

## 2 Implementation of structure-mapping inference by event-file 3 binding and action planning: a model of tool-improvisation 4 analogies

5 Chris Fields

6 Received: 19 March 2010 / Accepted: 18 May 2010  
7 © Springer-Verlag 2010

8 **Abstract** Structure-mapping inferences are generally  
9 regarded as dependent upon relational concepts that are  
10 understood and expressible in language by subjects capable  
11 of analogical reasoning. However, tool-improvisation infer-  
12 ences are executed by members of a variety of non-human  
13 primate and other species. Tool improvisation requires cor-  
14 rectly inferring the motion and force-transfer affordances of  
15 an object; hence tool improvisation requires structure map-  
16 ping driven by relational properties. Observational and  
17 experimental evidences can be interpreted to indicate that  
18 structure-mapping analogies in tool improvisation are  
19 implemented by multi-step manipulation of event files by  
20 binding and action-planning mechanisms that act in a lan-  
21 guage-independent manner. A functional model of lan-  
22 guage-independent event-file manipulations that implement  
23 structure mapping in the tool-improvisation domain is  
24 developed. This model provides a mechanism by which  
25 motion and force representations commonly employed in  
26 tool-improvisation structure mappings may be sufficiently  
27 reinforced to be available to inwardly directed attention and  
28 hence conceptualization. Predictions and potential experi-  
29 mental tests of this model are outlined.

### 30 Introduction

31 Analogical inference involves recognizing aspects of a  
32 remembered situation that are interesting like aspects of a  
33 novel situation, and applying knowledge of relations hold-  
34 ing in the remembered situation to explain behavior in or  
35 make predictions about the novel situation. Analogies are

distinguished by, and their often impressive explanatory 36  
power results from, the recognition and inferential use of 37  
similarities in relational structure between remembered and 38  
novel situations, as opposed to or in addition to similarities 39  
in the surface properties of the objects involved in the situa- 40  
tions (reviewed by Gentner, 2003; Holyoak, 2005). In con- 41  
ceptual analogies presented in language, the inferential 42  
steps of recognizing the structural similarity between a 43  
remembered “base” or “source” situation and a novel “tar- 44  
get” situation and then mapping the relational structure of 45  
the source situation onto the target situation are experimen- 46  
tally separable; the recognition step involves a frontal-pari- 47  
etal working memory (WM) network (Green, Fugelsang, 48  
Kraemer, Shamosh, & Dunbar, 2006), while the mapping 49  
step involves regions of rostral prefrontal cortex (RPFC; 50  
Green et al., 2006; Morrison et al., 2005) that are also 51  
implicated in multi-tasking (Dreher, Koehlin, Tierney, & 52  
Grafman, 2008; Sigman & Dehaene, 2006) and allocating 53  
attention between externally driven perception and internal 54  
imaginative processes (Burgess, Simons, Dumontheil, & 55  
Gilbert, 2007; Gilbert, Frith, & Burgess, 2005). 56

Structure-mapping inferences are typically explicated in 57  
terms of manipulations of relational concepts expressible in 58  
language. Gentner (2003) places relational concepts 59  
expressible in language at the center of analogical capabil- 60  
ity, claiming that “acquisition of relational language is 61  
instrumental in the development of analogy” (p. 219). 62  
Gentner and Christie (2008) advance the arguably stronger 63  
claim that “possession of an elaborated symbol system— 64  
such as human language—is necessary to make our rela- 65  
tional capacity operational” (p. 136). The dependence of 66  
analogical capability on relational language capability is 67  
evident in young children, who become progressively more 68  
able to recognize analogies between situations as their 69  
relational vocabularies increase and the meanings they 70

A1 C. Fields (✉)  
A2 Apdo. 363-4013, Atenas 20501, Costa Rica  
A3 e-mail: fieldsres@gmail.com

71 attach to relational terms approach the meanings generally  
 72 assigned by adults (reviewed by Gentner, 2005). Consistent  
 73 with the view that recognition of the relational similarities  
 74 that drive structure mapping depends upon relational concepts  
 75 expressible in language, analogies presented in language  
 76 dominate research on the mechanisms of structure mapping.  
 77 The experimental design employed by Green et al. (2006)  
 78 to functionally localize the analogical mapping process, for  
 79 example, depends on the manipulation of language-based  
 80 semantic relations. When analogies between pictures are  
 81 used experimentally, for example by Morrison et al. (2005),  
 82 the interpretation of the results typically relies on the  
 83 assumption that subjects are retrieving concepts expressible  
 84 in language to interpret the pictures as structurally  
 85 analogous.


86 While non-human animals are clearly capable of recognizing  
 87 similarities between situations, they are generally regarded  
 88 as being incapable of recognizing analogies. Penn et al. (2008),  
 89 for example, argue that non-human animals are incapable  
 90 of true analogical reasoning, i.e., reasoning in which  
 91 similarities between relations holding in two situations,  
 92 not similarities between surface features of objects, provide  
 93 the basis for an inference that one situation is like another.  
 94 They attribute this lack of analogical ability to an inability  
 95 to represent and carry out inferences about relations,  
 96 concluding that “only humans are able to reason about  
 97 higher-order relations in a structurally systematic and  
 98 inferentially productive fashion” (p. 128). Gentner (2003)  
 99 reviews evidence that chimpanzees are capable of symbolic  
 100 relation-matching tasks only if given specific training in the  
 101 use of symbols. She concludes that chimpanzees are capable  
 102 of relational reasoning, but can perform it “only if they  
 103 learn relational language” (p. 219). The common denominator  
 104 between these analyses is the claim that explicit representations  
 105 of the relations holding in pairs of situations, whether in a  
 106 natural language (Gentner, 2003) or in a “language of  
 107 thought” supporting reinterpretation of perceived relations  
 108 between particular entities as instances of conceptualized  
 109 abstract relations (Penn et al., 2008), are required for  
 110 structure mapping driven by relational similarity.

111 This paper challenges the claim that concepts expressible  
 112 in language—either a public language or a language of  
 113 thought—are prerequisites for inference by structure mapping.  
 114 It focuses on a particular class of inferences from a  
 115 remembered to a novel situation that are performed by both  
 116 humans and non-human animals and that appear *prima facie*  
 117 to involve relational knowledge: the inferences involved  
 118 in spontaneous tool improvisation. The improvisation or  
 119 invention of a novel tool to support a goal-driven activity,  
 120 previously performed using only parts of an animal’s own  
 121 body, requires the construction of a novel action plan in  
 122 which the motions and forces required to use the tool  
 123 replace the motions and forces previously employed.

“Introduction” reviews the phenomenology of tool improvisation  
 both in mammalian and in avian species, and shows that tool-  
 improvisation inferences are instances of structure mapping in  
 which the structures being mapped are goal-directed action plan  
 templates that encode both kinematic (specifying motion) and  
 dynamic (specifying force transfer) relations between objects.  
 The broad phylogenetic distribution of tool improvisation  
 suggests that such inferences may be the most ancient instances  
 of structure mapping, and that the highly developed capability  
 for structure mapping observed in humans may be significantly  
 based on an ancient capability broadly shared across species,  
 but restricted in its application, in non-humans, to tool  
 improvisation. “Structure mapping in tool improvisation”  
 reviews data indicating that the structure-mapping inferences  
 supporting tool improvisation are implemented by event-file  
 binding (Hommel, 2004) and pre-motor action planning  
 (Johnson-Frey, Newman-Norland, & Grafton, 2005; Lewis,  
 2006) networks that are substantially shared by humans and  
 macaques. In contrast, conscious simulation-based evaluation  
 and comparison of action plans, as well as the ability to  
 experience and hence to report that two action plans are  
 analogous, depend on attention-switching functions of  
 RPFPC that are evolutionarily recent and probably human-  
 specific (Burgess et al., 2007). A functional model of  
 structure-mapping inferences in the tool-improvisation domain  
 is proposed that requires manipulation of event files and  
 pre-motor action planning, but not conscious conceptual  
 understanding of motions or forces. “Consequences of the  
 event-file manipulation model: functional dependence of  
 motion concepts on structure mapping” shows that in this  
 event-file manipulation model of structure mapping in tool  
 improvisation, the direction of functional dependency is  
 reversed from that claimed by Gentner (2003, 2005) and by  
 Penn et al. (2008): kinematic and dynamic concepts  
 expressible in human language require, instead of being  
 required by, the capability for inference by structure  
 mapping. This proposal is consistent with the hypothesis that  
 human language-based concepts are at least partially  
 derived from pre-existing visuo-motor representations  
 (Barsalou, 2008; Fiebach & Schubotz, 2006; Gallese &  
 Lakoff, 2005). Both anecdotal and experimental evidences  
 supporting this conjecture are discussed. “Testing the  
 proposed model of tool-improvisation structure-mapping  
 inferences” outlines a number of predictions derived from the  
 proposed model of structure-mapping inferences, and  
 reviews observations bearing on them.

### Structure mapping in tool improvisation

Humans, chimpanzees (Whiten et al., 2001), orangutans  
 (van Schaik et al., 2003), gorillas (Breuer, Ndoundou-

	Large 426	290	xxxx	Dispatch: 29.5.10	No. of Pages: 14
	Journal	Article	MS Code	LE <input type="checkbox"/>	TYPESSET <input type="checkbox"/>
				CP <input checked="" type="checkbox"/>	DISK <input checked="" type="checkbox"/>

174 Hockemba, & Fishlock, 2005), capuchin monkeys (Ottoni,  
175 Dogo de Resende, & Izar, 2005), bottlenose dolphins  
176 (Krutzen, Mann, Heithaus, Connor, Bejder, & Sherwin,  
177 2005), elephants (Byrne, Bates, & Moss, 2009), crows  
178 (Hunt & Grey, 2003, 2004) and finches (Tebich & Bshary,  
179 2004) exhibit tool improvisation in the wild. The most  
180 familiar tools of any animal are its own limbs, and the most  
181 fundamental cases of tool improvisation involve using an  
182 object common to the animal's environment to augment the  
183 reach or force of an animal's limbs. New Caledonian crows  
184 employ manufactured hooks to extend the reach of their  
185 beaks (Hunt & Grey, 2003, 2004), modifying them as  
186 needed for particular tasks (Weir & Kacelnik, 2006).  
187 Woodpecker finches use cactus spines and sticks as probing  
188 tools (Tebich & Bshary, 2004). Bottlenose dolphins adapt  
189 sponges as head-mounted fishing tools (Krutzen et al.,  
190 2005; Mann et al., 2008). Elephants manufacture and  
191 employ tools for personal hygiene (Byrne et al., 2009).  
192 Capuchin monkeys use stones to crack nuts (Ottoni et al.,  
193 2005; Visalberghi, Fragaszy, Ottoni, Izar, de Oliveira, &  
194 Andrade, 2007). Gorillas use stout sticks as walking sticks,  
195 canes and bridges (Breuer et al., 2005). Chimpanzees and  
196 orangutans use many kinds of objects as tools, modifying  
197 them as needed (Sanz & Morgan, 2007; van Schaik et al.,  
198 2003; Whiten et al., 2001); distinct choices of objects to  
199 employ as tools and distinct methods and objectives of tool  
200 use among these primates are among the principal markers  
201 of community-specific cultures in wild primate communi-  
202 ties (reviewed by Whiten & van Schaik, 2007), as they are  
203 among humans. Paleo-anthropological evidence indicates  
204 proto-human use of modified stone tools from at least  
205 2.5 million years ago (Plummer, 2004; Wynn, 2002). Mod-  
206 ern humans immersed in a tool-rich technological culture  
207 continue to practice tool improvisation, from the cobbling  
208 together of prototypes of new technologically sophisticated  
209 tools to meet novel requirements to the casual use of screw-  
210 driver handles, crowbars or suitable stones in the place of  
211 forgotten hammers.

