

1 **Music predictability and liking enhance pupil dilation and promote motor learning in non-musicians**

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11 **Abstract**

12 Humans can anticipate music and derive pleasure from it. Expectations facilitate movements associated
13 with anticipated events, and they are linked with reward, which may also facilitate learning of the
14 anticipated rewarding events. The present study investigates the synergistic effects of predictability and
15 hedonic responses to music on arousal and motor-learning in a naïve population. Novel melodies were
16 manipulated in their overall predictability (predictable/unpredictable) as objectively defined by a model
17 of music expectation, and ranked as high/medium/low liked based on participants' self-reports collected
18 during an initial listening session. During this session, we also recorded ocular pupil size as an implicit
19 measure of listeners' arousal. During the following motor task, participants learned to play target notes
20 of the melodies on a keyboard (notes were of similar motor and musical complexity across melodies).
21 Pupil dilation was greater for liked melodies, particularly when predictable. Motor performance was
22 facilitated in predictable more than unpredictable melodies, but liked melodies were learned even in the
23 unpredictable condition. Low-liked melodies also showed learning but mostly in participants with higher
24 scores of task perceived competence. Taken together, these results suggest that effects of predictability
25 on learning can be overshadowed by effects of stimulus liking or task-related intrinsic motivation.

26 **1 Introduction**

27 Through passive exposure to music, we implicitly develop models about its structure ^{1,2}. These models
28 allow both listeners to generate expectations about upcoming musical events ^{3–5}, and trained musicians
29 to better plan and learn musical actions ^{6–9}. Expectations are also linked to the experience of musical
30 pleasure ^{10–12}, as neuroimaging evidence shows that musically evoked pleasure relies on the cross-talk
31 between neural systems responsible for prediction with those responsible for reward ¹³. Importantly,
32 rewarding stimuli increase arousal ¹⁴, and are also better learned ¹⁵. Given the inherent link between
33 predictability and pleasure in music, here we aim to assess their contributions to learning with the
34 hypothesis that motor learning in naïve population may benefit from the implicit musical expectations
35 and hedonic responses derived from music.

36 A large body of theoretical and experimental work suggests that agents use internal psychological
37 models to make sense of perceptual inputs and respond to them. Through implicit statistical learning,
38 the brain continuously scans the environment for regularities and acquires probabilistic models of the
39 world without deliberate effort or awareness ^{16–18}. For example, based on these internal models, the
40 brain can compute the statistical distribution of sequential phenomena ^{19,20} enabling it to predict
41 unfolding sensory events, thereby reducing its uncertainty ²¹. In particular, according to predictive coding
42 theory ²², by comparing top-down predictions with the actual sensory input, internal predictive models
43 contribute to the selection of relevant bottom-up inputs ²³, and error signals are used to update
44 predictions, construct new models, and guide subsequent actions. Internal models can aid motor
45 planning ^{24,25} and it has even been shown that visual statistical learning can inform the motor system to
46 better predict upcoming actions ^{26,27}.

47 Similar cognitive mechanisms are likely to play a role in musical perception and performance. When
48 listening to music, humans entertain a number of predictions, or hypotheses, about future musical
49 events ^{1,28}, and these predictions are subject to refinement and learning on different time scales and at
50 different levels of sophistication. For example, musical predictions vary as a function of the musical
51 culture one is exposed to ^{29–31} and can adapt to novel musical styles ³². Furthermore, musical predictions
52 are optimized by expertise ³³, and can even vary across the lifespan ³⁴. With regard to performance, it
53 has been shown that musical regularities facilitate movement selection ^{35,36}. This facilitation is supported
54 by associations between movement and ensuing effects formed through coupling of motor and sensory
55 cortices ^{36–39}. Moreover, studies in trained musicians show that internal predictive models allow long-
56 range motor planning of entire musical sequences ^{6,7,40–43}. Thus, motor performance in experts appears
57 to be guided by predictions based on learned internal models of music, suggesting that also initial stages
58 of learning may benefit from them.

59 Expectations are also linked to the experience of musical pleasure ^{10–12,44}. Pleasurable music strikes a
60 balance between predictable events, which allow listeners to form expectations, and moderately

61 unpredictable events that produce surprise. For example, a single repeating note is very predictable, but
62 may not be very enjoyable, and similarly, transitions to unrelated notes may be perceived as unpleasant
63 ^{45,46}. Indeed, musical pleasure seems to vary with stimulus complexity – e.g., harmonic or rhythmical
64 predictability – as an inverted-U function, with maximum liking occurring at intermediate levels of
65 complexity ^{47–50}. Neural evidence suggests that musical surprises induces liking by engaging the reward
66 system ⁵¹, via distinct phases of dopamine transmission during the anticipation and enjoyment of
67 listeners' favorite musical moments ⁵². Connectivity analyses suggest that it does so in cooperation with
68 fronto-auditory systems responsible for predictions during listening ¹³. The role of dopamine, which has
69 been widely linked to the 'wanting' and 'learning' components of reward ⁵³, has been also directly linked
70 with abstract hedonic responses to music (i.e., liking) ⁵⁴. Furthermore, a recent study showed that
71 pleasant musical moments activate the noradrenergic arousal system, as revealed by increased pupil
72 dilation during passive listening ¹⁴. A concomitant increase of arousal and reward systems during
73 pleasant music seems plausible given the known anatomical and functional link between dopaminergic
74 and noradrenergic subcortical nuclei ^{55,56}, and may explain findings of better memory for rewarding than
75 neutral musical excerpts ⁵⁷. Although rarely used to measure response to long stimuli – such as a melody
76 ^{58,59}, pupil dilation may thus be powerful to continuously track hedonic responses to unfolding music ⁶⁰.

77 These evidence that musical expectations contribute to increase of arousal and pleasure suggest a
78 possible link with learning based on the relevance of rewarding stimuli and reward-related dopamine
79 circuits ¹⁵. For example, animal studies show that brain plasticity associated with auditory learning is
80 greater when the information to be learned is rewarded ⁶¹. Further, pairing a tone with stimulation of
81 dopamine circuits in the brainstem increased the selectivity of responding in auditory neurons tuned to
82 the same tone ⁶². Importantly, dopamine has also been shown to modulate motor learning in humans
83 and animals both directly ^{63,64}, and indirectly through monetary reward ^{65,66}. Based on this body of work,
84 we test the idea that, by carrying abstract reward, music that is better liked could be associated with
85 greater arousal and learning.

86 As the motivation to learn a musical excerpt can derive from perceived pleasure, learning can also be
87 motivated by its inherent challenge. Psychological and computational accounts of motivation distinguish
88 extrinsic from intrinsic motivation: whilst the first is based on external reward or pressures outside the
89 individual, the latter is defined as doing an activity for its inherent satisfactions, that have the appeal of
90 aesthetic value or challenge for the individual ^{67,68}. It is possible that learning becomes pleasant for
91 actions that lead to decrease of uncertainty and improvement of internal predictive models ^{69–71}. Beside
92 this, motivation to learn may be also driven by individual's feeling of competence in achieving a self-
93 determined goal just for the challenge entailed by the task ⁶⁸. Therefore, individual differences in intrinsic
94 motivation should also be assessed to predict learning.

95 We tested participants with no-to-little musical training with pupillometry in a listening task, followed
96 by a melody learning task (Figure 1). We composed novel melodies which varied in their structural

97 predictability (from overall predictable to unpredictable melodies). We could formally determine the
98 complexity of the stimuli by means of a variable-order Markov model of melodic expectation (IDyOM)¹:
99 this model acquires knowledge of musical structure through unsupervised statistical learning and uses
100 this knowledge to estimate the probability of musical notes in a given melody. The expectedness of each
101 note in the melody is expressed in units of information content (IC), where high and low IC values
102 correspond respectively to less and more predictable notes. During the listening task, we recorded the
103 ocular pupil dilation response, and participants were asked to rate how much they liked each melody on
104 a seven-point scale. In the melody learning task, participants learned to play the last four notes of the
105 melodies on a piano-type keyboard (learning phase), and they were tested thereafter (test phase).
106 Importantly, the last four notes did not differ in predictability or motor requirements (See Figure 1), but
107 only in the predictability of the preceding musical context.

108 First, we hypothesized that moderately predictable melodies would be better liked, consistent with the
109 inverted-U hypothesis. Further, we predicted that sustained pupil dilation would be greater for melodies
110 that were better liked. For the melody learning task, we expected that predictable musical contexts
111 would result in better motor implementation of target notes, and that better-liked melodies would
112 potentially be better learned even in the unpredictable condition. Because there are large inter-
113 individual differences in music reward sensitivity and intrinsic motivation to perform a task, these
114 characteristics were assessed via questionnaires: Barcelona Music Reward Questionnaire;⁷² and the
115 Intrinsic Motivation Inventory (with focus on interest/ enjoyment, and perceived competence
116 subscale)⁷³.

