Title

Tradeoffs and cultural diversity

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Abstract

Culture is humanity's prime adaptation. Which cultural traits contribute to adaptive
value at the level of cultural groups, and how they do so, is, however, unclear and
debated. Here, we apply an approach from systems biology, known as Pareto task
inference (ParTI), to bring a fresh perspective to these questions. ParTI considers
systems that need to perform multiple tasks. No system can be optimal at all tasks at
once, leading to a fundamental tradeoff. These tradeoffs result in specific patterns in
multivariate trait data: data falls inside polygons in trait space, whose vertices are trait
combinations that are adaptive for each of the tasks. ParTI can infer the number of
adaptive tasks and their nature without need for *a-priori* assumptions, based on the
vertices of these polygons. Here, we applied ParTI to two datasets of cultural traits, on
Austronesian cultures and modern hunter-gatherers, adjusting for phylogeny and
spatial diffusion effects. We find evidence that these independent datasets show a
tradeoff between the same three tasks. We identify the tasks as resource defense,
resource competition, and mobility/exchange. Specific combinations of cultural traits
are adaptive for each of these tasks. This approach can be widely applied to any large
multivariate dataset to study trade-offs in cultural evolution.

<u>Introduction</u>

Culture has helped humans to adapt to diverse environments. Cultural traits vary

widely between groups, and a central goal of the field of cultural evolution is to understand

the processes that lead to this diversity. A central theory to understand the evolution of

cultural traits is the theory of cultural group selection (CGS), presented by Boyd and

Richerson (1). CGS proposes that processes of cultural evolution, acting on group-level

variation in cultural traits, lead to group-level adaptedness (2). Group-level variation in

cultural traits is transmitted, inherited, and maintained by various mechanisms, which include

biased social learning, coordination payoffs, punishment of deviant behaviors and group-

level symbolic boundaries (3). Intergroup competition, in the form of either natural selection,

selective imitation of successful groups, or selective migration between groups, acts on this

variation and promotes cultural traits beneficial on the group level (2,4–8). Cultural traits that

contribute to the fitness of the group tend to be selected (copied, maintained), and CGS

therefore predicts that cultural traits can be adaptive at the level of cultural groups.

Testing the hypothesis of CGS requires identifying which cultural traits are adaptive

at the level of the group, and what is their group-level function. This is difficult, since unlike

biological traits such as morphological features (3) or gene expression (4), the group-level

function of cultural traits cannot be readily measured or manipulated in a controlled setting.

Current methods mostly rely on correlations between traits and environmental features (4,9).

It is therefore important to develop new methods to rigorously test whether cultural traits are

adaptive, and, if so, what group-level function they serve.

We address this by developing an approach to infer the group-level function of

cultural traits directly from cross-cultural datasets of cultural traits. This method is applicable

to the case in which performance at several tasks contributes to group-level fitness. Thus,

following the notion of performance introduced by Arnold for organic evolution (10), fitness

as a function of traits $F(\vec{T})$ can be written in terms of performance functions $P_1, P_2, \ldots P_k$, one

for each task, such that $F(\vec{T}) = F(P_1(\vec{T}), P_2(\vec{T}), \ldots, P_k(\vec{T}))$. The idea is that each cultural

trait may contribute to (or reduce) performance in each task, and because no single

combination of traits can be optimal at all tasks at once, there is a fundamental trade-off.

A recent theory, called Pareto task inference (ParTI) (11,12), addressed this situation

and provides a way to infer the number and identity of the tasks (Figure 1). ParTI theory

shows that, under mild assumptions, the trait combinations that maximize F must fall inside a

low-dimensional polygonal shape in trait space (12). Each vertex of the polygon is a set of

traits   that is optimal for one of the tasks *i* (that is, the trait vector   that maximizes one of

the performance functions   ). Thus, the number of vertices is equal to the number of tasks:

For two tasks, the data should fall in a line segment between the two vertices; for three tasks,

it should fall in a triangle with the three vertices at the vertices and so on. Specialists at each

task lie near a vertex, and generalists lie in the center of the polygon. The intuitive reason for

the polygon is that any point outside of the polygon has a point inside the polygon that is

closer to all vertices, and hence has better performance at all tasks and hence higher fitness.