212 The inference that a novel object A can functionally  
213 substitute for a more familiar object B in the context of a  
214 goal-directed action is non-trivial. Consider the case of  
215 capuchins (Ottoni et al., 2005; Visalberghi et al., 2007) or  
216 chimpanzees (Biro, Inoue-Nakamura, Tonooka, Yamakoshi,  
217 Sousa, & Matsuzawa, 2003; Carvalho, Cunha, Sousa, &  
218 Matsuzawa, 2008) using stones to crack nuts. Both species  
219 are familiar with food sources with husks and peels, and  
220 with the removal of these coverings with the hands, but  
221 their hands are not capable of removing the hard shells of  
222 nuts. Some individuals of both species are observed to  
223 select stones from the local environment and use them as  
224 tools to crack nuts so that the shells can be removed. Tool-  
225 using individuals are capable of selecting from among mul-  
226 tiple stones those that are appropriate for use as tools

(Carvalho et al., 2008; Schrauf, Huber, & Visalberghi, 2008; Visalberghi et al., 2007, 2009). Young individuals of both species learn, through a combination of observation of older stone-using conspecifics and practice, to select stones appropriate for use in cracking nuts from a variety of available candidates, and to execute the positioning and striking motions necessary to crack nuts with the selected stones. Tool-using individuals do not merely learn that specific stones are useful as tools, but rather that stones with particular properties, including size, shape, weight and hardness, are useful as tools (Carvalho et al., 2008; Schrauf et al., 2008; Visalberghi et al., 2009). While these primates do not modify stones used for nut cracking, chimpanzees do modify other tools (Whiten et al., 2001) including pointed sticks used as spears (Pruetz & Bertolani, 2007) and concrete disks used as projectiles (Osvath, 2009). Orangutans (van Schaik et al., 2003), gorillas (Breuer et al., 2005), elephants (Byrne et al., 2009) and crows (Hunt & Grey, 2004; Weir & Kacelnik, 2006) also modify tools. Selection of potential tools using general and functionally relevant criteria, modification of selected objects to better satisfy functionally relevant criteria and learning of group-specific tool selection and use practices (Whiten & van Schaik, 2007) all indicate that non-human animal tool use involves non-trivial causal inferences as opposed to simple associations (Penn & Povinelli, 2007). Hence while available evidence does not support the claim that non-human animals understand concepts, such as applied force in the abstract (Penn, Holyoak & Povinelli, 2008; Penn & Povinelli, 2007), it does support the claim that, at least in tool-improvisation contexts, they execute inferences that require representations of physical parameters, such as size, weight, flexibility, tensile strength and sharpness that are relevant to the functioning of tools.

260 The existence of distinct tool-use cultures in neighboring  
261 bands of chimpanzees (Biro et al., 2003; Sanz & Morgan,  
262 2007) indicates that tool improvisation by individual chimpanzees is not uncommon. The first, "discovery" instance of using a novel tool need not involve a structure-mapping inference: a lucky capuchin or chimpanzee might, for example, fortuitously drop a rock onto a nut and crack it, revealing a food source inside. Positive affective tags associated with food discovery would be expected to increase the likelihood that such an event would be remembered. However, incorporating the remembered event into the repertoire of food-seeking action patterns requires inference; in the nut cracking case, it requires linking the goal of obtaining food to both the novel source and to the sequential actions of searching for an appropriate stone to use as a cracking tool and manipulating it in an appropriate way. While some experiments have been interpreted as indicating planning based on experienced episodic memories in chimpanzees and orangutans (Osvath & Osvath, 2008), most observations do not support such capabilities in non-

280 human animals (Suddendorf & Corballis, 2007; Sudden- 333  
 281 dorf, Corballis, & Collier-Baker, 2009). Inference from a 334  
 282 chance discovery, or from observation of tool use by a 335  
 283 mentor, is probably unconscious and automated, in capu- 336  
 284 chins even if not in chimpanzees. That such inferences are 337  
 285 non-trivial is indicated by the fact that multiple demonstra- 338  
 286 tions are typically required for learning behaviors, such as 339  
 287 nut cracking in both species (Biro et al., 2003; Marshall- 340  
 288 Pescini & Whiten, 2009; Ottoni et al., 2005). The primary 341  
 289 hypothesis of this paper is that the construction of novel 342  
 290 goal-directed action patterns involving tool use is accom- 343  
 291 plished by a particular kind of unconscious, but non-trivial 344  
 292 inference: structure mapping. 345

293 From a phenomenological perspective, tool-improvisa- 346  
 294 tion inferences satisfy the definitional criteria of structure 347  
 295 mapping. Nut-cracking capuchins or chimpanzees, for 348  
 296 example, appear to execute a structure-mapping analogy 349  
 297 stone:nut::hand:fruit. The source case for this analogy is an 350  
 298 action plan—hold the fruit so that it does not move and 351  
 299 remove the covering of the fruit by movements of the 352  
 300 hand—that has a specific goal, obtaining the food inside 353  
 301 the fruit. The target case is a similar action plan—secure 354  
 302 the nut so that it does not move and remove the covering by 355  
 303 movements of the hand holding the stone—with a similar 356  
 304 specific goal, obtaining the food inside the nut. When the 357  
 305 action encoded by either of these action plans is executed 358  
 306 successfully, the food that was previously hidden is 359  
 307 exposed and visible. Thus, source and target cases share (1) 360  
 308 their application to objects containing food; (2) their encod- 361  
 309 ings as action plans that involve visually coordinated force- 362  
 310 ful hand movements; (3) their goals of obtaining the hidden 363  
 311 food contained in the objects to which they are applied; and 364  
 312 (4) their observable successful outcomes of making visible 365  
 313 what was previously invisible. They differ in the details of 366  
 314 the objects to which they are applied, the hand movements 367  
 315 that are employed, and what the dominant hand is holding: 368  
 316 nothing in one case and a stone in the other. In the context 369  
 317 of the action plan, this last difference is encoded by differ- 370  
 318 ences in muscle configurations and movements and by two 371  
 319 parameters: the felt weight of the hand grasping the stone, 372  
 320 and the force required to move that weighted hand with 373  
 321 sufficient velocity to crack the nut (Brill, Dietrich, Foucart, 374  
 322 Fuwa, & Hirata, 2009). Mappings between source and tar- 375  
 323 get cases that preserve long-range organizing relations, 376  
 324 such as goals or outcomes while allowing variations in the 377  
 325 superficial details of objects and motions and in the values 378  
 326 of properties and parameters are structure mappings 379  
 327 (Gentner, 2003; Holyoak, 2005). Tool-improvisation analo- 380  
 328 gies in general share these defining characteristics of 381  
 329 structure mappings. 382

330 Non-trivial analogies are not just structure-mapping 383  
 331 inferences, but structure-mapping inferences in which 384  
 332 relations, not surface similarities, carry the inferential 385

weight (Gentner, 2003; Holyoak, 2005). Thus, it might 333  
 be objected that tool-improvisation inferences, while 334  
 qualifying as structure mappings, fail to qualify as analo- 335  
 gies because they are driven by surface similarities, not 336  
 relational similarities. This is not, however, the case. 337  
 Stones, for example, have few surface similarities with 338  
 hands, and do not functionally substitute for hands in 339  
 contexts involving grasping, manipulating, climbing, 340  
 grooming or locomotion. Stones only functionally substi- 341  
 tute for hands in contexts that call for a tool or a weapon, 342  
 i.e., contexts that involve the application of mechanical 343  
 force to another object. Utility for the application of 344  
 mechanical force is a relational criterion. While there is 345  
 no evidence that primates other than humans understand 346  
 this criterion in the abstract (Penn et al., 2008), the 347  
 marked preferences of both chimpanzees (Carvalho 348  
 et al., 2008) and capuchins (Schrauf et al., 2008; 349  
 Visalberghi et al., 2009) for stones with shapes, weights 350  
 and hardness suitable to the dynamic requirements of nut 351  
 cracking indicates that they are sensitive to this relational 352  
 requirement. The centrality of relational requirements 353  
 involving force (i.e., weight), tensile strength, rigidity 354  
 and particular details of shape is a general feature of tool- 355  
 improvisation structure mappings. Non-human animals, 356  
 like humans, select objects for use as tools that satisfy 357  
 functional criteria, not objects that merely share surface 358  
 features. Gorillas, for example, test branches or sticks for 359  
 strength before using them as supports (Breuer et al., 360  
 2005). Crows modify twigs so that the final shape differs 361  
 from the original shape in ways that contribute to func- 362  
 tion (Hunt & Grey, 2004). Chimpanzees sharpen sticks to 363  
 be used as spears with their teeth, achieving impressive 364  
 points (Pruetz & Bertolani, 2007). Tamarin monkeys, 365  
 although they apparently do not use tools in the wild, 366  
 differentiate functionally relevant from functionally irrel- 367  
 evant features of candidate tools in captivity, even in 368  
 infancy (Hauser, Pearson, & Seelig, 2002). The objects 369  
 that are selected as satisfying tool-improvisation struc- 370  
 ture mappings are thus selected, or selected and then 371  
 modified, on the basis of criteria directly relevant to the 372  
 principle organizing relation of the structure mapping, 373  
 the utility of the object employed as a tool in achieving 374  
 the result that motivates the structure-mapping inference. 375  
 Tool-improvisation structure mappings therefore qualify 376  
 as analogies in the strict sense of inferences driven by 377  
 relational similarities, not surface similarities. As dis- 378  
 cussed above, selection and modification on the basis of 379  
 functional, relational criteria do not imply conscious 380  
 understanding of these criteria, or of the concepts of 381  
 force or of utility to achieve an end in the abstract, but do 382  
 imply at least an implicit representation of such criteria, 383  
 and do require that these criteria trump functionally irrel- 384  
 evant surface similarities in the selection process. 385

386 Tool-improvisation analogies executed by non-human,  
387 and therefore language-lacking animals pose both a diffi-  
388 culty and an opportunity for functional models of structure  
389 mapping. The difficulty is that existing models of structure  
390 mapping depend on the manipulation of concepts express-  
391 ible in language, either a public natural language or an  
392 internal, comprehended language of thought. The opportu-  
393 nity is that the representation of tool use in the primate  
394 brain is considerably better understood than the representa-  
395 tion of abstract conceptual reasoning; hence tool-improvi-  
396 sation analogies may provide insights into how brains  
397 implement structure mappings, at least those structure map-  
398 pings that depend on kinematic and dynamic relations  
399 between objects.