117 **Figure 1**

118 **2 Results**

119 **2.1 Liking ratings of the melodies as function of Predictability**

120 Based on theoretical and empirical work⁴⁷⁻⁴⁹, we expected an inverted U-shape relationship between
121 subjective liking and stimulus complexity. Therefore, a parabola was fitted in a model describing
122 participant's ratings (scaled by subject) as a function of IC of melodies (averaged across notes). Figure 2
123 shows the quadratic relationship between melody IC and participants' liking ratings ($\chi^2(1) = 4.513$, $p =$
124 $.033$), indicating that liking was higher for moderately predictable melodies, but it decreases when
125 melodies become more complex (high IC). Possibly because of a relative narrow range of IC across
126 melodies, a linear model could also fit the relationship between liking and IC ($\chi^2(1) = 4.174$, $p = .041$),
127 which yielded similar model fit to the quadratic term ($\Delta AIC = .6$).

128 **Figure 2**

129 **2.2 Pupil diameter during listening**

130 During listening, participants provided liking ratings to each of the melodies whilst pupil dilation was
131 measured. We analyzed the effects of predictability (Predictable/Unpredictable) and liking
132 (High/Medium/Low) and their interaction on participants' pupil size change over 13 time-bins,
133 corresponding to the 13 notes in the melodies (Figure 3). We found an effect of predictability by time
134 bin on the sustained pupil response ($\chi^2(1) = 5.291, p = .021$). Although the differential increase over time
135 did not reach statistical significance after multiple comparisons, post hoc paired t-test showed overall
136 greater pupil dilation for predictable compared with unpredictable melodies [P-U: $t(1,22) = 3.137, p =$
137 $.005$]. Moreover, we found an interaction of liking and time bin ($\chi^2(2) = 29.257, p < .001$): this indicated
138 that high-liked melodies induced greater dilation than medium-liked (high-medium: $b = 5.942, SE =$
139 $1.458, p < .001$), or than low-liked melodies (high - low: $b = 3.928, SE = 1.469, p = .021$). A three-way
140 interaction of predictability, liking and time bin ($\chi^2(2) = 6.152, p = .046$) showed that pupil dilation for
141 the high-liked condition increased more for predictable than unpredictable melodies (P - U: $b = 4.846,$
142 $SE = 2.107, p = .021$). No predictability effect was indeed found in the low-liked (P - U: $b = -1.113, SE =$
143 $2.05, p = .587$), or medium-liked condition (P - U: $b = -1.783, SE = 2.19, p = .377$). These results suggest
144 that the increase of pupil dilation as a function of liking was greater when the music was predictable.

145 Finally, individual scores of music reward sensitivity positively predicted pupil size change across all trials
146 regardless of condition ($\rho = .320, p = .005$) (Figure 3; right panel), showing that individuals who are more
147 sensitive to musical reward have a greater physiological response.

148

Figure 3

149 **2.3 Motor performance**

150 In the motor task participants listened to the first nine *context* notes of each melody through
151 headphones and then played the last four *target* notes on a piano-type keyboard. Accuracy and
152 asynchrony were entered in separate mixed effects regressions for the training and the following test
153 phase.

154 In the training phase, we found increase of *accuracy* across repetition trials ($\chi^2(3) = 22.498, p < .001$),
155 and no other effect involving Predictability or Liking ($p > .111$). For *asynchrony* (Figure 4), there was a
156 main effect of Predictability ($\chi^2(1) = 11.677, p < .001$), and an interaction with repetition trial ($\chi^2(1) =$
157 $5.758, p = .016$), such that predictable trials were better executed, and that this advantage were greater
158 in the early trials (P – U across repetition trials: $b = 5.883, SE = 2.753, p = .033$). An interaction between
159 Liking and Predictability ($\chi^2(2) = 9.092, p = .01$) indicated that these effect of predictability mainly
160 regarded the medium-liked melodies (P - U: $b = -27.872, SE = 11.454, p = .015$), but not the high-liked (P
161 - U: $b = -5.675, SE = 11.659, p = .131$), and low-liked ones (P - U: $b = -17.484, SE = 11.574, p = .131$). The
162 poorer performance for medium-liked melodies in the unpredictable condition suggests that the
163 unpredictability of the stimulus is compensated when strong (positive or negative) rather than mild
164 (medium) emotional responses are at play. We indeed found a significant main effect of Liking ($\chi^2(2) =$

165 16.755, $p < .001$), and an interaction with repetition trial ($\chi^2(2) = 21.263, p < .001$) such that learning was
166 better for the high-liked and the low-liked melodies when compared with the medium-like ones (across
167 repetition trials high-medium: $b = -12.718, SE = 3.401, p < .001$; low-medium: $b = -7.998, SE = 3.313, p =$
168 .016; high-low: $b = -4.720, SE = 3.398, p = .165$).

169 Taken together, these findings indicate that learning of the target-notes was greater after predictable
170 than unpredictable contexts in presence of mild emotional responses. However, the disadvantage due
171 to the unpredictable contexts was minimized for high- and low-liked melodies, which showed similar
172 learning regardless of predictability.

173 **Figure 4**

174 We further investigated the role of other factors than stimulus-induced affective responses in learning,
175 as inter-individual differences associated with intrinsic motivation. Specifically, we tested the
176 relationship of the general enjoyment and task-perceived competence IMI scales, with the learning
177 slopes – measured as reduction of asynchrony across repetition trials – in the high-like and low-like
178 conditions. To estimate the individual performance improvement across repetition trials as a function of
179 liking, we ran mixed model testing for only the effect of liking (High/Medium/Low) across repetition
180 trials, and extracted residual individual slopes for the low-/high-liked condition, after adjustment for the
181 effects of the performance change in the medium like condition at the intercept. More negative slopes
182 (faster learning) associated with low-liked condition were predicted by higher scores of individuals'
183 perceived competence in succeeding the task (Fig. 4 right panel, $\rho = -.488, p = .015$), and there was a
184 trend with the general enjoyment scale ($\rho = .393, p = .085$). The learning slope for high-liked melodies
185 did not correlate with perceived-competence scores ($\rho = -.224, p = .292$), nor with general enjoyment
186 scale ($\rho = .147, p = .535$). Thus, the learning effect observed in low-liked melodies was particularly
187 driven by participants with high task perceived competence, whilst liked music was learned regardless.
188 This suggests that perceived competence helped the participants overcome low liking to learn the
189 melodies, but it didn't play a role in highly liked stimuli as participants were already motivated by the
190 pleasure carried by music.

191 Furthermore, we tested if learning highly liked stimuli was predicted by pupil response to those stimuli.
192 The correlation between the mean amplitude of pupil diameter in high-like trials and the learning slope
193 of high-like condition did not yield significant results ($\rho = .044, p = .84$). This may be due to the fact that
194 the range of liking values was relatively limited.

195 In the test phase, we didn't observe any significant effect of liking nor predictability on accuracy and
196 asynchrony (all $p > .389$). This non-finding in the test phase may be explained by the fact that only one
197 trial per melody was not sufficient for seeing an effect. Also, the increased number of errors committed
198 in the test phase compared with the last repetition trial of the training phase (pitch and timing errors
199 combined, mean averaged across participants $10.74 \pm 26.60\%$, in the last repetition trial; $35.51 \pm 24.04\%$,

200 in the test phase; $t (25) = -5.940, p < .001$, Cohen's $d = -2.376$), suggests that more repetitions may be
201 desirable for stabilization of learning.

202 **3 Discussion**

203 The present study investigated the contribution of predictability and liking on arousal and learning in
204 non-musicians. First, we found an inverted U-shaped relationship between music complexity and liking,
205 showing that moderately predictable melodies were more liked than highly predictable and
206 unpredictable melodies. Further, we showed a synergistic effect of predictability and hedonic response
207 to music on arousal, as reflected by sustained pupil dilation. We also observed that pupil dilation was
208 overall greater in individuals with higher sensitivity to musical reward, suggesting that it is a good marker
209 of responsiveness to music. With regard to melody learning, performance was facilitated for predictable
210 compared to unpredictable melodies in medium-liked music, indicating that musical expectations can
211 facilitate auditory-motor predictions and movement preparation, even in non-musicians. This effect of
212 predictability was overshadowed by musical reward as liked melodies were better learned, even when
213 unpredictable. Finally, we found that not-liked melodies were also learned and that this effect was
214 correlated with individuals' task perceived competence, suggesting that, beyond the musical reward,
215 other factors carrying reward — such as individual's task-related intrinsic motivation— contribute to
216 learning.

217 Our results link quantitative measures of stimulus complexity (as music predictability) with liking
218 response of listeners as an inverted-U-shaped function, whereby moderately predictable melodies were
219 more liked than highly predictable and unpredictable melodies. Importantly, listeners were unfamiliar
220 with the stimuli, and complexity was objectively characterized by the IDyOM model which has been
221 shown to optimally predict subjective perceptual expectations⁷⁴, and perceived complexity of musical
222 structure⁷⁵. The inverted-U model was first proposed by Berlyne (1971) to reflect a general relationship
223 between aesthetic appreciation and structural complexity in art. But, it has also been shown to be a
224 general property of complex stimuli including visual shapes⁷⁶, music and rhythm^{44,49}. In line with a
225 predictive account, liking may derive from an intrinsic reward which occurs whenever an internal
226 predictive model improves by decreasing uncertainty^{69,71}. Because the potential for decreasing
227 uncertainty is maximal when music is moderately complex, so it should be for the liking. The inverted-U
228 model has received empirical support in some music experiments^{47,48,50}, but not in others^{77,78}, possibly
229 because it is often difficult to generate ecologically valid stimuli that cover the full range of complexity,
230 or because other psychological mechanisms triggered by the stimulus such as familiarity, imagery,
231 memory or associations^{1,79} interact with expectation-based emotions.