ParTI algorithms determine the best-fit polygon (line, triangle, tetrahedron, etc.) that

describes the data and its vertices. The algorithm uses statistical tests for the quality of the fit

to the polygon, by comparing the data to shuffled data (Methods). It thus discovers the

number of tasks, and, based on the cultures closest to each vertex, provides information that

can be used to infer the tasks.

ParTI was used to define evolutionary tasks and tradeoffs in several biological

domains, including gene expression in cells (13,14), development (15), animal morphology

(16), and life-history strategies (17) (Supplementary Information 1). Due to the generality of

the assumptions underlying ParTI, and their alignment with CGS theory, we reasoned that it

might also apply to cultural evolution. In this case, the input data is multivariate cross-

cultural trait datasets. Here we use ParTI to test hypothesis regarding adaptedness at the level

of groups, and to infer the number and nature of the tasks that groups face, and which cultural

traits are associated with these tasks.

Figure 1. Schema of ParTI algorithm to infer tasks and trade-offs from multivariate trait data. ParTI

receives as input a dataset where rows are cultures and columns are traits. ParTI tests the dimensionality of the

dataset, and whether it is well-described by a polygon according to statistical tests. If the data indeed falls on a

low-dimensional polygon, it infers its vertices. The number of vertices is equal to the number of tasks to be

inferred. The cultures closest to each vertex are assumed to specialize in a certain task, and information about

these cultures and the traits that make up the vertex is used to infer that task.

Methods

Cross-cultural datasets. The *Pulotu* (18) dataset on traits of 113 Austronesian cultures,

linked to a language-based family tree, was downloaded from https://pulotu.econ.mpg.de/.

We used 46 traits that are cultural (i.e. not physical) and belong to the traditional time focus

as defined by Watts et al. (18) (see Supplementary Information 2 for data preparation).

Binford's Hunter-Gatherer (BHG) (19) dataset, that includes cultural traits for 339 modern

hunter-gatherer cultures, was downloaded from http://ajohnson.sites.truman.edu/data-and-

program/ (see Supplementary Information 3 for data preparation). The two datasets differ

both in the mode of subsistence of societies that are included (*Pulotu* includes mostly

agricultural societies) and in the cultural traits measured, which are largely non-overlapping

between the datasets. In Supplementary Tables 1, 2 we list the traits used for our analysis

(codebooks for traits are available in the links above). Importantly, we verified that the

results are not confounded by the structure of the dataset. That is, the empty regions in the

trait-space of both datasets consist of (*a-priori*) feasible trait combinations (see

Supplementary Information 5 for discussion).

Language tree phylogenies. To generate a phylogenetic null model for principal component

analysis and for polygon statistical significance, we used a tree with informative branch

lengths for 93 Austronesian cultures from the *Pulotu* dataset (see Watts et al. (20)), `tree1`. For

phylogenetic adjustment of enrichment calculations, which does not require informative

branch lengths, we constructed a larger tree, `tree2`, by adding the other Austronesian cultures

according to their language families (`tree2`, provided in Supplementary Information 8).

Similarly, we constructed a phylogenetic tree for the cultures in *Binford's Hunter-Gatherer*

according to language families (`tree3`, provided in Supplementary Information 9).

Number of significant principal components. To test the effective dimensionality of

Pulotu, we determined the number of principal components that cannot be explained by a

phylogenetic null model in which cultural traits are acquired and lost at random based on

`tree1` (see Supplementary Information 4 for details).