#### 400 **Neurocognitive implementation of structure mappings** 401 **for tool use: evidence and functional model**

402 Non-human animals lack human language; they must there-  
403 fore implement tool-improvisation analogies with neuro-  
404 cognitive mechanisms that do not rely on human language.  
405 This requirement has two parts: first, non-human animals  
406 must have non-language-based representations of the goals,  
407 objects and action plans involved both in the source and in  
408 the target cases; second, they must have a non-language-  
409 dependent inferential mechanism capable of executing  
410 structure-mapping inferences, at least for source and target  
411 cases in the tool-use domain. Humans do have human lan-  
412 guage, and clearly execute analogies, such as the political  
413 analogies described by Holyoak (2005), that appear to be  
414 explicable only in terms of language-dependent inferences.  
415 One can, however, ask also in the case of humans how  
416 goals, objects and action plans involved specifically in tool  
417 use are represented, and how structure-mapping inferences  
418 specifically involving tool improvisation are executed.

419 To develop a language-independent model of tool-  
420 improvisation inferences, it is useful to consider the neuro-  
421 cognitive implementation of tool-use actions and action  
422 planning. A considerable body of experimental evidence  
423 indicates that humans represent actions involving tools in a  
424 left-hemisphere-dominated praxis network that includes  
425 posterior-parietal multi-modal binding areas, somatosen-  
426 sory areas and premotor areas (reviewed by Culham &  
427 Valyear, 2006; Johnson-Frey et al., 2005; Lewis, 2006;  
428 Martin, 2007). This frontoparietal network is activated not  
429 only by performing actions with tools, but also by panto-  
430 miming actions with tools and imagining actions with tools.  
431 It overlaps significantly with the mirror-neuron system  
432 (MNS) that maps observations of others performing motor  
433 acts onto motor plans (reviewed by Puce & Perrett, 2003;  
434 Rizzolatti & Craighero, 2004). Mirror neurons respond to  
435 non-biological motions that are kinematically similar to

biological motions, such as motions of reaching or pound- 436  
ing tools, as well as to biological motions (Engel, Burke, 437  
Fiehler, Bien, & Rosler, 2007; Schubotz & van Cramon, 438  
2004); rigid tool motions are represented separately within 439  
the praxis network (Martin, 2007). Planning tool use cou- 440  
ples this frontoparietal action representation to areas of lat- 441  
eral prefrontal cortex involved in learning motor responses 442  
to visual stimuli (Boettiger & D'Esposito, 2005), maintain- 443  
ing representations of task requirements as motions are exe- 444  
cuted (Cole & Schneider, 2007; Courtney, 2004; Tanji & 445  
Hoshi, 2008), and associating task requirements with pre- 446  
motor-encoded information about movement capabilities 447  
(Johnson-Frey et al., 2005). Increasing the complexity of 448  
tool-use actions increases activation of more rostral areas of 449  
prefrontal cortex, as demonstrated in experiments in which 450  
novices (Stout & Chaminade, 2007) and experts (Stout 451  
et al., 2008) manufactured replicas of early stone-age tools. 452  
Relatively simple motions used by novices to construct rel- 453  
atively simple stone tools activated the frontoparietal net- 454  
work supporting perceptual control of motor actions, but 455  
not prefrontal executive areas (Stout & Chaminade, 2007), 456  
while the more complex sequences of motions used by 457  
experts to construct more sophisticated tools activated both 458  
lateral and rostral prefrontal areas, including language-pro- 459  
duction areas (Stout et al., 2008; Stout & Chaminade, 460  
2009). 461

Comparative studies of human and non-human primate 462  
tool use indicate broad similarities in the encoding of tool- 463  
use actions across primates. Tool use both in macaque 464  
monkeys and in humans leads to specificity changes in 465  
interparietal sulcus (IPS) neurons implementing visual to 466  
somatosensory binding that effectively extend the body to 467  
incorporate the tool (reviewed by Maravita & Iriki, 2004), 468  
while maintaining a body-tool distinction (Povinelli, 469  
Reaux, & Frey, 2009). Monkey and human IPS are highly 470  
anatomically and functionally homologous, implementing 471  
multi-modal sensory binding to construct spatial layouts, 472  
binding action plans to the representations of such layouts, 473  
and controlling motions relevant to objects in a layout 474  
(reviewed by Grefkes & Fink, 2005). Mirror neurons spe- 475  
cific to observations of tool use have been identified in 476  
macaque monkeys (Ferrari, Rozzi, & Fogassi, 2005). The 477  
specificities of these tool-use-specific mirror neurons 478  
develop slowly over months of training and experience 479  
with tool-like objects, consistent with both the time course 480  
of tool-use learning in wild primates (Biro et al., 2003; 481  
Otoni et al., 2005) and the general plasticity of mirror-neu- 482  
ron specificities observed in humans (Catmur, Gillmeister, 483  
Bird, Liepelt, Brass, & Heyes, 2008; Catmur, Walsh, & 484  
Heyes, 2007). Multi-step actions are planned, sequenced 485  
and controlled by areas of lateral prefrontal cortex in 486  
macaques as they are in humans (reviewed by Hoshi, 487  
2006); in macaques lateral prefrontal cortex appears to 488

489 encode control for all action sequences regardless of com- 542  
 490 plexity (reviewed by Tanji & Hoshi, 2008) with more ros- 543  
 491 tral prefrontal cortex reserved to decision-making based on 544  
 492 affective and sensory (primarily olfactory) cues (Averbach 545  
 493 & Seo, 2008).

494 While activation of the frontoparietal praxis network by 546  
 495 tool-improvisation inferences has not been observed 547  
 496 directly, the involvement of this network in imagining and 548  
 497 planning tool use (Lewis, 2006) indicates that it would be 549  
 498 active in tool-improvisation inferences if they involve 550  
 499 either imagining or planning tool use. The overlapping, 551  
 500 multimodal nature of the representation of tool-use actions 552  
 501 and tool-use planning in the frontoparietal praxis network 553  
 502 indeed suggests that this network itself may implement 554  
 503 structure-mapping inferences in the tool-use domain. 555  
 504 A functional model of the implementation of two structure- 556  
 505 mapping inferences, the stone:nut::hand:fruit analogy 557  
 506 discussed above and the common human backpacker's 558  
 507 tool-improvisation analogy stone:tent-stake::hammer:nail, 559  
 508 based on their implementation by the praxis network is 560  
 509 shown in Fig. 1. This model proposes that (1) the represen- 561  
 510 tational structures that are mapped in tool-improvisation 562  
 511 analogies are event files (Hommel, 2004) implemented as 563  
 512 activation patterns centered on IPS; and (2) mapping of 564  
 513 event files is executed in two phases by two distinct binding 565  
 514 processes. The first of these processes involves retrieval of 566  
 515 an action instance or minimally abstracted action schema 567  
 516 that serves as the source case, and induces mapping of the 568  
 517 object and motion components of the task environment into 569  
 518 a source-case-based action plan. The second process 570  
 519 involves the embedding of additional action components 571  
 520 into the partially mapped action plan, and induces mapping 572  
 521 of the tool components of the task environment to create a 573  
 522 fully mapped target-case action plan. In the final step of this 574  
 523 second process, the fully mapped target-case action plan is 575  
 524 executed, confirming or disconfirming the adequacy of the 576  
 525 structure mapping.

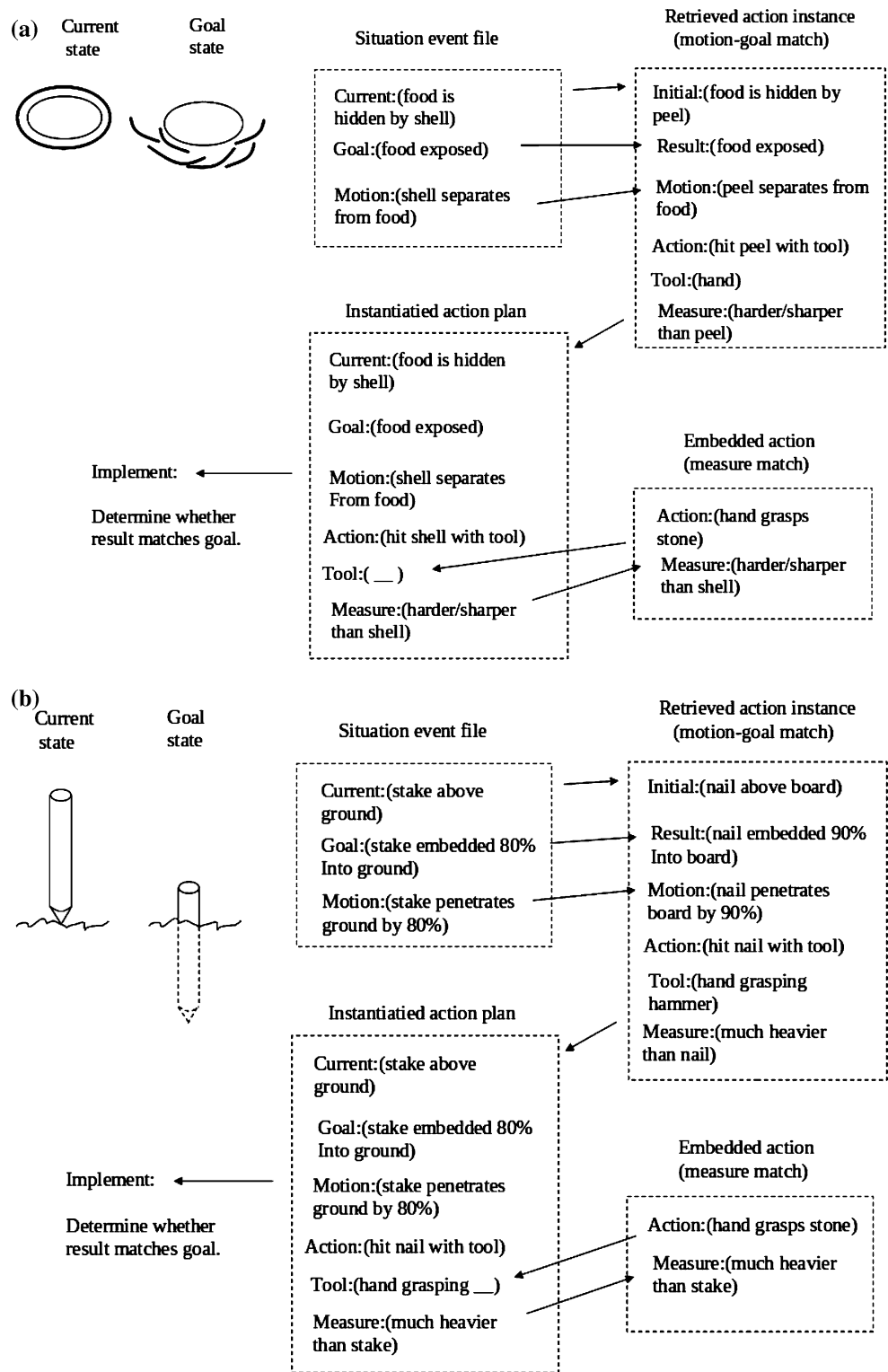
526 In the model shown in Fig. 1, the task environment 577  
 527 explicitly specifies the current layout of task-relevant 578  
 528 objects and implicitly specifies a goal layout in which the 579  
 529 position or orientation of one or more objects has changed. 580  
 530 This task environment is represented by an event file 581  
 531 (Hommel, 2004) binding the current layout, the goal layout 582  
 532 and the motion(s) required to resolve the spatial discrep- 583  
 533 ancy between the two layouts. Such event files are con- 584  
 534 structed hierarchically from lower-level object-motion 585  
 535 bindings, in a process that is sensitive to priming by long- 586  
 536 term memory (LTM) resident representations encoding 587  
 537 relationships between objects or features in the current per- 588  
 538 ceived situation (Colzato, Raffone, & Hommel, 2006). 589  
 539 Event files, thus, provide a level of representation at which 590  
 540 relational priming could drive implicit analogical inference, 591  
 541 as proposed by Leech, Mareshal and Cooper (2008) for 592  
 593  
 594