232 The effect of liking on subject's arousal is in line with previous literature¹⁴, in that pupil response during
233 listening was greater for liked melodies. Importantly, we extend this finding by showing that pupil
234 dilation increased for high-liked melodies, but only when they were predictable. These results are novel

235 because they address for the first time both the effect of subjective pleasure and musical complexity on
236 sustained pupil response over relatively long stimuli. They are compatible with the interpretation that
237 sustained pupil dilation is modulated by both changes in attentional engagement due to stimulus
238 structure tracking ^{80,81}, and to subjective affective evaluation or reward ^{14,82}. In support of this
239 interpretation, electrophysiological evidence has established a link between pupillary response and
240 norepinephrine activity in the nucleus locus coeruleus ⁸³ that has synergistic connections with subcortical
241 dopaminergic nuclei involved in reward, and prefrontal areas involved in stimulus evaluation processes
242 ^{55,56}. Predictable melodies may thus result in greater attentional engagement – enhanced noradrenergic
243 activity – as they conform to listeners' prior expectations and allow them to form precise predictions
244 about the incoming stimulus ¹. Such effect of predictability may explain effects of greater sensitivity and
245 memory of music from one's own culture, or of simple more than complex excerpts ^{31,32}. Conversely,
246 unpredictable melodies may down-weigh predictions from a model that does not match the incoming
247 stimulus, resulting in attentional disengagement, and lower pupil response. The interaction effect of
248 liking and melody predictability on pupil dilation suggests that positive evaluative processes build on
249 successful tracking of the stimulus structure. One proposed mechanism is based on a hypothesized
250 feedforward loop between forebrain regions associated with reward evaluation and the concerted
251 action of the noradrenergic and dopaminergic systems ⁵⁵. As a positive subjective evaluation is formed
252 throughout the melody, succeeding valid predictions gain greater reward value through dopamine-
253 mediated response, which in turn boosts norepinephrine-mediated attention. Future investigations
254 combining pupillometry and brain imaging are necessary to identify this circuit, and the proposed
255 dynamic interaction during response to music.

256 Learning of the target notes was facilitated for predictable compared with unpredictable melodic
257 contexts in medium-liked music, demonstrating that musical structure promotes predictions and motor
258 encoding in naïve performers. Importantly, better learning cannot be explained by differences in
259 predictability or motor complexity intrinsic to the target notes, because these were similar across all
260 melodies. They only differed because they were embedded in contexts that allowed better or worse
261 prediction of the most likely continuation of the melody. Further, learning effects were observed for
262 temporal accuracy of the movements, not note accuracy – which was high in all conditions. This is
263 important because the stimuli varied in melodic expectations, but not in timing. Thus, better temporal
264 accuracy for more predictable melodic contexts indicates that musical structure promotes motor
265 prediction and planning by heightening the precisions of the movements. This result is in line with the
266 notion of 'active inference' ^{22,24}: by relying on models of the environment with a high level of precision
267 in predictable contexts, the brain can select a narrower set of information to predict the future
268 sensorimotor state and to reduce uncertainty ^{84,85}. Given that perception and action are intertwined,
269 perceptual and motor networks may also interact during the generation of predictions about the most
270 likely next state ²⁶. Accordingly, in music models of musical structure inform the sensory system to
271 anticipate the most predictable sound ⁸⁶, and they also drive the motor systems to facilitate the

272 movement required to produce it ^{37,39,42,87}. Moreover, there is recent evidence that non-musicians
273 rapidly form sensorimotor representations of anticipated events after short-term motor training ^{36,88}.
274 Our results suggest that even in naïve performers, predictions based on experience in the auditory
275 modality affect predictions in the motor domain, irrespective of previous training linking sounds to
276 actions. A possible underlying mechanism may be the rapid formation of sensorimotor associations at
277 the first attempts of execution, which result in facilitated performance in the following repetition trials
278 ⁸⁹. They may also be based on long-term priors –the so-called SMARC effect (Spatial Musical Association
279 of Response Codes) – which shows that even for individuals without training, higher pitches facilitate
280 upward or rightward responses, and low pitches facilitate downward or leftward responses ⁹⁰.
281 Alternatively, in line with the view that sensory and motor systems act as independent “emulators” of
282 upcoming events ⁹¹, predictive models in the motor domain may be independently generated based on
283 existing models of music built through auditory perception.

284 Liking a melody reduced the disadvantage in performance due to the unpredictable contexts, suggesting
285 that music-induced hedonic response promotes learning by overshadowing the effect of predictability.
286 A possible underlying mechanisms may be an interaction between dopamine-mediated reward induced
287 by music ⁷¹ and dopamine-mediated learning mechanisms ¹⁵. This is consistent with work reporting
288 enhanced motor learning and retention in presence of external incentives, such as monetary reward
289 ^{65,66}. In line with the idea the reward value of music may act as a reinforcement signal for learning ⁵¹, our
290 results foster the link between reward and motor learning in a more complex task and for an abstract
291 stimulus-related incentive.

292 The motor learning benefit associated with preferred music may be indirectly linked with general greater
293 attention, as reflected by increase of pupil in liked melodies. The well-known interaction between the
294 noradrenergic system – underlying pupil dilation – and the dopamine system – associated with reward
295 – ^{55,92}, suggest that the concerted action of these two systems may mediate the beneficial effect of music
296 reward on memory and motor learning. We did not find evidence to relate motor learning and pupil
297 response to liking ratings, probably because of the limited range of response elicited by the stimuli used
298 here. Previous studies using stimuli that induce musical chills have shown that they induce greater pupil
299 dilation ¹⁴, and are also better remembered ⁵⁷, consistent with the key role of the reward system in
300 stimulus encoding ⁹³. Future studies, likely in trained musicians, could use stimuli which induce more
301 intense pleasure to examine their effect on learning.

302 Non-liked melodies were also learned similarly to liked melodies, irrespective of music predictability. The
303 learning of non-liked melodies was driven by participants with higher task-achievement motivation
304 (perceived-competence scale), as opposed to a general learning effect of high-liked melodies. This is in
305 line with the definition of competence where the achieving process, rather than goal being achieved, is
306 central ⁶⁷. Thus, our results suggest that when the music is not rewarding per se, people with greater
307 general task-related motivation succeed better in learning it.

308 Predictability and liking are inherently linked in music. Their intertwined effect was evident in the
309 pupillary response, which was enhanced both by musical expectations and subjective music reward.
310 However, what are their contributions to learning when assessed separately? We observed that when
311 operationalized as information content, effects of predictability on learning can be over-shadowed by
312 effects of liking or of intrinsic motivation. One implication of this is that liking in music should neither be
313 reduced to “mere” liking – as it can drive learning, thus acting “as” a reinforcer, nor to mere predictability
314 – because unpredictable melodies were learned equally well when liked. This may also in part be due to
315 the fact that subjective liking plausibly involves many dimensions beyond predictability, such as
316 familiarity, imagery, memory or idiosyncratic associations ^{1,79}. These results reinforce the view that
317 emotional and motivational factors have powerful impact on learning not only for cognitive tasks, but
318 also for procedural and motor-skill learning ⁹⁴. In conclusion, this study provides an important first step
319 in understanding how motor learning benefits from the contributions of implicit musical expectations
320 and the derived emotional response. Future research in this direction may shed light on their additional
321 benefits on rehabilitation in clinical populations.

322 4 Material and methods

323 4.1 Participants

324 Twenty-seven individuals with no previous piano training took part in the study (18 female; Age: $M =$
325 24.74 ± 5.03). Participants had on average less than one year ($M = 0.8$ years ± 1.5) of formal music
326 training, which did not take place in the last 10 years (note that one participant with 16 years of training
327 in another instrument was excluded from the analysis). All participants were neurologically normal, were
328 not taking any medication that could affect motor performance, and had normal hearing, and normal or
329 corrected to normal visual acuity. All participants were naïve with regard to the purpose of the study
330 and provided written informed consent. The Concordia University Human Research Ethics committee
331 approved the study (30007730) and conducted adhering to the Canadian Tri-council Policy on ethical
332 conduct for research involving humans ⁹⁵.

333 4.2 Stimuli and Procedure

334 4.2.1 Stimuli

335 Sixteen different melodies of 13 notes were newly composed specifically for the experiment according
336 to the rules of classical Western tonal music. All of them began on the first beat, and were notated with
337 a time signature of 3/4 that was thought to be easier to count along. In order to focus specifically on
338 pitch expectations, each note had the same duration and equivalent inter-onset interval of 428 ms (140
339 bpm). Melodies were created with MuseScore program (version 2.0.2) and synthesized with a piano
340 sound (generated using Ableton Live 8) with the same loudness for all notes and melodies. Each melody
341 had a total duration of 6.4 s.