Polytope inference and statistical significance. To infer the vertices of the polytopes

(polytopes are the generalization of polygons and polyhedra to higher dimensions) we

applied Principal Convex Hull Analysis (PCHA) (11,21) as implemented in the python

package *py_pcha*. Given an *n*-dimensional dataset and a number of vertices *k*, PCHA

computes a convex hull with *k* (*n*-dimensional) vertices that best accounts for the variance in

the data (11). This differs from PCA, which attempts to recover the orthogonal axes that

explain the most variance, which need not correspond to the relevant trade-off space. We

determined the number of vertices to test, k, by using the estimated dimension D of the data

estimated by PCA, by *k=D+1*. Thus, two-dimensional data is fit to a triangle with three

vertices, one-dimensional data to a line segment with two vertices, etc. We tested statistical

significance of polygons by using the t-ratio test as specified by Shoval et al. (12). The t-ratio

is the ratio between the areas (or volume) of the convex hull of the data and the area (or

volume) of the minimal enclosing polygon. The closer the polygon fits the data, the closer the

t-ratio is to one. The idea is that a good fit of a polygon means that the external outline of the

data cloud (its convex hull) has almost the same volume as the polygon (there is little empty

space between the polygon and the data). A p-value was computed by the probability that the

t-ratio of data shuffles is smaller or equal to the t-ratio of the real data based on 10,000 data

shuffles (see Supplementary Information 5). In cases where we adjusted for phylogeny and

spatial effects, the data was shuffled using null models hat preserve the phylogenetic and

geographic structure (Supplementary Information 5).

Trait regression according to distance from vertices. To infer tasks, we reasoned that traits

relevant to a task should appear most frequently in the cultures closest to the vertex

associated with that task. We tested this by performing regression, where the response

variable is the trait and the predictor variable is the Euclidean distance from vertex in the

space of the first two PCs. To adjust for spatial and phylogenetic effects, we added to each

regression, and for each trait, covariates for the average trait value among the 10 nearest

phylogenetic neighbors (tree2 and tree3) and 10 nearest spatial neighbors (see Supplementary

Information 6 for details), similar to the adjustment performed by Botero et al. (22). For

binary variables, we performed logistic regression, and for other variables, we performed

linear regression. We controlled for multiple hypotheses testing errors using FDR

(Supplementary Information 6).

<u>Results</u>

Figure 2. ParTI analysis of *Pulotu* dataset suggests three group-level tasks. (A) *Pulotu* cultural traits are

well described by a triangle (p<10⁻³) in the plain of the first two principal components. The three vertices of the

triangle correspond to adaptive trait complexes for the putative tasks of mobility/exchange, resource defense and

resource competition. (B) To test for statistical significance, we used the t-ratio test (12). The t-ratio is the ratio

between the areas of the convex hull (pink) and the area of the minimal enclosing triangle. A t-ratio close to 1

means the data fills the triangle. Shuffled datasets, on the other hand, do not fill their enclosing triangles as

tightly. (C) A consensus language tree for Austronesian languages. Cultures are color coded according to their

relative distance from each vertex of the triangle. (D) A geographical map, with color coding as in C. (E) To

infer tasks for each vertex, we seek traits that appear at higher frequency in the data points closest to the vertex.

Such strongly enriched traits are listed. Sign (+/-) indicates positive or negative enrichment. Nine of the

enriched traits are visualized on the data points, with red (yellow) corresponding to a high (low) level of the

trait. For example, *no myth of mans' creation* is found predominantly in cultures near the mobility/exchange

vertex, whereas *deified ancestor worship* is frequent in cultures closest to the resource-competition vertex.

ParTI analysis of the Pulotu dataset of Austronesian cultures

Using ParTI analysis, we find that the distribution of traits for the *Pulotu* cultures is

well described by a triangle (p<10⁻³) (Figure 2A, Supplementary Information 5). We plot this

triangle in the plane of the first two principal components, which explain 20% of the variance

and are significant compared to a phylogenetic null model (Methods). The triangle is

populated almost uniformly by cultures. Many a-priori possible combinations of traits do not

seem to occur, resulting in the empty trait-space around the triangle. Shuffled data does not

fill a triangle (Figure 2B), also when taking into account phylogeny (Supplementary

Information 5).