analogies between concepts expressible in language. Con- 542  
 struction of a task-environment event file requires representa- 543  
 tion of the goal layout as a manipulable image, and inference 544  
 of the required motion(s) from the spatial discrepancy 545  
 between the perceived current layout and an imagined goal 546  
 layout. It is important to emphasize that neither the goal lay- 547  
 out nor the inferences of motion need be consciously experi- 548  
 enced. Even in humans, such representations and inferences 549  
 are not experienced during expert "flow-like" performance of 550  
 familiar tasks (Dietrich, 2004; Ericsson & Lehmann, 1996). 551

552 The first phase of structure mapping is initiated by acti- 553  
 554 vation of an LTM resident representation of a previously 554  
 555 executed or observed action instance or minimally 555  
 556 abstracted action schema that encodes both a result and a 556  
 557 motion sufficiently similar to the goal and motion encoded 557  
 558 by the task-environment representation. This retrieved 558  
 559 action instance is thus both a goal-result and a kinematic 559  
 560 match to the event file representing the task environment. 560  
 561 On the basis of this goal-motion alignment, the retrieved 561  
 562 instance is bound to the event file representing the task 562  
 563 environment. This binding step replaces the object and 563  
 564 motion representations of the retrieved action instance with 564  
 565 those of the task-environment event file to produce a par- 565  
 566 tially mapped, partially instantiated action plan that shares 566  
 567 the goal of and satisfies the kinematic requirements of the 567  
 568 task environment, but still encodes the dynamic, i.e., force- 568  
 569 application, parameters of the retrieved action instance. 569  
 570 Such non-intentional—in fact fully unconscious—replacement 570  
 571 of components of retrieved representations by components of 571  
 572 current perceptual representations by structure-mapping 572  
 573 mechanisms has been observed in verbal analogies (Day & 573  
 574 Gentner, 2007).

574 The appropriate application of force is critical to suc- 574  
 575 cessful tool improvisation; as discussed above, it is only in 575  
 576 the context of such dynamic constraints that a tool can be 576  
 577 said to be analogous to a part of the body. The proposed 577  
 578 model requires that LTM-resident action instances encode 578  
 579 applied force in two ways: as a reproducible sensation of 579  
 580 muscular effort and as a parametric representation of the 580  
 581 resulting motion. Choice of and use of tools by chimpan- 581  
 582 zees indicate that they are sensitive to these representations 582  
 583 of force (Brill et al., 2009). Calibration of these two repre- 583  
 584 sentations to achieve expert ability in fine motor control 584  
 585 requires extensive practice (Ericsson & Lehmann, 1996). 585  
 586 Studies of expert athletes indicate that fine adjustments in 586  
 587 motor control driven by representations of muscular force 587  
 588 are performed unconsciously in response to unconscious 588  
 589 perceptions of movement requirements (Kibele, 2006), 589  
 590 consistent both with their common encoding at the event- 590  
 591 file level and the independence of force-motion inferences 591  
 592 from deliberate conceptual processing. In the model shown 592  
 593 in Fig. 1, applied force is represented parametrically with 593  
 594 respect to the object to which force is applied, while motion

**Fig. 1** Frame-based representation of structure-mapping steps in tool-improvisation analogies: **a** the stone:nut::hand:fruit analogy and **b** the stone:tent-stake::hammer:nail analogy. Arrows indicate retrieval and mapping steps. The use of frames is heuristic only and is not meant to imply that the representations implemented by the praxis system encode concepts expressible in either public language or an internally comprehended language of thought



595 is represented qualitatively in terms of the final dispositions  
 596 of relevant objects.  
 597 A partially mapped action plan may be executed, but  
 598 will fail in cases requiring tool improvisation. In the  
 599 stone:nut::hand:fruit case, implementation of the partially  
 600 mapped plan fails because hands are not hard and sharp

enough to open nuts, as some, but not all wild chimpanzees  
 eventually comprehend (Biro et al., 2003). In the  
 stone:tent-stake::hammer:nail case, the partially mapped  
 plan typically fails because the backpacker has not brought  
 a hammer. In either case, failure of the partially mapped  
 action plan initiates the second phase of structure mapping.

607 In this phase, one or more LTM-resident action instances  
 608 are activated that encode force measures similar to that of  
 609 the partially mapped action plan. Similarities in applied  
 610 force are relational, not surface, similarities. The retrieved  
 611 action plan is embedded into the partially mapped action  
 612 plan, inducing replacement of the insufficient tool with the  
 613 object of the embedded action. The result of this embed-  
 614 ding is a fully mapped action plan incorporating the objects  
 615 and motions of the task environment and the alternative  
 616 tool retrieved for its ability to meet the force requirements  
 617 of the task environment. This model of action embedding  
 618 as a method of action-plan generation is similar to that  
 619 employed in some robotic action planners (e.g., Beaudry  
 620 et al., 2005). Action plan embedding involves holding at  
 621 least two action plans in WM simultaneously, a form of  
 622 multitasking; hence capability in action-plan embedding  
 623 and therefore in tool improvisation would be expected to  
 624 increase with increased development of rostral prefrontal  
 625 cortex, which supports multitasking (Dreher et al., 2008;  
 626 Green et al., 2006), consistent with the observed capability  
 627 gradients from simians to great apes to humans and from  
 628 children to adults. Interestingly, young chimpanzees are  
 629 more efficient learners of some tool-use tasks than are  
 630 human children (Horner & Whiten, 2005), suggesting that  
 631 they may be more efficient tool improvisers as well.

632 Instantiation of the fully mapped action plan provides  
 633 the criteria necessary for a visual and tactile search for an  
 634 object to serve as the alternative tool followed by dynamic  
 635 testing of the object to determine whether it actually meets  
 636 the force-application requirements of the action plan. Heft-  
 637 ing a stone to assess its weight or bending a stick to assess  
 638 its rigidity are dynamic tests of this kind. Tool modification  
 639 may follow testing. The fully mapped action plan is then  
 640 executed and its results observed. Successful plans are  
 641 those for which the result of execution matches the goal.

642 The binding and memory-access steps proposed by this  
 643 model of structure mapping would be expected to engage  
 644 areas of the temporal-parietal junction (binding), pre-motor  
 645 cortex (mirror-neuron action representation and motor  
 646 planning), anterior cingulate cortex (process monitoring  
 647 and conflict detection), dorso-lateral prefrontal cortex (goal  
 648 maintenance and WM management) and rostral prefrontal  
 649 cortex (attentional control). Left-hemisphere activation  
 650 would be expected to dominate, consistent with the left-  
 651 hemisphere specialization for sequential actions (Fiebach &  
 652 Schubotz, 2006) and tool-related actions in particular  
 653 (Lewis, 2006). Such activation would contrast with the  
 654 right-hemisphere activation associated with general seman-  
 655 tic representations (Bar, 2008), which is observed specifi-  
 656 cally when subjects solve word-association problems  
 657 involving distant semantic connections (Bowden et al.,  
 658 2006; Sandkühler & Bhattacharya, 2008; Jung-Beeman  
 659 et al., 2004; Kounios & Beeman, 2009). Specific tests of

660 this model would require either imaging or magnetic deac-  
 661 tivation of specific praxis-network areas while subjects per-  
 662 formed tool-use relevant analogies not presented in, and  
 663 hence not potentially confounded by, language. Activity  
 664 patterns generated while subjects performed analogies pre-  
 665 sented in language involving tool use, tools but no tool-use  
 666 motions, bodily motions but no tools and neither motions  
 667 nor tools would be suitable comparisons. Testing these lat-  
 668 ter conditions separately would provide a more sensitive  
 669 analysis than that of Green et al. (2006), who employed  
 670 some analogy problems involving descriptions of physical  
 671 motions.

#### 672 **Consequences of the event-file manipulation model:** 673 **functional dependence of motion concepts on structure** 674 **mapping**

675 The functional model outlined above and illustrated in  
 676 Fig. 1 describes tool improvisation as structure mapping at  
 677 three levels. First, the objects in the task environment are  
 678 mapped to objects in the retrieved action instance using  
 679 motion and goals or results as structuring relations. Second,  
 680 tools in the retrieved action instance are mapped to tools in  
 681 the embedded action by using a force measure as the struc-  
 682 turing relation. Finally, the observed result of executing a  
 683 successful fully mapped action plan is related, in practice,  
 684 to the result of the original retrieved action instance by the  
 685 functional composition of the two previous structure map-  
 686 pings. Thus, the criterion of systematicity that characterizes  
 687 good analogies (Gentner, 2005; Holyoak, 2005) can be rig-  
 688 orously defined in the case of tool improvisation as coher-  
 689 ent scaling of both the kinematic and dynamic requirements  
 690 between source and target cases. An “analogy” in which the  
 691 forces applied cannot produce the motion required to  
 692 achieve the goal is not a good analogy; applying too much  
 693 force—swatting a fly with an axe—generally produces bad  
 694 results as well.