342 The predictability of each melody was objectively defined using the information dynamics of music
343 model, IDyOM ¹⁹ and based on the average information content values of each note. This model is
344 trained through a process of unsupervised learning on a large training set of 903 Western tonal melodies.
345 IDyOM first analyses the statistical structure of the training set, represented as sequences of pitch and
346 note's scale degree relative to the key of the melody. In a new sequence, it then estimates the probability
347 of each note, based on a combination of the training set's statistics and those of the sequence at hand,
348 which it learns dynamically. The output is a note-by-note measure of information content (IC, the
349 negative logarithm, to the base 2, of the probability of an event occurring), which IDyOM uses instead
350 of raw probability for greater numerical stability and a meaningful information-theoretic interpretation
351 in terms of redundancy and compression. We manipulated the mean predictability/IC of each melody by
352 varying the number of out-of-key notes over the first 9 *context* notes – whereby an out-of-key note
353 results in high IC (Figure 5A). The predictability/IC of the four *target* notes designed to be similar across
354 melodies (Figure 5B). An ANOVA with factors Predictability (P/U) and Note type (Context/Target Notes)
355 on the IC of each note yielded a significant interaction of Predictability and Note type [$F_{(1, 14)} = 11.60, p =$
356 $.004, \eta_p^2 = .45$], indicating that IC for Context but not Target notes differed significantly between
357 predictable and unpredictable melodies [Main effect of predictability on context notes only: $F_{(1, 14)} =$
358 $52.88, p < 0.001, \eta_p^2 = .79$; Main effect of predictability on target notes only: $F_{(1, 14)} = 2.50, p = .136, \eta_p^2 =$
359 $.15$]. Based on the IC measure the 16 melodies were ranked from high to low probability ($M = 4.48 \pm 1.56$,
360 range = 1.7-9.1), and then they were divided into Predictable or Unpredictable based on the median split
361 (Figure 5A).

362 **Figure 5**

363 **4.2.2 Liking ratings and pupillometry**

364 Participants listened to all 16 melodies, one at a time, and rated how much they liked them using a 7-
365 point Likert scale (1 being not all and 7 being very much) at the end of each melody. Liking ratings for
366 each melody were then scaled by subject, and ranked as high, medium and low (Figure 5C).

367 Pupil dilation was measured during Listening using the EyeLink 1000 head-supported infrared optical
368 eye-tracking system (running host software ver. 4.56, SR Research, Ottawa, ON, Canada) in binocular
369 1000Hz sampling configuration, connected to an Apple iMac (Mac OS X 10.12). The EyeLink system was
370 used in the Pupil-Corneal Reflection tracking mode. Participants were seated in a comfortable chair with
371 their head stabilized in a chin and forehead rest, facing the computer monitor (View sonic G225fb 21"
372 CRT, 1024 \times 768pixel resolution, 100 Hz refresh rate, linear gamma correction for luminance with mean
373 luminance = 60 cd/m²) at a distance of 70 cm, in a quiet, moderately lit room (40 cd/m²). To calibrate
374 the eye-tracker, a circular target (1 degree of angle) appeared in random order at one of 6-points on the
375 screen (HV6, in the default Eyelink screen locations), followed by a separate validation. Calibration and
376 validation were repeated until the average error across all calibration targets was below 0.5 degrees of
377 visual angle, and the maximum error at any one calibration point was below 1 degree of visual angle.

378 The order of presentation of the melodies was randomized across participants. Two seconds of baseline
379 pupil data was acquired before and after each melody was played. After each melody, two seconds were
380 given for the liking rating, which was followed by a 2-seconds blank grey screen before the next trial.
381 Participants were instructed to continuously fixate on a cross at the center of the screen (size: 9.4cm x
382 9.4cm, corresponding to a 4.5° visual angle at a viewing distance of 70 cm; RGB: 75,75,75; background
383 grey color, RGB: 150,150,150), not to move their heads for the duration of the eye tracking component
384 of the experiment, and to avoid blinking while the melody was playing. The total duration of the Listening
385 task was approximately 20 minutes.

386 **4.2.3 Melody Learning**

387 In this task participants listened to the first nine *context* notes of each melody through headphones and
388 then played the last four *target* notes on a piano-type keyboard using the four fingers of the right hand:
389 thumb, index, middle and ring finger which were assigned to a fixed white key to which all notes were
390 mapped. This ensured that motor demands for the notes to be played were matched across conditions.
391 Target notes were cued using visual display representing the keyboard (see Figure 1) where a dot
392 appeared sequentially (IOI 428 ms) to indicate which key to play. Participants heard the notes they
393 produced through the headphones. To facilitate accurate playback timing, melodies were accompanied
394 by a metronome beat at the beginning of every bar. The fourth metronome beat cued participants to
395 begin playing back the target notes. The notes played and their timing were recorded from the keyboard
396 and used to score accuracy and synchronization. The 16 melodies were presented in a randomized order,
397 and each melody was repeated 5 times with an ISI of 1 second. At the end of the Learning task,
398 participants performed a final recall block where each of the sixteen melodies was played back once in
399 a random order.

400 Before training, participants were familiarized with the playback task in a brief practice block of four
401 trials in which they had to count 3 metronome beats (corresponding to 3 bars), and then perform 4
402 keypresses cued by the visual display. The familiarization trials contained all finger transitions that were
403 to be encountered in the Melody Learning task. No auditory feedback was provided. Presentation of the
404 melodies and recording of the responses was controlled custom-written Python software running on a
405 PC Linux computer.

406 At the end of the experiment, each participant's music reward sensitivity (i.e. how important music is in
407 his/her life) was assessed via the Barcelona Music Reward Questionnaire ⁷². Participant's intrinsic
408 motivation to perform the task was assessed via the standard, 22-items version of Intrinsic Motivation
409 Inventory ⁷³. The entire experiment lasted approximately 60 minutes.

410 **4.3 Data analysis**

411 **4.3.1 Analysis of pupil diameter**

412 Pupil diameter was measured in arbitrary units. Blinks were identified by identifying in each trial samples
413 without data due to blinks, removing 100 ms before and after the edges of the non-data points to make
414 sure that all artefacts of the pupil size algorithm were removed. For each blink, four equally spaced time
415 points (t_2 = blink onset; t_3 = blink offset; $t_1=t_2-t_3+t_2$; $t_4=t_3-t_2+t_3$) were interpolated by using a cubic-
416 spline fit and the original signal was replaced by the cubic spline, leaving the signal unchanged except
417 for the blink period. Random sample artefacts were removed using a median Hampel filter (from EEGLab
418 software, version 14.1), after which data were smoothed using a Savitzky-Golay Filter over an 11-ms
419 timeframe to remove the high-frequency noise in the pupil without time-delaying the pupil signal. Then,
420 each trial was baseline-corrected against the median pupil size in the 400 ms before the onset of the
421 melody, and then divided in 13 bins corresponding to the onset of each note. Trials during which
422 participants blinked for more than 15% of the total trial duration were excluded (two participants'
423 datasets, and a mean of 0.92 ± 2.53 for the rest of participants). Baseline-corrected pupil size change
424 was then analyzed by using linear mixed-effects regressions testing for the effects of predictability
425 (Predictable/Unpredictable), Liking (High/Medium/Low), time-bins (1-13) and their full interaction.

426 **4.3.2 Analysis of motor performance**

427 Participants' performances were examined off-line to evaluate key errors (MIDI note number) and
428 response times for the four keystrokes relative to the four last target notes of each melody. Trials were
429 considered invalid and excluded from the analyses if participants pressed more or fewer than 4 notes
430 per trial. On these data, two indexes of performance were computed: *trial accuracy* (i) was quantified
431 by counting the total number of errors. These were defined either by an incorrect keystroke (key identity
432 error), or by an absolute response time larger than 428 ms (timing error; for values outside this range,
433 keystrokes occurred within the range of the note preceding or following the one with which it was
434 supposed to be synchronized). *Asynchrony* (ii) was quantified only on correct trials by measuring the
435 time difference between the actual keystroke of a note and the expected onset of that given note.

436 Statistical analyses were performed separately for the two performance indexes (i.e., trial accuracy and
437 asynchrony) and for the training (where each melody was performed for 5 consecutive times) and the
438 test phase (where each melody was played only one time). We used mixed effects regression analyses
439 testing for the effects of predictability (Predictable/Unpredictable), liking (High/Medium/Low),
440 repetition trial (1-4), keystroke (1-4), and the full interaction between liking, predictability and repetition
441 trial. Keystroke was introduced as an effect of no interest to account for known motor execution
442 differences between initiation (first key press) and completion (following 3 keypresses) of sequential
443 movements⁹⁶. Keystrokes for the first repetition trial were initially analyzed, but then excluded because
444 of too many invalid trials ($M= 55.04 \pm 26.60$ % of invalid trials across participants). The analysis on the
445 test phase looked for learning stabilization, and estimated the effects of predictability
446 (Predictable/Unpredictable) and liking (High/Medium/Low).