According to Pareto theory, this triangle indicates three tasks that have tradeoffs

between them. The three vertices of the triangle are combinations of cultural traits that are

potentially adaptive for these three tasks (see Table S1). Below, we infer the three putative

tasks. We label the three vertices by these tasks – Resource Competition, Mobility/Exchange

and Resource Defense - and describe how the tasks were inferred below.

In order to study the relation between the traits and language phylogeny, we displayed

the cultures on a language tree. Cultures are color-coded according to their distances from

the vertices of the triangle (Figure 2C). For example, cultures colored red are closest to the

Resource Defense vertex on the triangle. We note an early-late phylogenetic gradient, from

Resource Defense to Mobility/Exchange to Resource Competition. Similarly, we displayed

the cultures on a geographical map (Figure 2D), showing an east-west geographical gradient

from Resource Defense to Mobility/Exchange to Resource Competition. There are notable

exceptions in which cultures that are close geographically and phylogenetically are not close

on the triangle (have different color codes in Figure 2CD), suggesting that geographical

proximity and phylogenetic inertia are not the only forces at play. Below, we adjusted for

phylogenetic and geographic gradients in our analysis.

Characterization of vertices for *Pulotu*

To characterize the underlying tasks, we asked which cultural traits characterize the

cultures closest to each vertex. For this purpose, we used regression for each trait on its

distance on the triangle from each vertex, adjusted for spatial and phylogenetic effects

(Methods), correcting for multiple hypotheses testing errors. Each of the three vertices shows

distinct cultural traits (Figure 2E).

The cultures closest to the Resource Competition vertex are the stratified Polynesian

cultures (23,24). Closeness to this vertex is strongly associated with *mana* (supernatural

power), especially when linked with descent and social status, and with the *worship of deified*

ancestors. These traits are related to colonization and conquest - founders of colonies become

revered ancestors and their descendants enjoy high status ("founder-focused ideology")

(23,25). Other traits found near this vertex include *low egalitarianism* and the *existence of a*

social hierarchy tapu.

The Mobility/Exchange vertex is associated with cultures that live in areas of low

resource density. The cultures closest to this vertex include the sea-nomad cultures of *Sama*

Dilaut and *Moken* and the sea dwelling people of *Manus*. The Polynesian and Micronesian

cultures that lie on low atolls are also close to this vertex (Supplementary Information 7). The

Mobility/Exchange vertex is enriched with *infrequent conflict between communities*, *high*

reliance on trade and wage labor rather than on agriculture, and *large ritual social groups*

that transcend political communities. It is enriched with *piercing as a feature of rite*, as well

as with *belief in ancestral spirits* (which have a smaller sphere of influence than do *deified*

ancestors). Otherwise, this vertex is characterized by its lacks in religious complexity – *lack*

of mana, *lack of human sacrifice*, *lack of a myth of man's creation*, and *lack of belief in*

deified ancestors and in *nature gods*.

The third vertex, Resource Defense, is enriched with highlander cultures that rely on

agriculture and *land-based hunting* as opposed to fishing and water-based gathering, as well

as with *belief in nature gods*, and with the *belief that the actions of oneself affect the nature*

of one's afterlife.

ParTI analysis of the *BHG* dataset of modern hunter-gatherer cultures

To test the generality of these findings, we analyzed an independent dataset of

modern hunter-gatherer cultures (19) (Figure 3). Considering the first two principal

components, which explain 33% of the variance of this dataset, we again find a distribution

inside a triangle ($p < 10^{-3}$) with three well-defined vertices (Figure 3A, Supplementary

Information 5 and Table S2). For reasons discussed below, we assign the same three labels to

these vertices. As with the *Pulotu* dataset, the traits show geographical trends (Figure 3B),

and hence we performed regression adjusted for phylogenetic and spatial effects

(Supplementary Information 6).