695 If this model of tool improvisation is correct, two promi-  
 696 nent claims regarding analogical inference require revision.  
 697 First, the claim that structure-mapping analogy is a  
 698 uniquely human capability, which has been based on the  
 699 poor performance of animals on abstract and conceptual  
 700 analogy tasks (Gentner, 2003; Penn et al., 2008) must be  
 701 rejected in the case of tool-improvisation analogies, which  
 702 members of many non-human species perform with facility  
 703 in the wild. Second, the claim that structure-mapping anal-  
 704 ogy is dependent on relational language (Gentner, 2003) or  
 705 on explicit access to relational concepts in a language of  
 706 thought (Penn et al., 2008) must also be rejected in the case  
 707 of tool-improvisation analogies, both for animals lacking  
 708 such language, and for humans who may implement such  
 709 analogies using language-independent, event-file-based




710 binding and action-planning mechanisms. Indeed, the  
711 model predicts exactly the reverse functional dependency:  
712 that a natural class of motion and force concepts express-  
713 ible in language are functionally dependent on the struc-  
714 ture-mapping capabilities of the event-file manipulation  
715 and action-planning systems.

716 The neurocognitive representation of abstract concepts,  
717 such as “tool”, “motion” or “force” is not well understood  
718 (Martin, 2007). However, humans can clearly focus suffi-  
719 cient attention on the representations of such concepts, in  
720 the absence of relevant perceptual input, to activate overt  
721 behaviors including speech. Alert attentional focus on inter-  
722 nal representations in the absence of perceptual input is  
723 managed by an area of medial rostral prefrontal cortex  
724 proximal to areas implementing the self-other distinction  
725 and hence the capacity for autoegetic episodic memory  
726 (Simons, Henson, Gilbert, & Fletcher, 2008; Turner,  
727 Simons, Gilbert, Frith, & Burgess, 2008); the apparent  
728 human-specificity of both experienced abstract conceptual  
729 understanding (Penn et al., 2008) and experienced autoe-  
730 getic memory (Suddendorf & Corballis, 2007) may result  
731 from the evolutionarily recent elaboration of this region of  
732 cortex (Burgess et al., 2007). Not all possible abstractions  
733 of motions and forces, however, are expressed by abstract  
734 concepts in natural languages: most such abstractions are  
735 expressible only in the artificial, technical languages of ana-  
736 lytical mathematics and physics. The unconscious execu-  
737 tion of structure-mapping inferences by the binding and  
738 premotor systems provides a mechanism by which some  
739 particular motion and force abstractions, those activated in  
740 tool use and in recognizing the utility of objects as tools,  
741 would be sufficiently selectively reinforced by everyday  
742 life to make them available for attentional amplification  
743 even in the absence of relevant perceptual input. An expect-  
744 ation of the model outlined here is, therefore, that the force  
745 and motion concepts expressible in natural languages, and  
746 hence those employed in “folk physics,” will be those that  
747 would be activated in unconscious structure mappings  
748 involving tool use.

749 It is well known that children naturally develop  
750 (Karmaloff-Smith, 1995) and adults routinely employ  
751 (Gentner, 2002) a “folk physics” with essentially Aristote-  
752 lian concepts of force and motion. These concepts include  
753 the notion that motion continues only as long as force is  
754 applied and the notion that the shapes of curvilinear trajec-  
755 tories are preserved by “curvilinear momentum”. These  
756 concepts conflict with classical Newtonian mechanics, but  
757 are easily understood from the perspective of tool manipu-  
758 lation. Using tools requires applying force, force that is felt  
759 as feedback from the muscles. Hence, tool use would tend  
760 to reinforce the Aristotelian and folk-physics notion that  
761 continuing motion requires continuing application of force.  
762 Hand-held tools that move in curvilinear trajectories do so

763 because they are swung by arms moving forcefully on fixed 763  
764 pivots, the shoulders. Hence, forceful curvilinear motions 764  
765 with tools would tend to reinforce the notion of curvilinear 765  
766 momentum, and as well as the intuitive notion of centrifugal 766  
767 force. Such felt muscular forces and typical resulting 767  
768 trajectories and force-application capabilities are the 768  
769 relations that drive tool-improvisation structure mappings 769  
770 of the kind illustrated in Fig. 1. The folk physics concepts 770  
771 of continuing force for continuing motion, curvilinear 771  
772 momentum and centrifugal force are, therefore, the very 772  
773 concepts that would be expected if the human paradigms of 773  
774 physical motions are tool-use motions and the paradigms of 774  
775 forces are the muscular forces employed to assess whether 775  
776 an object is suitable as a tool, and then to use it as such. 776  
777 These folk physics concepts of motion and force are rou- 777  
778 tinely employed to solve practical problems in contexts in 778  
779 which subjects cannot later fully enunciate either a comple- 779  
780 te and correct description of the task environment or of 780  
781 the rules being employed, suggesting that problem solving 781  
782 is being performed by visuo-motor simulation, not explicit 782  
783 conceptual reasoning (Hegarty, 2004; Wolff, 2007), consis- 783  
784 tent with a functional dependence of the concepts as con- 784  
785 sciously understood and expressed in language on 785  
786 underlying pre-motor capabilities. Children identify situa- 786  
787 tions in which hidden mechanisms cause unexpected 787  
788 behavior unattributable to animate agency at around 4 years 788  
789 of age (Sobel, Yoachim, Gopnik, Meltzoff, & Blumenthal, 789  
790 2007), well before they possess a conceptual understanding 790  
791 of mechanical systems, suggesting that they are capable of 791  
792 an implicit analysis of motions and implied forces. Activa- 792  
793 tion of the praxis system in qualitative numerosity judg- 793  
794 ments (Cantlon, Brannon, Carter, & Pelphrey, 2006) and in 794  
795 algebraic equation-solving (Qin et al., 2004) provides addi- 795  
796 tional suggestive evidence for the involvement of motor 796  
797 simulation in what on the surface appears to be purely con- 797  
798 ceptual problem solving. 798

799 Even in formalized, mathematical physics, analogies and 799  
800 metaphorical representations that directly conflict with 800  
801 established theory and hence with conceptual understand- 801  
802 ing are routinely relied upon and employed both practically 802  
803 and pedagogically. Perhaps, the best-known example is the 803  
804 Rutherford atom analogy electrons:nucleus::planets:sun, 804  
805 which was employed by Green et al. (2006) as a canonical 805  
806 test case for analogical reasoning. Ernest Rutherford’s 806  
807 (1911) model of the atom as consisting of a small, heavy 807  
808 central nucleus orbited by much lighter electrons was pro- 808  
809 posed to account for the results of experiments in which 809  
810 gold atoms were bombarded by high-energy alpha particles. 810  
811 Most of the alpha particles passed straight through the gold 811  
812 foil target, but others were deflected backwards, suggesting 812  
813 collisions with a small heavy object and thoroughly con- 813  
814 tradicting the then-dominant Thompson or “plum pudding” 814  
815 model of atoms as spheres containing a uniform mixture of 815

	Large 426	290	xxxx	Dispatch: 29.5.10	No. of Pages: 14	
	Journal	Article	MS Code	LE <input type="checkbox"/>	TYPESSET <input type="checkbox"/>	CP <input checked="" type="checkbox"/>

816 positively charged material and electrons (Rutherford, 1911  
817 and Randall, 2005 briefly review the relevant history from a  
818 physicist's perspective; Mehra and Rechenberg (1982) pro-  
819 vide a more detailed historical review). Rutherford's model  
820 was revolutionary in that it proposed an atom consisting  
821 mostly of empty space, in which the positive charges were  
822 concentrated in the center and the negative charges (the  
823 electrons) occupied the distant periphery. However, while  
824 the Thompson model with its statically embedded electrons  
825 was consistent with classical electrodynamics, the Ruther-  
826 ford orbital model directly contradicted existing theory:  
827 classical electrons moving in the electric field of the posi-  
828 tively charged nucleus would radiate away their kinetic  
829 energy in much less than a second, and the Rutherford atom  
830 would explosively collapse. This tension was resolved by  
831 Bohr's (1913) proposal of quantized electron orbits, but at  
832 the price of altogether removing the classical concept of  
833 motion from the physical description of events at atomic  
834 scales.

835 The staying power of the Rutherford atom with orbiting  
836 electrons, an image so ubiquitous as to be iconic, is prima  
837 facie evidence that experienced motions and forces are cen-  
838 tral to the understanding of even such abstract concepts as  
839 atoms. The popularity of Feynman diagrams as illustrations  
840 of elementary particle interactions provides further such  
841 evidence. Physicists greatly prefer Feynman diagrams to  
842 the complex path integrals that they represent, employing  
843 them in professional publications and pedagogy; Randall  
844 (2005) is a case in point. Such diagrams are, however,  
845 grossly misleading if taken literally. They depict particles  
846 as having well-defined trajectories, and depict the "virtual"  
847 particles that carry forces in quantum field theory as being  
848 emitted and absorbed at well-defined locations along these  
849 trajectories. Both of these depictions are flatly inconsistent  
850 with quantum mechanics. As in the case of the Rutherford  
851 atom, depictions consistent with the motions and forces of  
852 everyday tool use and folk physics are maintained as cogni-  
853 tive aids, even when they are inconsistent with conceptual  
854 knowledge. Pedagogical research in physics indicates that  
855 such graphic aids and the manipulations that they invoke  
856 nonetheless significantly aid conceptual learning (Lasry &  
857 Aulls, 2007). The utility of manipulations in conceptual  
858 learning is corroborated by recent experiments in which  
859 activation of components of the praxis network is directly  
860 measured. Subjects briefly trained to manipulate novel  
861 objects as if they were tools later classify them as tools, as  
862 indicated by activation of tool-specific areas of left-hemi-  
863 sphere TPJ and pre-motor cortex (Martin, 2007; Weisberg,  
864 van Turenhout, & Martin, 2007). Manipulating tools and  
865 other common objects facilitates verbal descriptions of  
866 their shapes in the absence of visual input, again accompa-  
867 nied by activation of tool-use relevant areas of TPJ (Oliver,  
868 Geiger, Lewandowski, & Thompson-Schill, 2009). In both

of these cases, as apparently in the cases of atoms and ele- 869  
mentary-particle interactions, learning and use of object 870  
concepts is facilitated by the kinds of manipulations that 871  
provide input to pre-motor structure-mapping inferences. 872