447 **4.3.3 Statistical analysis**

448 All data analyses were conducted in MATLAB R2015b (Mathworks, Natick, MA, USA), except for the linear
449 mixed model analysis that was implemented in R environment Version 0.99.320 using the 'lmer' function
450 from package lme4 to build the models ⁹⁷ and the ANOVA function from package car to obtain
451 significance tests ⁹⁸. In contrast to a more traditional approach with data aggregation and repeated-
452 measures ANOVA analysis, linear mixed effects regression allows controlling for the variance associated
453 with random factors without data aggregation (see ⁹⁹). By using random effects for subjects and stimuli
454 item, we controlled for the influence of different mean responses associated with these variables.
455 Moreover, we also included by-participant random slopes for the effects of interest (predictability, liking
456 and their interaction), which accounted also for differences in how predictability and liking affected
457 participants' responses (random slopes). Contrasts were carried out using the 'emmeans' package in R
458 ¹⁰⁰. We report unstandardized effect sizes (unstandardized regression coefficients, indicated as 'b' for
459 the statistical tests) which is in line with general recommendations of how to report effect sizes in linear
460 mixed models ¹⁰¹. Significance of the fixed effects of these models were evaluated with the
461 Satterthwaite approximation ¹⁰², and *p* values were adjusted for multiple comparisons using the
462 multivariate t method.

463

464 **5 References**

- 465 1. Pearce, M. T. Statistical learning and probabilistic prediction in music cognition: Mechanisms of
466 stylistic enculturation. *Ann. N. Y. Acad. Sci.* **1423**, 378–395 (2018).
- 467 2. Rohrmeier, M., Rebuschat, P. & Cross, I. Incidental and online learning of melodic structure.
468 *Conscious. Cogn.* **20**, 214–222 (2011).
- 469 3. Tillmann, B. & Lebrun-Guillaud, G. Influence of tonal and temporal expectations on chord
470 processing and on completion judgments of chord sequences. *Psychol. Res.* **70**, 345–58 (2006).
- 471 4. Tillmann, B., Bigand, E. & Madurell, F. F. Local versus global processing of harmonic cadences in
472 the solution of musical puzzles. *Psychol. Res.* **61**, 157–174 (1998).
- 473 5. Guo, S. & Koelsch, S. Effects of veridical expectations on syntax processing in music: Event-
474 related potential evidence. *Sci. Rep.* **6**, 19064 (2016).
- 475 6. Palmer, C. & Pfördresher, P. Q. Incremental planning in sequence production. *Psychol. Rev.* **110**,
476 683–712 (2003).
- 477 7. Bianco, R. *et al.* Syntax in action has priority over movement selection in piano playing: an ERP
478 study. *J. Cogn. Neurosci.* **28**, 41–54 (2016).

479 8. Pfordresher, P. Q., Palmer, C. & Jungers, M. K. Speed, accuracy, and serial order in sequence
480 production. *Cogn. Sci.* **31**, 63–98 (2007).

481 9. Mathias, B., Tillmann, B. & Palmer, C. Sensory, Cognitive, and Sensorimotor Learning Effects in
482 Recognition Memory for Music. *J. Cogn. Neurosci.* **28**, 1111–1126 (2016).

483 10. Huron, D. *Sweet Anticipation : Music and the Psychology of Expectation by David Huron. Sweet*
484 *Anticipation: Music and the Psychology of Expectation.* (2006).

485 11. Zatorre, R. J. & Salimpoor, V. N. From perception to pleasure: music and its neural substrates.
486 *Proc. Natl. Acad. Sci.* **110**, 10430–7 (2013).

487 12. Berlyne, D. E. *Aesthetics and Psychobiology.* (Appleton- Century-Crofts, 1971).

488 13. Salimpoor, V. N. *et al.* Interactions between the nucleus accumbens and auditory cortices
489 predict music reward value. *Science (80-.*). **340**, 216–9 (2013).

490 14. Laeng, B., Eidet, L. M., Sulutvedt, U. & Panksepp, J. Music chills: The eye pupil as a mirror to
491 music's soul. *Conscious. Cogn.* **44**, 161–178 (2016).

492 15. Schultz, W. Multiple reward signals in the brain. *Nat. Rev. Neurosci.* **1**, 199–207 (2000).

493 16. Cleeremans, A., Destrebecqz, A. & Boyer, M. Implicit learning: news from the front. *Trends Cogn.*
494 *Sci.* **2**, 406–416 (1998).

495 17. Saffran, J. R., Johnson, E. K., Aslin, R. N. & Newport, E. L. Statistical learning of tone sequences by
496 human infants and adults. *Cognition* **70**, 27–52 (1999).

497 18. Perruchet, P. & Pacton, S. Implicit learning and statistical learning: one phenomenon, two
498 approaches. *Trends Cogn. Sci.* **10**, 233–8 (2006).

499 19. Pearce, M. T. The Construction and Evaluation of Statistical Models of Melodic Structure in
500 Music Perception and Composition. *Dissertation* 267 (2005).

501 20. Skerritt-Davis, B. & Elhilali, M. Detecting change in stochastic sound sequences. *PLoS Comput.*
502 *Biol.* **14**, 1–24 (2018).

503 21. Hasson, U. The neurobiology of uncertainty : implications for statistical learning Neurobiological
504 approaches to uncertainty and statistical learning : Interfaces and. *Phil. Trans. R. Soc. B.* **372**, 1–
505 12 (2017).

506 22. Friston, K. The free-energy principle: a unified brain theory? *Nat. Rev. Neurosci.* **11**, 127–138
507 (2010).

508 23. Kanai, R., Komura, Y., Shipp, S. & Friston, K. Cerebral hierarchies: Predictive processing, precision
509 and the pulvinar. *Philos. Trans. R. Soc. B Biol. Sci.* **370**, (2015).

510 24. Pezzulo, G. & Cisek, P. Navigating the Affordance Landscape : Feedback Control as a Process
511 Model of Behavior and Cognition. *Trends Cogn. Sci.* **20**, 414–424 (2016).

512 25. Friston, K. *et al.* Active inference and epistemic value. *Cogn. Neurosci.* **6**, 187–224 (2015).

513 26. Monroy, C. D., Meyer, M., Schröer, L., Gerson, S. A. & Hunnius, S. The infant motor system
514 predicts actions based on visual statistical learning. *Neuroimage* **185**, 947–954 (2019).

515 27. Monroy, C. D., Gerson, S. A. & Hunnius, S. Translating visual information into action predictions:
516 Statistical learning in action and nonaction contexts. *Mem. Cogn.* **46**, 600–613 (2018).

517 28. Koelsch, S., Vuust, P. & Friston, K. Predictive Processes and the Peculiar Case of Music. *Trends
518 Cogn. Sci.* **xx**, 1–15 (2018).

519 29. Loui, P. Statistical learning – What can music tell us? *Stat. Learn. Lang. Acquis.* 433–462 (2012).
520 doi:10.1515/9781934078242.433

521 30. Hansen, N. C., Vuust, P. & Pearce, M. 'If you have to ask, you'll never know': Effects of
522 specialised stylistic expertise on predictive processing of music. *PLoS One* **11**, 1–20 (2016).

523 31. Morrison, S. J., Demorest, S. M. & Stambaugh, L. A. Enculturation Effects in Music Cognition. *J.
524 Res. Music Educ.* **56**, 118–129 (2008).

525 32. Hannon, E. E., Soley, G. & Ullal, S. Familiarity overrides complexity in rhythm perception: A cross-
526 cultural comparison of American and Turkish listeners. *J. Exp. Psychol. Hum. Percept. Perform.*
527 **38**, 543–548 (2012).

528 33. Hansen, N. C. & Pearce, M. T. Predictive uncertainty in auditory sequence processing. *Front.
529 Psychol.* **5**, 1052 (2014).

530 34. Daikoku, T. Depth and the Uncertainty of Statistical Knowledge on Musical Creativity Fluctuate
531 Over a Composer's Lifetime. *Front. Comput. Neurosci.* **13**, (2019).

532 35. Drost, U. C., Rieger, M., Brass, M., Gunter, T. C. & Prinz, W. Action-effect coupling in pianists.
533 *Psychol. Res.* **69**, 233–41 (2005).

534 36. Stephan, M. A., Lega, C. & Penhune, V. B. Auditory prediction cues motor preparation in the
535 absence of movements. *Neuroimage* **174**, 288–296 (2018).

536 37. Zatorre, R. J., Chen, J. L. & Penhune, V. B. When the brain plays music: auditory-motor
537 interactions in music perception and production. *Nat. Rev. Neurosci.* **8**, 547–558 (2007).

538 38. Prinz, W. What re-enactment earns us. *Cortex* **42**, 515–517 (2002).

539 39. Novembre, G. & Keller, P. E. A conceptual review on action-perception coupling in the musicians'
540 brain: what is it good for? *Front. Hum. Neurosci.* **8**, 603 (2014).

