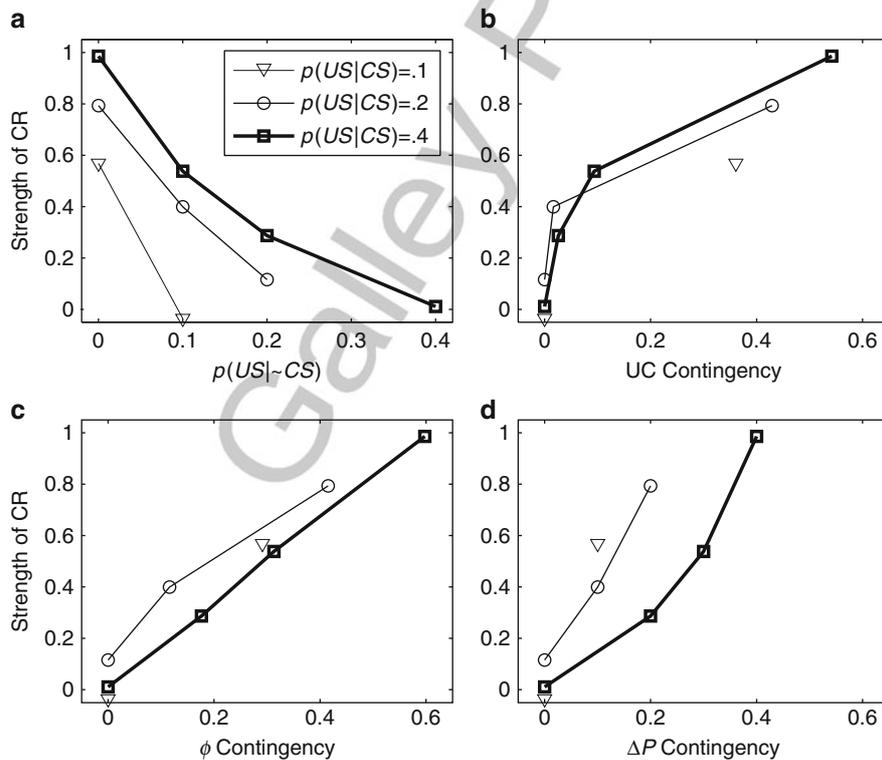
263 Allan, L. G., Hannah, S. D., Crump, M. J., & Siegel, S. (2008).
264 The psychophysics of contingency assessment. *Journal of Experimen-*
265 *tal Psychology: General*, 137(2), 226–243.
266 Balsam, P., & Gallistel, C. R. (2009). Temporal maps and informativeness
267 in associative learning. *Trends in Neurosciences*, 32(2), 73–78.
268 Gallistel, C. R., & Gibbon, J. (2000). Time, rate, and conditioning.
269 *Psychological Review*, 107(2), 289–344.

Gibbon, J., Berryman, R., & Thompson, R. L. (1974). Contingency 270
spaces and measures in classical and instrumental conditioning. 271
Journal of the Experimental Analysis of Behavior, 21(3), 585–605. 272
Rescorla, R. A. (1968). Probability of shock in the presence and absence of 273
CS in fear conditioning. *Journal of Comparative and Physiological* 274
Psychology, 66(1), 1–5. 275

Galley Proof

t1.1 Contingency in Learning. Table 1 2x2 contingency table

	#US	#~US	Row totals
t1.2 #CS:	a	b	$a+b$
t1.3 #~CS:	c	d	$c+d$
t1.4 Col totals:	$a+c$	$b+d$	



Contingency in Learning. Fig. 1 (a). The strength of the CR on first test trial as a function of $p(US|CS)$ and $p(US|\sim CS)$ in Rescorla's (1968) experiment on the role of CS-US contingency as against temporal pairing. Although in each of the three conditions, the temporal pairing of US and CS [hence $p(US|CS)$] was held constant, the strength of the CR declined to zero as the contingency was degraded by increasing $p(US|\sim CS)$. **(b)** Performance data in **a** plotted against the uncertainty coefficient (UC) measure of contingency. **(c)** Performance data in **a** plotted against the ϕ measure of contingency. **(d)** Performance data in **a** plotted against the ΔP measure of contingency

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AU1	Please check sentence starting "A broadly useable measure..." for completeness.	it was incomplete because improperly punctuated; I have corrected it