873 Additionally, albeit highly indirect evidence for the  
874 dependence of motion and force concepts on a small num-  
875 ber of abstractions of experienced motions and forces is  
876 provided by the relative paucity of words for motions and  
877 mechanical forces in the vocabularies of natural languages.  
878 Natural languages typically include words naming high-  
879 level abstractions: "move" for physical motion, "push" and  
880 "pull" for mechanical force, "put" and "take" for manipula-  
881 tions involving force-transferring actions. However, precise  
882 specifications of motions, even of the human body, tend to  
883 be specialized technical names or descriptive phrases.  
884 Reproducibly and correctly identifying the referents of such  
885 specialized terms typically requires extensive specialized  
886 training and practice; they are not "natural" parts of human  
887 languages. Two of the oldest such specialized vocabularies  
888 available for study are those of yoga and chi-gung. Both  
889 vocabularies employ richly descriptive metaphorical lan-  
890 guage to name precisely specified motions and postures,  
891 i.e., particular proprioceptive images. Both require exten-  
892 sive physical training and practice to correctly identify the  
893 referents of the these terms; specialized phrases such as  
894 "chaturanga" (a motion) or "downward dog" (a posture)  
895 name concepts that are learned by learning to recognize  
896 particular dynamic or static proprioceptive images. Posture  
897 names are far more common than motion names in the  
898 vocabularies of yoga and chi-gung, as they are in natural  
899 languages. Why are there not ubiquitous, natural concepts  
900 and hence names for many if not most of the motions avail-  
901 able to the human body, including those practiced in  
902 ancient disciplines, such as yoga and chi-gung? Perhaps,  
903 because these motions and the forces felt while performing  
904 them do not play common roles in pre-motor structure map-  
905 pings, and hence are not sufficiently reinforced to be avail-  
906 able to the internally directed attention required for  
907 conceptualization.


#### 908 **Testing the proposed model of tool-improvisation** 908 909 **structure-mapping inferences** 909

910 The event-file manipulation model of structure-mapping 910  
911 inference in tool improvisation proposed here generates a 911  
912 number of experimentally testable predictions in addition to 912  
913 the predicted praxis-network activations discussed above. 913  
914 Additional evidence relevant to any of these would serve to 914  
915 confirm or disconfirm the model as presented. 915


916 A primary prediction of the model is dissociability of 916  
917 conceptual comprehension of tool-improvisation analogies 917  
918 from their implementation. Patients exhibiting motor-imag- 918

919	ery apraxias that spare semantic memory would be	forces as organizing relations, whether or not they involve	972
920	expected to be capable of comprehending verbal explanations	tool improvisation.	973
921	of tool-improvisation analogies, but not of executing		
922	such analogies if they are presented in modalities other than		
923	language. Functional dissociation of tool-use abilities from		
924	conceptual knowledge of tools and their uses in human		
925	apraxias (reviewed by Johnson-Frey, 2004; Petreska, Adri-		
926	ani, Blanke, & Billard, 2007) provides support for this pre-		
927	dition. Conversely, patients exhibiting aphasias disrupting		
928	semantic memory for tools and tool uses, but not apraxia,		
929	would be expected to be incapable of understanding verbal		
930	descriptions of tool-improvisation analogies, but capable of		
931	executing them if presented graphically or with actual candi-		
932	date tools. The practical intelligence displayed by Susan		
933	Schaller's language-less subject Ildefonso, who appears to		
934	have lacked conscious conceptual knowledge (Schaller,		
935	1995), is consistent with this prediction.		
936	The event-file manipulation model of structure mapping		
937	also predicts that cognitively normal subjects would com-		
938	plete tool-improvisation analogy tasks more rapidly and if		
939	time-limited, more accurately if the analogy problems were		
940	presented graphically, visually or tactily as compared to		
941	verbally. It predicts that chimpanzees and possibly orangu-		
942	tans may exhibit higher-than-expected analogical ability if		
943	presented with tasks requiring the analogical transfer of		
944	causal knowledge from one context to another, as compared		
945	with the symbolic analogy tasks reviewed by Gentner		
946	(2003). The performance of young chimpanzees, which		
947	used the same tools and methods to extract a food reward		
948	from an opaque box as they had used to extract a similar		
949	reward from a similar transparent box (Horner & Whiten,		
950	2005) is consistent with this prediction.		
951	The mechanism of action embedding postulated by the		
952	model predicts that RPFC activation in tool-improvisation		
953	tasks will scale with the number of independent motions,		
954	and hence the number of independent embedded actions,		
955	required to complete the task. It similarly predicts enhanced		
956	RPFC activation in analogy tasks in which multiple embed-		
957	dable actions conflict, compared to tasks in which action-		
958	embedding conflict is minimal.		
959	The model would also predict that individuals scoring in		
960	the low range on tests of systemizing bias (Baron-Cohen,		
961	2002; Baron-Cohen et al., 2003) will exhibit worse perfor-		
962	mance on tool-improvisation analogies than individuals		
963	with matched total or verbal IQ, but with higher systemiz-		
964	ing bias. This prediction is consistent with general observa-		
965	tions of correlations between sex, gender orientation,		
966	systemizing bias and mechanical skills (Goldenfeld, Baron-		
967	Cohen, & Wheelwright, 2006; Nettle, 2007; Baron-Cohen,		
968	2008).		
969	Finally, the considerations outlined in the previous sec-		
970	tion suggest that the above predictions may extend to other		
971	or even all analogies involving motion and mechanical		
		<b>Conclusions</b>	974
		Structure-mapping analogy is a fundamental inferential and	975
		learning mechanism. It has been regarded as concept-	976
		dependent and human-specific (Gentner, 2003; Penn et al.,	977
		2008). The model developed here is based on the hypothe-	978
		sis that structure-mapping analogies in tool improvisation	979
		are implemented by manipulations of event files (Hommel,	980
		2004) and do not require awareness or understanding of	981
		relational concepts expressible in language. Considerable	982
		observational and experimental evidences support this	983
		event-file manipulation model, suggesting that tool-impro-	984
		visation analogies are neither concept-dependent nor	985
		human-specific. This result renders human analogical capa-	986
		bilities continuous with those of other species, and provides	987
		an evolutionary path from higher-primate tool-improvisa-	988
		tion capability through proto-human tool-improvisation	989
		capability to modern-human tool-improvisation and possi-	990
		bly more general motion-and-force-involving analogy	991
		capabilities. It moreover suggests that at least some con-	992
		cepts common to natural languages, those referring to expe-	993
		rienced motions and forces, are functionally dependent on	994
		structure-mapping capabilities of the event-file binding and	995
		pre-motor planning systems. If correct, this functional	996
		dependence provides a mechanistic basis for proposals,	997
		such as that of Gallese and Lakoff (2005): visuo-motor sim-	998
		ulation underlies language abilities, and raises the possibil-	999
		ity that the human ability to focus attention on internally	1000
		generated representations (Burgess et al., 2007), not a	1001
		human-specific inferential capacity, is primarily responsi-	1002
		ble for the impressive analogical abilities of <i>Homo sapiens</i> .	1003
		<b>Acknowledgments</b> Two anonymous referees provided valuable	1004
		comments on an earlier version of this paper.	1005
		<b>Conflict of interest statement</b> The author states that he has no	1006
		conflicts of interest relevant to the reported research.	1007
		<b>References</b>	1008
		Bar, M. (2008). The proactive brain: Using analogies and associations to	1009
		derive predictions. <i>Trends in Cognitive Sciences</i> , 11(7), 280–289.	1010
		Baron-Cohen, S. (2002). The extreme male brain theory of autism.	1011
		<i>Trends in Cognitive Sciences</i> , 6(6), 248–254.	1012
		Baron-Cohen, S. (2008). Autism, hypersystemizing, and truth. <i>The</i>	1013
		<i>Quarterly Journal of Experimental Psychology</i> , 61(1), 64–75.	1014
		Baron-Cohen, S., Richler, J., Bisarya, D., Gurunathan, N., & Wheel-	1015
		wright, S. (2003). The systemizing quotient: An investigation of	1016
		adults with Asperger syndrome or high-functioning autism, and	1017
		normal sex differences. <i>Philosophical Transactions of the Royal</i>	1018
		<i>Society of London B</i> , 358, 361–374.	1019