541 40. Mathias, B., Pfördresher, P. Q. & Palmer, C. Context and meter enhance long-range planning in
542 music performance. *Front. Hum. Neurosci.* **8**, 1–15 (2015).

543 41. Novembre, G. & Keller, P. E. A grammar of action generates predictions in skilled musicians.
544 *Conscious. Cogn.* **20**, 1232–43 (2011).

545 42. Sammler, D., Novembre, G., Koelsch, S. & Keller, P. E. Syntax in a pianist's hand: ERP signatures
546 of 'embodied' syntax processing in music. *Cortex* **49**, 1325–1339 (2013).

547 43. Bianco, R., Novembre, G., Keller, P. E., Villringer, A. & Sammler, D. Musical genre-dependent
548 behavioural and EEG signatures of action planning. A comparison between classical and jazz
549 pianists. *Neuroimage* **169**, 383–394 (2018).

550 44. Vuust, P. & Witek, M. a. G. Rhythmic complexity and predictive coding: a novel approach to
551 modeling rhythm and meter perception in music. *Front. Psychol.* **5**, 1111 (2014).

552 45. Steinbeis, N., Koelsch, S. & Sloboda, J. a. The role of harmonic expectancy violations in musical
553 emotions: evidence from subjective, physiological, and neural responses. *J. Cogn. Neurosci.* **18**,
554 1380–1393 (2006).

555 46. Koelsch, S., Fritz, T. & Schlaug, G. Amygdala activity can be modulated by unexpected chord
556 functions during music listening. *Neuroreport* **19**, 1815–9 (2008).

557 47. Heyduk, R. G. Rated preference for musical compositions as it relates to complexity and
558 exposure frequency. *Percept. Psychophys.* **17**, 84–91 (1975).

559 48. Witek, M. A. G., Clarke, E. F., Wallentin, M., Kringelbach, M. L. & Vuust, P. Syncopation, body-
560 movement and pleasure in groove music. *PLoS One* **9**, (2014).

561 49. Chmiel, A. & Schubert, E. Back to the inverted-U for music preference: A review of the literature.
562 *Psychol. Music* **45**, 886–909 (2017).

563 50. Matthews, T. E., Witek, M. A. G., Heggli, O. A., Penhune, V. B. & Vuust, P. The sensation of
564 groove is affected by the interaction of rhythmic and harmonic complexity. *PLoS One* **14**, 1–17
565 (2019).

566 51. Gold, B. P. *et al.* Musical reward prediction errors engage the nucleus accumbens and motivate
567 learning. *Proc. Natl. Acad. Sci.* **116**, 3310–3315 (2019).

568 52. Salimpoor, V. N., Benovoy, M., Larcher, K., Dagher, A. & Zatorre, R. J. Anatomically distinct
569 dopamine release during anticipation and experience of peak emotion to music. *Nat. Neurosci.*
570 **14**, 257–62 (2011).

571 53. Berridge, K. C., Robinson, T. E. & Aldridge, J. W. Dissecting components of reward: 'liking',
572 'wanting', and learning. *Curr. Opin. Pharmacol.* **9**, 65–73 (2009).

573 54. Ferreri, L. *et al.* Dopamine modulates the reward experiences elicited by music. *Proc. Natl. Acad. Sci.* **116**, 3793–3798 (2019).

575 55. Sara, S. J. The locus coeruleus and noradrenergic modulation of cognition. *Nat. Rev. Neurosci.* **10**, 211–223 (2009).

577 56. Aston-Jones, G. & Cohen, J. D. An integrative theory of locus coeruleus-norepinephrine function: Adaptive Gain and Optimal Performance. *Annu. Rev. Neurosci.* **28**, 403–450 (2005).

579 57. Ferreri, L. & Rodriguez-Fornells, A. Music-related reward responses predict episodic memory performance. *Exp. Brain Res.* **235**, 3721–3731 (2017).

581 58. Weiss, M. W. *et al.* Pupils Dilate for Vocal or Familiar Music. *J. Exp. Psychol. Hum. Percept. Performance.* **42**, 1061 (2016).

583 59. Graham, D. J., Lee, Y. & Gingras, B. The Eye is Listening : Music-Induced Arousal and Individual Differences Predict Pupillary Responses. *Front Hum Neurosci* **9**, 619 (2015).

585 60. Laeng, B., Sirois, S. & Gredeback, G. Pupillometry: A Window to the Preconscious? *Perspect. Psychol. Sci.* **7**, 18–27 (2012).

587 61. David, S. V., Fritz, J. B. & Shamma, S. A. Task reward structure shapes rapid receptive field plasticity in auditory cortex. *Proc. Natl. Acad. Sci.* **109**, 2144–2149 (2012).

589 62. Bao, S., Chan, V. T. & Merzenich, M. M. Cortical remodelling induced by activity of ventral tegmental dopamine neurons. *Nature* **412**, 79 (2001).

591 63. Flöel, A. *et al.* Dopaminergic influences on formation of a motor memory. *Ann. Neurol.* **58**, 121–130 (2005).

593 64. Tremblay, P. L. *et al.* Movement chunking during sequence learning is a dopamine-dependant process: A study conducted in Parkinson's disease. *Exp. Brain Res.* **205**, 375–385 (2010).

595 65. Abe, M. *et al.* Reward improves long-term retention of a motor memory through induction of offline memory gains. *Curr. Biol.* **21**, 557–562 (2011).

597 66. Galea, J. M., Mallia, E., Rothwell, J. & Diedrichsen, J. The dissociable effects of punishment and reward on motor learning. *Nat. Neurosci.* **18**, 597–602 (2015).

599 67. Oudeyer, P. Y. & Kaplan, F. What is intrinsic motivation? A typology of computational approaches. *Front. Neurorobot.* **1**, 1–14 (2007).

601 68. Ryan, R. M. & Deci, E. L. Intrinsic and Extrinsic Motivations: Classic Definitions and New Directions. *Contemp. Educ. Psychol.* **25**, 54–67 (2000).

603 69. Schmidhuber, J. Developmental robotics, optimal artificial curiosity, creativity, music, and the

604 fine arts. *Conn. Sci.* **18**, 173–187 (2006).

605 70. Koelsch, S. Brain correlates of music-evoked emotions. *Nat. Rev. Neurosci.* **15**, 170–180 (2014).

606 71. Salimpoor, V. N., Zald, D. H., Zatorre, R. J., Dagher, A. & McIntosh, A. R. Predictions and the
607 brain: How musical sounds become rewarding. *Trends in Cognitive Sciences* **19**, 86–91 (2015).

608 72. Mas-Herrero, E., Marco-Pallares, J., Loreno-Seva, U., Zatorre, R. J. & Rodriguez-Fornells, A.
609 Individual Differences in Music Reward Experiences. *Music Percept.* **31**, 118–138 (2013).

610 73. McAuley, E., Duncan, T. & Tammen, V. Psychometric properties of the Intrinsic Motivation
611 Inventory in a competitive sport setting: a confirmatory factor Analysis. *Res Q Exerc Sport.* **60**,
612 48–58 (1989).

613 74. Pearce, M. T., Ruiz, M. H., Kapasi, S., Wiggins, G. a & Bhattacharya, J. Unsupervised statistical
614 learning underpins computational, behavioural, and neural manifestations of musical
615 expectation. *Neuroimage* **50**, 302–13 (2010).

616 75. Erola, T. Expectancy-violation and information-theoretic models of melodic complexity. *Empir.
617 Musicol. Rev.* **11**, 2–17 (2016).

618 76. Güçlütürk, Y., Jacobs, R. H. A. H. & Lier, R. van. Liking versus Complexity: Decomposing the
619 Inverted U-curve. *Front. Hum. Neurosci.* **10**, 1–11 (2016).

620 77. Orr, M. G. & Ohlsson, S. Relationship Between Complexity and Liking as a Function of Expertise.
621 *Music Percept. An Interdiscip. J.* **22**, 583–611 (2005).

622 78. Egermann, H., Pearce, M. T., Wiggins, G. A. & McAdams, S. Probabilistic models of expectation
623 violation predict psychophysiological emotional responses to live concert music. *Cogn. Affect.
624 Behav. Neurosci.* **13**, 533–553 (2013).

625 79. Juslin, P. N. & Västfjäll, D. Emotional responses to music: the need to consider underlying
626 mechanisms. *Behav. Brain Sci.* **31**, 559–621 (2008).

627 80. Kang, O. & Wheatley, T. Pupil dilation patterns reflect the contents of consciousness. *Conscious.
628 Cogn.* **35**, 128–135 (2015).

629 81. Bradley, M. B., Miccoli, L. M., Escrig, M. a & Lang, P. J. The pupil as a measure of emotional
630 arousal and automatic activation. *Psychophysiology* **45**, 602 (2008).

631 82. Kuchinke, L., Trapp, S., Jacobs, A. M. & Leder, H. Pupillary responses in art appreciation: Effects
632 of aesthetic emotions. *Psychol. Aesthetics, Creat. Arts* **3**, 156–163 (2009).