Figure 3. ParTI analysis of *BHG* dataset suggests three adaptive complexes. (A) Data plotted on the first two principal components of *BHG* dataset is well-described by a triangle ($p<10^{-3}$). The three vertices of the triangle are hypothesized to be adaptive combinations of traits for three tasks. (B) Cultures are color coded according to their distance from each vertex on a geographical map. (C) List of traits associated with cultures near each vertex. Sign (+/-) indicates positive or negative enrichment. Six of the traits are displayed on the data distribution, with red (yellow) corresponding to a high (low) level of the trait. For example, *social class distinctions* are strongest close to the resource-competition vertex, whereas *elaborate puberty rites* occur predominantly near the resource-defense vertex.

Characterization of vertices for *BHG*

The hunter-gatherer cultures closest to the Resource-Competition vertex are the stratified societies of the North-American Pacific Northwest, with strong enrichments for *prerogatives of leadership*, *intense and aggressive warfare* and *political complexity*, as well as *sedentism* (Figure 3C). As with the Austronesian cultures, the hunter-gatherers that are the closest to the Mobility/Exchange vertex come from areas with the lowest resource density – the arctic tundra and deserts. This vertex is enriched with *high mobility*, *low inter-group violence* and *low religious and political complexity*. Last, the cultures closest to the Resource

Defense vertex are Australian cultures from the relatively fertile Arnhem Land. Closeness to

this vertex is associated with *low dependence on food storage*, *elaborate male puberty*

rituals, and other traits that relate to *marriage patterns*.

We next asked whether the tasks of the three vertices found in the two datasets might

be similar. This comparison is challenging due to the small overlap in traits between the two

datasets, and the differences in subsistence strategies. We note that in both datasets, the

Resource Competition vertex is enriched with traits that relate to territorial conquest and

social dominance and the cultures near the Mobility/Exchange vertex live in areas with lower

resource density. We tentatively conclude that the vertices might correspond to comparable

tasks, and used information from both datasets to provide an interpretation of these tasks.

Economic defendability theory and the tasks

We find that the three tasks in both datasets can be interpreted using economic

defendability theory, a classic theory from human behavioral ecology which relates variation

in territorial behavior with differences in resource density and predictability (26). This theory

observes that only when a resource is dense and predictable is it worthwhile to engage in

costly behaviors to defend it. Otherwise, other behaviors, such as mobility and exchange, will

be preferred. Economic defendability was first used to explain the evolution of territorial

systems in birds (27), and has been applied to study-small scale human societies (26,28,29).

In agreement with economic defendability theory, the Mobility/Exchange vertex,

which is associated with cultures that live in areas with low resource density, is enriched with

lack of competitive cultural traits. The cultures nearest this vertex are highly mobile; their

reliance on trade or wage labor may compensate for poor and unpredictable resources;

prevalence of piercing may be explained by findings that extreme rituals promote pro-

sociality and inclusive social identity (30); and their large ritual groups may reflect the need

to reinforce cultural identity in the face of dispersal (31). For these reasons, we conclude that

this vertex includes traits that are adaptive for the task of mobility and exchange.

Conversely, when resources are sufficiently dense and predictable, territorial

strategies should emerge. Territorial strategies require two separate tasks (32) – resource

defense and resource competition. The Resource Competition vertex is enriched in both

datasets with socially competitive traits, which are associated with conquest and colonization,

whereas the Resource Defense vertex is associated with traits that are more cooperative.

Resource competition may become more important than resource defense when there

are strong resource gradients (33), unpredictable natural disasters (34) or due to population

pressure (35). The enrichments for hierarchy-related traits near the Resource Competition

vertex may be due to the advantages of hierarchy for group-coordination (36) or for outgroup

exploitation (37).