- 1020 Barsalou, L. (2008). Grounded cognition. *Annual Review of Psychol-*  
1021 *ogy*, 59, 617–645.
- 1022 Beaudry, E., Brosseau, Y., Cote, C., Raievisky, C., Letourneau, D.,  
1023 Kabanza, F., & Michaud, F. (2005). Reactive planning in a moti-  
1024 vated behavioral architecture. In *Proceedings of the National*  
1025 *Conference on Artificial Intelligence AAAI-05*. [http://www-edu.](http://www-edu.gel.usherbrooke.ca/cotc2204/publications/AAAI2005Beaudry.pdf)  
1026 [gel.usherbrooke.ca/cotc2204/publications/AAAI2005Beaudry.](http://www-edu.gel.usherbrooke.ca/cotc2204/publications/AAAI2005Beaudry.pdf)  
1027 [pdf](http://www-edu.gel.usherbrooke.ca/cotc2204/publications/AAAI2005Beaudry.pdf). Accessed 1 Feb 2009.
- 1028 Biro, D., Inoue-Nakamura, N., Tonooka, R., Yamakoshi, G., Sousa, C.,  
1029 & Matsuzawa, T. (2003). Cultural innovation and transmission of  
1030 tool use in wild chimpanzees: Evidence from field experiments.  
1031 *Animal Cognition*, 6, 213–223.
- 1032 Boettiger, C., & D'Esposito, M. (2005). Frontal networks for learning  
1033 and executing arbitrary stimulus-response associations. *Journal*  
1034 *of Neuroscience*, 25(10), 2723–2732.
- 1035 Bohr, N. (1913). On the constitution of atoms and molecules, Part II—  
1036 Systems containing only a single nucleus. *Philosophical Maga-*  
1037 *zine*, 26, 476–502.
- 1038 Breuer, T., Ndongou-Hockemba, M., & Fishlock, V. (2005). First  
1039 observation of tool use in wild gorillas. *PLOS Biology*, 3(11),  
1040 e380.
- 1041 Brill, B., Dietrich, G., Foucart, J., Fuwa, K., & Hirata, S. (2009). Tool  
1042 use as a way to assess cognition: How do captive chimpanzees  
1043 handle the weight of the hammer when cracking a nut? *Animal*  
1044 *Cognition*, 12, 217–235.
- 1045 Burgess, P. W., Simons, J., Dumontheil, I., & Gilbert, S. (2007). The  
1046 gateway hypothesis of rostral prefrontal cortex (area 10) function.  
1047 In J. Duncan, L. Phillips & P. McLeod (Eds.), *Measuring the mind:*  
1048 *speed, control, and age* (pp. 217–248). Oxford University Press.
- 1049 Byrne, R. W., Bates, L. A., & Moss, C. J. (2009). Elephant cognition  
1050 in primate perspective. *Comparative Cognition and Behavior*  
1051 *Reviews*, 4, 1–15.
- 1052 Cantlon, J., Brannon, E., Carter, E., & Pelphrey, K. (2006). Functional  
1053 imaging of numerical processing in adults and 4-y-old children.  
1054 *PLOS Biology*, 4(5), 0844–0854.
- 1055 Carvalho, S., Cunha, E., Sousa, C., & Matsuzawa, T. (2008). Chaînes  
1056 opératoires and resource-exploitation strategies in chimpanzee  
1057 (*Pan troglodytes*) nut cracking. *Journal of Human Evolution*, 55,  
1058 148–163.
- 1059 Catmur, C., Gillmeister, H., Bird, G., Liepelt, R., Brass, M., & Heyes,  
1060 C. (2008). Through the looking lass: Counter-mirror activation  
1061 following incompatible sensorimotor learning. *European Journal*  
1062 *of Neuroscience*, 28, 1208–1215.
- 1063 Catmur, C., Walsh, V., & Heyes, C. (2007). Sensorimotor learning  
1064 configures the human mirror system. *Current Biology*, 17, 1527–  
1065 1531.
- 1066 Cole, M. W., & Schneider, W. (2007). The cognitive control network:  
1067 Integrated cortical regions with dissociable functions. *NeuroIm-*  
1068 *age*, 37, 343–360.
- 1069 Colzato, L., Raffone, A., & Hommel, B. (2006). What do we learn from  
1070 binding features? Evidence for multilevel feature integration.  
1071 *Journal of Experimental Psychology: Human Perception and*  
1072 *Performance*, 32, 705–716.
- 1073 Courtney, S. (2004). Attention and cognitive control as emergent prop-  
1074 erties of information representation in working memory. *Cogni-*  
1075 *tive, Affective and Behavioral Neuroscience*, 4(4), 501–516.
- 1076 Culham, J., & Valyear, K. (2006). Human parietal cortex in action.  
1077 *Current Opinion in Neurobiology*, 16, 205–212.
- 1078 Day, S., & Gentner, D. (2007). Nonintentional analogical inference in  
1079 text comprehension. *Memory and Cognition*, 35(1), 39–49.
- 1080 Dietrich, A. (2004). Neurocognitive mechanisms underlying the experi-  
1081 ence of flow. *Consciousness and Cognition*, 13, 746–761.
- 1082 Dreher, J.-C., Koechlin, E., Tierney, M., & Grafman, J. (2008). Dam-  
1083 age to the fronto-polar cortex is associated with impaired multi-  
1084 tasking. *PLOS One*, 3(9), e3227.
- Engel, A., Burke, M., Fiehler, K., Bien, S., & Rosler, F. (2007). How  
1085 moving objects become animated: The human mirror system  
1086 assimilates non-biological movement patterns. *Social Neurosci-*  
1087 *ence*, 3, 368–387.
- Ericsson, K., & Lehmann, A. (1996). Expert and exceptional perfor-  
1089 mance: Evidence of maximal adaptation to task constraints.  
1090 *Annual Review of Psychology*, 47, 273–305.
- Ferrari, P., Rozzi, S., & Fogassi, L. (2005). Mirror neurons responding  
1092 to observation of actions made with tools in monkey ventral  
1093 premotor cortex. *Journal of Cognitive Neuroscience*, 17(2), 212–  
1094 226.
- Fiebach, C., & Schubotz, R. (2006). Dynamic anticipatory processing  
1096 of hierarchical sequential events: A common role for Broca's area  
1097 and ventral pre-motor cortex across domains? *Cortex*, 42, 499–  
1098 502.
- Gallese, V., & Lakoff, G. (2005). The brain's concepts: The role of  
1100 sensory-motor systems in conceptual knowledge. *Cognitive Neu-*  
1101 *ropsychology*, 22, 455–479.
- Gentner, D. (2002). Psychology of mental models. In N. J. Smelser &  
1103 P. B. Bates (Eds.), *International encyclopedia of the social and*  
1104 *behavioral sciences* (pp. 9683–9687). Amsterdam: Elsevier.
- Gentner, D. (2003). Why we're so smart. In D. Gentner & S. Goldin-  
1106 Meadow (Eds.), *Language and mind: Advances in the study of*  
1107 *language and thought* (pp. 195–235). Cambridge, MA: MIT  
1108 Press.
- Gentner, D. (2005). The development of relational category knowl-  
1110 edge. In L. Gershkoff-Stowe & D. Rakison (Eds.), *Building object*  
1111 *categories in developmental time* (pp. 245–275). Hillsdale, NJ:  
1112 Erlbaum.
- Gentner, D., Brem, S., Ferguson, R., Markman, A., Levidow, B.,  
1114 Wolff, P., et al. (1997). Analogical reasoning and conceptual  
1115 change: A case study of Johannes Kepler. *Journal of the Learning*  
1116 *Sciences*, 6(1), 3–40.
- Gentner, D., Christie, S., et al. (2008). Relational language supports  
1118 relational cognition in humans, apes (Comment on Penn, 2008).  
1119 *Behavioral and Brain Sciences*, 31(2), 136–137.
- Gilbert, S., Frith, C., & Burgess, P. (2005). Involvement of rostral pre-  
1121 frontal cortex in selection between stimulus-oriented and stimu-  
1122 lus-independent thought. *European Journal of Neuroscience*, 21,  
1123 1423–1431.
- Goldenfeld, N., Baron-Cohen, S., & Wheelwright, S. (2006). Empa-  
1125 thizing and systemizing in males, females, and autism. *Clinical*  
1126 *Neuropsychiatry*, 2, 338–345.
- Green, A., Fugelsang, J., Kraemer, D., Shamosh, N., & Dunbar, K.  
1128 (2006). Frontopolar cortex mediates abstract integration in anal-  
1129 ogy. *Brain Research*, 1096, 125–137.
- Grefkes, C., & Fink, G. (2005). The functional organization of the  
1131 intraparietal sulcus in humans and monkeys. *Journal of Anatomy*,  
1132 207, 3–17.
- Hauser, M., Pearson, H., & Seelig, D. (2002). Ontogeny of tool use in  
1134 cottontop tamarins, *Saguinus oedipus*: Innate recognition of func-  
1135 tionally-relevant features. *Animal Behavior*, 64, 299–311.
- Hegarty, M. (2004). Mechanical reasoning by mental simulation.  
1137 *Trends in Cognitive Sciences*, 8(6), 280–285.
- Holyoak, K. (2005). Analogy. In K. Holyoak & R. Morrison (Eds.),  
1139 *The Cambridge handbook of thinking and reasoning* (pp. 117–  
1140 142). Cambridge: Cambridge University Press.
- Hommel, B. (2004). Event files: Feature binding in and across percep-  
1142 tion and action. *Trends in Cognitive Sciences*, 8(11), 494–500.
- Horner, V., & Whiten, A. (2005). Causal knowledge and imitation/  
1144 emulation switching in chimpanzees (*Pan troglodytes*) and chil-  
1145 dren (*Homo sapiens*). *Animal Cognition*, 8, 164–181.
- Hunt, G., & Grey, R. (2003). Diversification and cumulative evolution  
1147 of New Caledonian crow tool manufacture. *Philosophical Trans-*  
1148 *actions of the Royal Society of London B*, 270, 867–874.


	Large 426	290	xxxx	Dispatch: 29.5.10		No. of Pages: 14	
	Journal	Article	MS Code	LE <input type="checkbox"/>	TYPESSET <input type="checkbox"/>	CP <input checked="" type="checkbox"/>	DISK <input checked="" type="checkbox"/>