633 83. Joshi, S., Li, Y., Kalwani, R. M. & Gold, J. I. Relationships between Pupil Diameter and Neuronal
634 Activity in the Locus Coeruleus, Colliculi, and Cingulate Cortex. *Neuron* **89**, 221–234 (2016).

635 84. Beukema, P. & Verstynen, T. Predicting and binding: interacting algorithms supporting the
636 consolidation of sequential motor skills. *Curr. Opin. Behav. Sci.* **20**, 98–103 (2018).

637 85. Friston, K., Mattout, J. & Kilner, J. Action understanding and active inference. *Biol. Cybern.* **104**,
638 137–60 (2011).

639 86. Tillmann, B., Janata, P., Birk, J. & Bharucha, J. J. Tonal centers and expectancy: Facilitation or
640 inhibition of chords at the top of the harmonic hierarchy? *J. Exp. Psychol. Hum. Percept.
641 Perform.* **34**, 1031–1043 (2008).

642 87. Bianco, R. *et al.* Neural networks for harmonic structure in music perception and action.
643 *Neuroimage* **142**, 454–464 (2016).

644 88. Stephan, M. A., Brown, R., Lega, C. & Penhune, V. Melodic priming of motor sequence
645 performance: The role of the dorsal premotor cortex. *Front. Neurosci.* **10**, 210 (2016).

646 89. Lahav, A., Saltzman, E. & Schlaug, G. Action representation of sound: audiomotor recognition
647 network while listening to newly acquired actions. *J. Neurosci.* **27**, 308–14 (2007).

648 90. Cho, Y. S., Bae, G. Y. & Proctor, R. W. Referential coding contributes to the horizontal SMARC
649 effect. *J. Exp. Psychol. Hum. Percept. Perform.* **38**, 726–734 (2012).

650 91. Schubotz, R. I. Prediction of external events with our motor system: towards a new framework.
651 *Trends Cogn. Sci.* **11**, 211–8 (2007).

652 92. Weinshenker, D. & Schroeder, J. P. There and back again: A tale of norepinephrine and drug
653 addiction. *Neuropsychopharmacology* **32**, 1433–1451 (2007).

654 93. Murty, V. P. & Adcock, R. A. Enriched encoding: Reward motivation organizes cortical networks
655 for hippocampal detection of unexpected events. *Cereb. Cortex* **24**, 2160–2168 (2014).

656 94. Wolpert, D. M., Diedrichsen, J. & Flanagan, J. R. Principles of sensorimotor learning. *Nat. Rev.
657 Neurosci.* **12**, 739–51 (2011).

658 95. Canadian Institutes of Health Research, Natural Sciences and Engineering Research Council of
659 Canada & Social Sciences and Humanities Research Council of Canada. *Tri-council policy
660 statement: Ethical conduct for research involving humans.* (2014).

661 96. Keller, P. E. & Koch, I. The planning and execution of short auditory sequences. *Psychon. Bull.
662 Rev.* **13**, 711–6 (2006).

663 97. Bates, D., Mächler, M., Bolker, B. & Walker, S. Fitting linear mixed-effects models using lme4. *J.
664 Stat. Softw.* **67**, 51 (2015).

665 98. Fox, J. & Weisberg, S. *An {R} Companion to Applied Regression.*, (2011).

666 99. Baayen, R. H., Davidson, D. J. & Bates, D. M. Mixed-effects modeling with crossed random
667 effects for subjects and items. *J. Mem. Lang.* **59**, 390–412 (2008).

668 100. Lenth, R. Emmeans: Estimated Marginal Means, aka Least-Squares Means. R package
669 <Https://CRAN.R-project.org/package=emmeans>, ge version 1.2.1. (2018).

670 101. Pek, J. & Flora, D. B. Reporting effect sizes in original psychological research: A discussion and
671 tutorial. *Psychol. Methods* **23**, 208–225 (2018).

672 102. Luke, S. G. Evaluating significance in linear mixed-effects models in R. *Behav. Res. Methods* **49**,
673 1494–1502 (2017).

674

675 **Competing Interests Statement**

676 The author(s) declare no financial AND non-financial interests competing interests.

677 **6 Author Contributions**

678 **RB** conceived performed and analyzed the experiments; wrote manuscript.
679 **BG** provided expertise and feedback; commented on manuscript draft.
680 **AJ** provided expertise and feedback on pupillometry and analysis; commented on manuscript draft.
681 **VP** supervised and administered the project; secured the funding; conceived the experiments; wrote the
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688 We are grateful to Joe Thibodeau for his assistance in developing the hardware and software interfaces
689 for this experiment. We acknowledge the contribution of Soraya Lalou in collecting the data. We also
690 thank Prof. Robert Zatorre and Dr. Gabriele Chierchia for inspiring discussions, and three anonymous
691 referees for very constructive suggestions.

692 **9 Data Availability Statement**

693 The datasets and the stimuli for this study can be found in the OSF repository (link:
694 https://osf.io/x42sz/?view_only=e75f0dd5b6964c3cb39f603095141885).

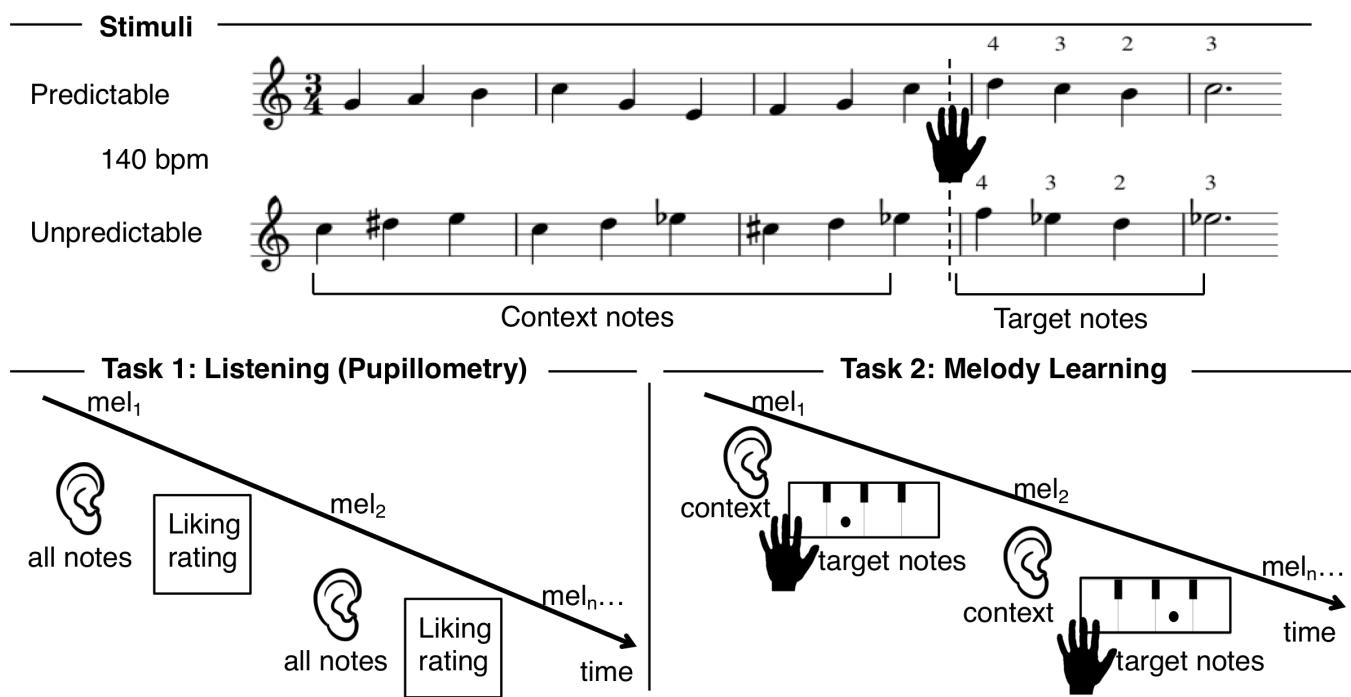
695 **10 Informed Consent Statement**

696 All participants provided written informed consent. The Concordia University Human Research Ethics
697 committee approved the study (30007730) and conducted adhering to the Canadian Tri-council Policy
698 on ethical conduct for research involving humans

699

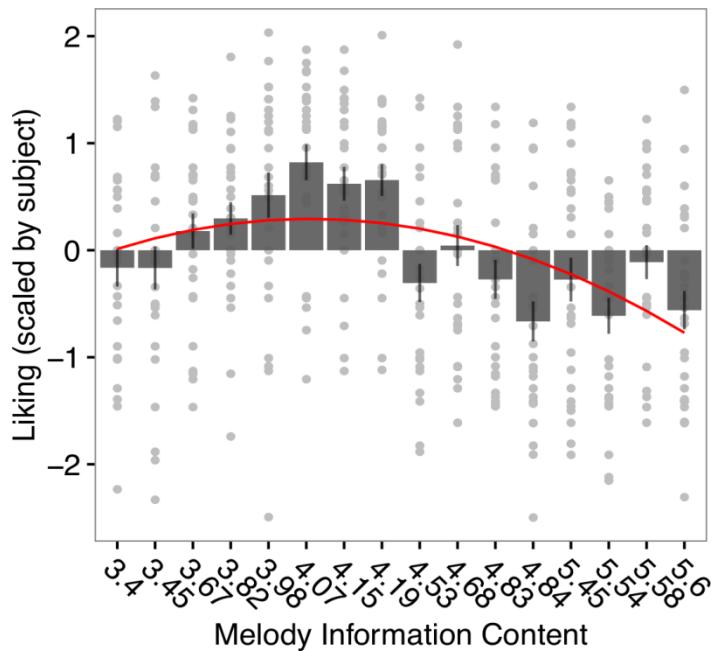
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Figures