As an example, the difference between the tasks can be illustrated by the headhunting

practices in two phylogenetically close cultures that lie on two sides of the same island:

Laboya from the western part of the island of Sumba, and *East Sumba* from its less fertile

eastern part. *Laboya* lies close to the Resource Defense vertex while *East Sumba* lies close to

the Resource Competition vertex. Both cultures used to practice headhunting (the practice of

killing people to obtain their heads); however, in West Sumba the motivation for headhunting

was reported to be defense of traditional territories, whereas in East Sumba its motivation

was reported to be territorial expansion (38), in accordance with the proposed tasks for these

vertices.

Discussion

In this study, we tested the hypothesis that functional tradeoffs are important for

adaptation at the level of cultural groups, by using an approach from systems biology to infer

tasks and tradeoffs and applying it multivariate datasets of cultural traits. We find a triangular

geometrical structure of two independent cultural datasets, for Austronesian cultures and

modern hunter-gatherers, in the space of traits. This triangle suggests a fundamental tradeoff

between three tasks at the level of cultural groups. Near each vertex are geographically and

linguistically distinct cultures that have shared features. These features provide clues about

the tasks. Using theory from human behavioral ecology, we identify the three tasks as

mobility/exchange, resource defense and resource competition. We detect specific

combinations of cultural traits that are predicted to be adaptive for each task. The present

approach can in principle be used to infer tasks and tradeoffs for other cultures using multi-

variate datasets.

The results provide numerous hypotheses regarding the adaptive role of cultural traits

with respect to these three putative tasks. While the functional role of many of the traits fits

well with the theory of economic defendability, other suggested associations are surprising.

For example, the association of *taboo* with resource competition rather than resource defense

suggests further investigation.

An important caveat is that by itself, ParTI cannot test whether a specific trait

provides adaptive value, or, instead, whether the trait is linked with another trait that provides

adaptive value. These hypotheses should therefore be tested in future studies that employ

methods such as phylogenetic analysis or experimental psychology. Such studies can, in turn,

help refine our understanding of the tasks and trade-offs that shape the cultural evolutionary
process.

The ParTI approach does not depend on the evolutionary mechanisms at play, nor
does it suppose that a culture is perfectly adapted to its environment. It provides a
quantifiable, geometrical signature for generating hypotheses on the adaptive tasks of human
social organization and cultural evolution. The empty space around the triangle that describes
the datasets corresponds to combinations of cultural traits that could *a-priori* have existed,
but are not found in the data. For example, a combination of small ritual groups (as in
resource defense vertex) together with reliance on trade/wage labor (as in mobility/exchange)
is rare in the *Pulotu* dataset. ParTI, adjusted for phylogeny and spatial diffusion, explains
both empty space and co-occurring traits based on tradeoffs between adaptive tasks.

ParTI might be useful to complement other methods of cultural evolutionary analysis.
It overcomes some of the difficulties of existing approaches. One difficulty is in the
interpretation of correlations between a given cultural trait and environmental factors (39).
For example, collectivist cultural values are correlated with pathogen prevalence, which led
to the hypothesis they may represent an anti-pathogen defense strategy (40). Interpreting such
environmental associations is, however, often ambiguous, since they often relate to the
latitudinal gradient and may be explained by other factors (9,39). ParTI uses multivariate
data, rather than single traits, allowing stronger statistical inference and the ability to adjust
for confounding geographical and phylogenetic factors.

ParTI might also address some of the limitations of other approaches to model the
function of cultural traits, namely optimality models from behavioral ecology (41,42) and co-
evolutionary analyses (43,44). Optimality models require information on costs and benefits
that is not always available. Co-evolutionary models follow traits along phylogenetic trees,
constructed for example from languages. They have been used to study associations such as
the link between human sacrifice and social stratification (20) and the link between livestock
and descent (45). Such models, however, require strong phylogenetic signals and are not
applicable to traits that have weak phylogenetic transmission.

The ParTI approach can apply to any cross-cultural dataset, and thus be used to test
hypotheses on the tradeoffs in cultural adaptations. It provides a top-down way to provide
hypotheses on the function of cultural traits and the trade-offs that shape cultural variation.

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