- 1150 Hunt, G., & Grey, R. (2004). The crafting of hook tools by wild  
1151 New Caledonian crows. *Philosophical Transactions of the Royal*  
1152 *Society of London B (Suppl)*, 271, S88–S90.
- 1153 Johnson-Frey, S. (2004). The neural bases of complex tool use in  
1154 humans. *Trends in Cognitive Sciences*, 8(2), 71–78.
- 1155 Johnson-Frey, S., Newman-Norland, R., & Grafton, S. (2005). A  
1156 distributed left-hemisphere network active during planning of  
1157 everyday tool use skills. *Cerebral Cortex*, 15, 681–695.
- 1158 Jung-Beeman, M., Bowden, E. M., Haberman, J., Frymiare, J. L.,  
1159 Arambel-Liu, S., Greenblatt, R., et al. (2004). Neural activation  
1160 when people solve verbal problems with insight. *PLOS Biology*,  
1161 2, 0500–0510.
- 1162 Karmaloff-Smith, A. (1995). *Beyond modularity: A developmental*  
1163 *perspective on cognitive science*. Cambridge, MA: MIT Press.
- 1164 Kibele, A. (2006). Non-consciously controlled decision making for  
1165 fast motor reactions in sports—A priming approach for motor  
1166 responses to non-consciously perceived movement features.  
1167 *Psychology of Sport and Exercise*, 7(6), 591–610.
- 1168 Kounios, J., & Beeman, M. (2009). The “Aha!” moment: The cogni-  
1169 tive neuroscience of insight. *Current Directions in Psychological*  
1170 *Science*, 18(4), 210–216.
- 1171 Króliczak, G., & Frey, S. H. (2009). A common network in the left  
1172 cerebral hemisphere represents planning of tool use pantomimes  
1173 and familiar intransitive gestures at the hand-independent level.  
1174 *Cerebral Cortex*, 19, 2396–2410.
- 1175 Krutzen, M., Mann, J., Heithaus, M., Connor, R., Bejder, L., &  
1176 Sherwin, W. (2005). Cultural transmission of tool use in bottle-  
1177 nose dolphins. *Proceedings of the National Academy of Sciences*  
1178 *USA*, 102(25), 8939–8943.
- 1179 Lasry, N., & Aulls, M. (2007). The effects of multiple internal repre-  
1180 sentations on context rich instruction. *American Journal of Phys-*  
1181 *ics*, 75, 1030–1037.
- 1182 Leech, R., Mareshal, D., & Cooper, R. (2008). Analogy as relational  
1183 priming: A developmental and computational perspective on the  
1184 origins of a complex cognitive skill. *Behavioral and Brain Sci-*  
1185 *ences*, 31, 357–378.
- 1186 Lewis, J. (2006). Cortical networks related to human use of tools. *The*  
1187 *Neuroscientist*, 12(3), 211–231.
- 1188 Mann, J., Sargeant, J. L., Watson-Capps, J. J., Gibson, Q. A., Heithaus,  
1189 M. R., Connor, R. C., et al. (2008). Why do dolphins carry spon-  
1190 ges? *PLOS One*, 3, 1–7.
- 1191 Maravita, A., & Iriki, A. (2004). Tools for the body (schema). *Trends*  
1192 *in Cognitive Sciences*, 8(2), 79–86.
- 1193 Marshall-Pescini, S., & Whiten, A. (2009). Social learning of nut-  
1194 cracking behavior in East African sanctuary-living chimpanzees  
1195 (*Pan troglodytes schweinfurthii*). *Journal of Comparative Psy-*  
1196 *chology*, 122, 186–194.
- 1197 Martin, A. (2007). The representation of object concepts in the brain.  
1198 *Annual Review of Psychology*, 58, 25–45.
- 1199 Mehra, J., & Rechenberg, H. (1982). *The historical development of*  
1200 *quantum theory, Vol 1: The quantum theory of Planck, Einstein,*  
1201 *Bohr and Sommerfeld: its foundation and the rise of its difficulties*  
1202 *1900–1925 (372 pp)*. Berlin: Springer.
- 1203 Morrison, R., Krawczyk, D., Holyoak, K., Hummel, J., Chow, T., Mill-  
1204 er, B., et al. (2005). A neurocomputational model of analogical  
1205 reasoning and its breakdown in frontotemporal lobar degenera-  
1206 tion. *Journal of Cognitive Neuroscience*, 16(2), 260–271.
- 1207 Nettle, D. (2007). Empathizing and systemizing: What are they, and  
1208 what do they contribute to our understanding of psychological sex  
1209 differences? *British Journal of Psychology*, 98, 237–255.
- 1210 Oliver, R. T., Geiger, E. J., Lewandowski, B. C., & Thompson-Schill,  
1211 S. L. (2009). Remembrance of things touched: How sensorimotor  
1212 experience affects the neural instantiation of object form. *Neuro-*  
1213 *psychologia*, 47, 239–247.
- 1214 Osvath, M. (2009). Spontaneous planning for future stone throwing by  
1215 a male chimpanzee. *Current Biology*, 19, R190–R191.
- Osvath, M., & Osvath, H. (2008). Chimpanzee (*Pan troglodytes*) and  
1216 orangutan (*Pongo abelii*) forethought: Self-control and pre-expe-  
1217 rience in the face of future tool use. *Animal Cognition*, 11, 661–  
1218 674.
- Otoni, E., Dogo de Resende, B., & Izar, P. (2005). Watching the best  
1220 nutcrackers: What capuchin monkeys (*Cebus apella*) know about  
1221 others' tool-using skills. *Animal Cognition*, 24, 215–219.
- Penn, D., Holyoak, K., & Povinelli, D. (2008). Darwin's mistake:  
1223 Explaining the discontinuity between human and nonhuman  
1224 minds. *Behavioral and Brain Sciences*, 31, 109–178.
- Penn, D., & Povinelli, D. (2007). Causal cognition in human and non-  
1226 human animals: A comparative, critical review. *Annual Review of*  
1227 *Psychology*, 58, 97–118.
- Petreska, B., Adriani, M., Blanke, O., & Billard, A. (2007). Apraxia: A  
1229 review. In C. von Hofsten & K. Rosander (Eds.), *Progress in*  
1230 *brain research* (Vol. 164, pp. 61–83).
- Plummer, T. (2004). Flaked stones and old bones: Biological and cul-  
1232 tural evolution at the dawn of technology. *Yearbook of Physical*  
1233 *Anthropology*, 47, 118–164.
- Povinelli, D. J., Reaux, J. E., & Frey, S. H. (2009). Chimpanzee's con-  
1235 text-dependent tool use provides evidence for separable represen-  
1236 tations of hand and tool even during active use within  
1237 peripersonal space. *Neuropsychologia*, 48, 243–247.
- Pruetz, J. D., & Bertolani, P. (2007). Savanna chimpanzees, *Pan trog-*  
1239 *lodytes versus*, hunt with tools. *Current Biology*, 17, 1–6.
- Puce, A., & Perrett, D. (2003). Electrophysiology and brain imaging of  
1241 biological motion. *Philosophical Transactions of the Royal Soci-*  
1242 *ety of London B*, 358, 435–445.
- Qin, Y., Carter, C., Silk, E., Stenger, V. A., Fissell, K., Goode, A., et al.  
1244 (2004). The change in brain activation patterns as children learn  
1245 algebra equation solving. *Proceedings of the National Academy*  
1246 *of Sciences USA*, 101(15), 5686–5691.
- Randall, L. (2005). *Warped passages: Unraveling the mysteries of the*  
1248 *universe's hidden dimensions*. New York: Harper Perennial.
- Rizzolatti, G., & Craighero, L. (2004). The mirror-neuron system.  
1250 *Annual Reviews of Neuroscience*, 27, 169–192.
- Rutherford, E. (1911). The scattering of alpha and beta particles by  
1252 matter and the structure of the atom. *Philosophical Magazine*, 21,  
1253 669–688.
- Sandkühler, S., & Bhattacharya, J. (2008). Deconstructing insight: EEG  
1255 correlates of insightful problem solving. *PLOS One*, 3, e1459.
- Sanz, C. M., & Morgan, D. B. (2007). Chimpanzee tool technology in  
1257 the Goulougo Triangle, Republic of Congo. *Journal of Human*  
1258 *Evolution*, 52, 420–433.
- Schaller, S. (1995). *A man without words*. Berkeley: University of Cal-  
1260 ifornia.
- Schrauf, C., Huber, L., & Visalberghi, E. (2008). Do capuchin mon-  
1262 keys use weight to select hammer tools? *Animal Cognition*, 11,  
1263 413–422.
- Schubotz, R., & van Cramon, D. Y. (2004). Sequences of abstract non-  
1265 biological stimuli share ventral premotor cortex with action  
1266 observations and imagery. *The Journal of Neuroscience*, 24(24),  
1267 5467–5474.
- Sigman, M., & Dehaene, S. (2006). Dynamics of the central bottle-  
1269 neck: Dual-task and task uncertainty. *PLOS Biology*, 4(7), e220.
- Simons, J. S., Henson, R. N. A., Gilbert, S. J., & Fletcher, P. C. (2008).  
1271 Separable forms of reality monitoring supported by anterior pre-  
1272 frontal cortex. *Journal of Cognitive Neuroscience*, 20(3), 447–457.
- Sobel, D., Yoachim, C., Gopnik, A., Meltzoff, A., & Blumenthal, E.  
1274 (2007). The blicket within: Preschooler's inferences about insides  
1275 and causes. *Journal of Cognitive Development*, 8(2), 159–182.
- Stout, D., & Chaminade, T. (2007). The evolutionary neuroscience of  
1277 tool making. *Neuropsychologia*, 45(5), 1091–1100.
- Suddendorf, T., & Corballis, M. C. (2007). The evolution of foresight:  
1279 What is mental time travel, and is it unique to humans? *Behav-*  
1280 *ioral and Brain Sciences*, 30, 299–351.

	Large 426	290	xxxx	Dispatch: 29.5.10		No. of Pages: 14	
	Journal	Article	MS Code	LE <input type="checkbox"/>	TYPESSET <input type="checkbox"/>	CP <input checked="" type="checkbox"/>	DISK <input checked="" type="checkbox"/>

- 1282 Suddendorf, T., Corballis, M. C., & Collier-Baker, E. (2009). How  
1283 great is great ape foresight? *Animal Cognition*, *12*, 751–754.
- 1284 Tanji, J., & Hoshi, E. (2008). Role of lateral prefrontal cortex in behav-  
1285 ioral control. *Physiological Review*, *88*, 37–57.
- 1286 Tebbich, S., & Bshary, R. (2004). Cognitive abilities related to tool use  
1287 in the woodpecker finch, *Cactospiza pallida*. *Animal Behavior*,  
1288 *67*, 689–697.
- 1289 Turner, M. S., Simons, J. S., Gilbert, S. J., Frith, C. D., & Burgess, P.  
1290 W. (2008). Distinct roles for lateral and medial rostral prefrontal  
1291 cortex in source monitoring of perceived and imagined events.  
1292 *Neuropsychologia*, *46*(5), 1442–1453.
- 1293 Van Schaik, C., Ancrenaz, M., Borgen, G., Galdikas, B., Knott, C.,  
1294 Singleton, I., et al. (2003). Orangutan cultures and the evolution  
1295 of material culture. *Journal of Human Evolution*, *36*, 719–741.
- 1296 Visalberghi, E., Addessi, E., Truppa, V., Spagnoletti, N., Ottoni, E.,  
1297 Izar, P., et al. (2009). Selection of effective stone tools by wild  
1298 bearded capuchin monkeys. *Current Biology*, *19*, 1–5.
- 1299 Visalberghi, E., Fragaszy, D., Ottoni, E., Izar, P., de Oliveira, M. G., &  
1300 Andrade, F. R. D. (2007). Characteristics of hammer stones and  
anvils used by wild bearded capuchin monkeys (*Cebus libidino-* 1301  
*sus*) to crack open palm nuts. *American Journal of Physical* 1302  
*Anthropology*, *132*, 426–444. 1303
- Weir, A., & Kacelnik, A. (2006). A New Caledonian crow (*Corvus* 1304  
*moneduloides*) creatively re-designs tools by bending or unbend- 1305  
ing aluminum strips. *Animal Cognition*, *9*(4), 317–334. 1306
- Weisberg, J., van Turenout, M., & Martin, A. (2007). A neural system 1307  
for learning about object function. *Cerebral Cortex*, *17*, 513–521. 1308
- Whiten, A., Goodall, J., McGrew, W., Nishida, T., Reynolds, V., 1309  
Sugiyama, Y., et al. (2001). Charting cultural variation in 1310  
Chimpanzees. *Behavior*, *138*(11–12), 1481–1516. 1311
- Whiten, A., & van Schaik, C. (2007). The evolution of animal 1312  
'cultures' and social intelligence. *Philosophical Transactions of* 1313  
*the Royal Society of London B*, *362*, 603–620. 1314
- Wolff, P. (2007). Representing causation. *Journal of Experimental* 1315  
*Psychology: General*, *136*, 82–111. 1316
- Wynn, T. (2002). Archeology and cognitive evolution. *Behavioral and* 1317  
*Brain Sciences*, *25*, 389–438. 1318

UNCORRECTED PROOF

	Large 426	290	xxxx	Dispatch: 29.5.10	No. of Pages: 14	
	Journal	Article	MS Code	LE <input type="checkbox"/>	TYPESSET <input type="checkbox"/>	CP <input checked="" type="checkbox"/> DISK <input checked="" type="checkbox"/>