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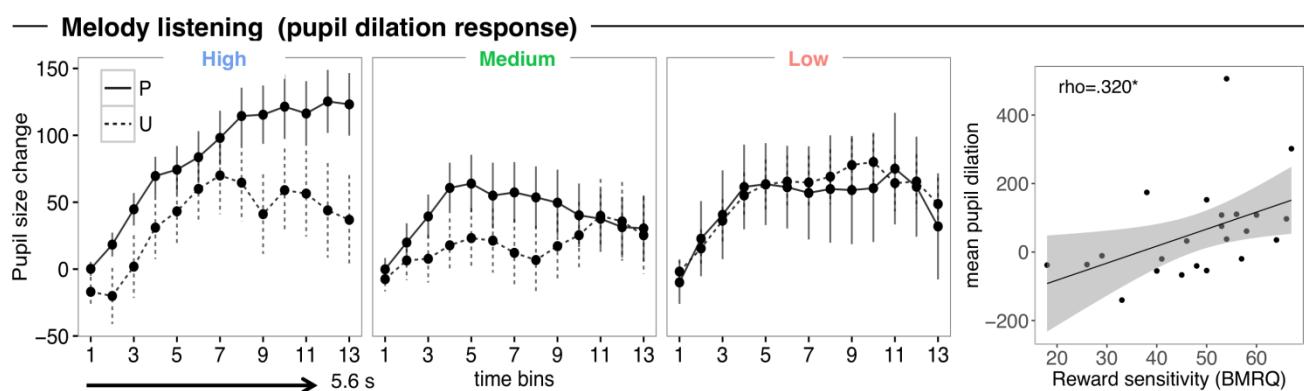
702 **Figure 1.** Schematic illustration of the experimental paradigm. **Stimuli.** The musical scores of 2 examples
703 – 1 more predictable and 1 less predictable based on the information content (IC) of the context notes.
704 Participants listened to the context notes and then played the target notes as guided by the visual
705 display. The IC of the context notes (before the dotted line) was manipulated to result in predictable
706 (low IC) or unpredictable (high IC) contexts. These were followed by four target ending notes with similar
707 IC between predictable and unpredictable melodies. Fingering for the target notes is indicated by the
708 numbers on the last two bars. Thumb, index, middle and ring finger were assigned to a fixed white key
709 to which different expected sounds were artificially mapped to. This ensured that motor demands for
710 the target ending notes were matched across conditions. **Task 1.** Listening and pupillometry: pupil
711 dilation was measured while participants listened to the entire melody and liking ratings (7-point scale)
712 were collected at the end of each trial. **Task 2.** Melody Learning: participants listened to the first nine
713 context notes of the melody and completed the melody by playing the last four target notes on a midi-
714 keyboard. The notes expected to play were cued by sequential dots drawn onto a keyboard on the
715 screen. Each note occurred at a tempo of 140 bpm, and a metronome sound at 46 bpm (every three
716 notes) guided participants' pace. Each trial was repeated 5 times during training and 1 time in a final
717 test-phase.



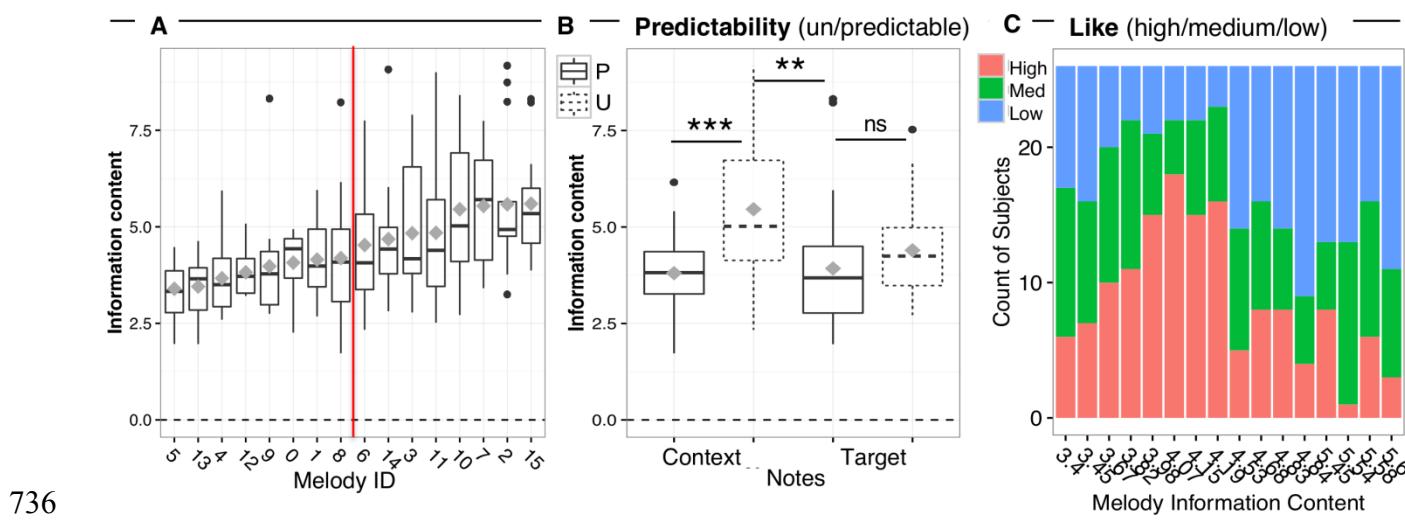
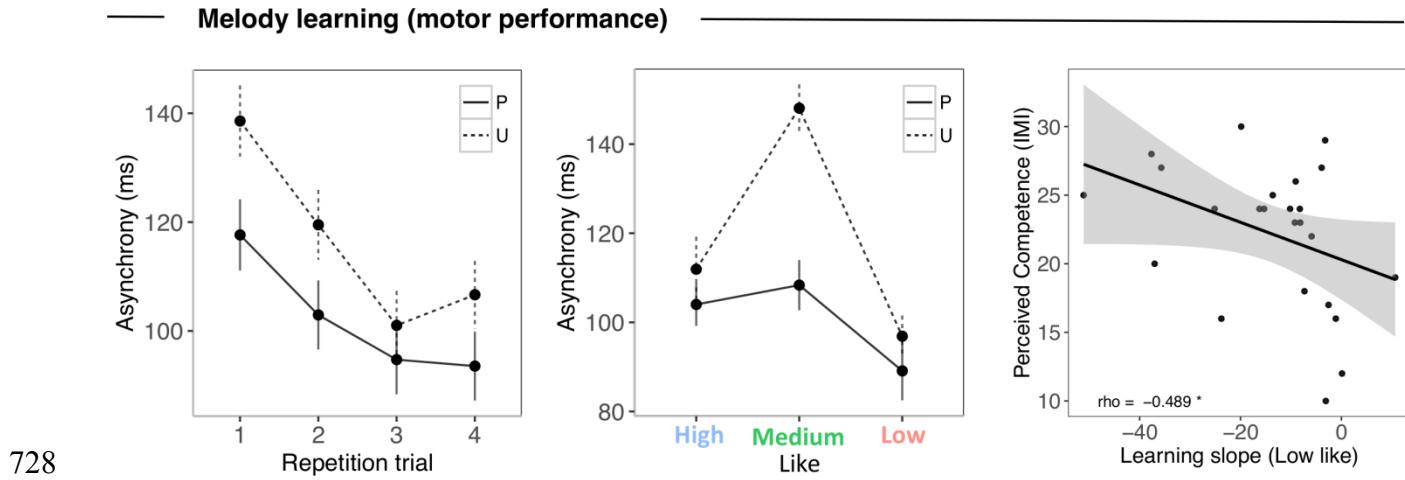
718

719 **Figure 2.** Inverted-U relationship between mean information content (IC) of each melody (ordered by
720 increasing mean IC values on the x axis) and subjective liking ratings (scaled by subjects). Each point
721 represents individual ratings for each melody. Error bars represent 1 s.e.m. of ratings for each melody.

722



724 **Figure 3.** Pupil size change across 13 time-bins (for each note) for predictable (P) and unpredictable (U)
725 melodies across different degrees of liking (high, medium and low). Error bars represent 1 s.e.m. of all
726 trials. (right panel) Scatter Plot showing the mean pupil dilation as a function of reward sensitivity score.
727 Each data point represents an individual participant. The diagonal indicates the line of best fit.



743