Innateness and Brain-Wiring Optimization: 
Non-Genomic Nativism

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Our experimental work in computational neuroanatomy has uncovered distinctly efficient layout of wiring in nervous systems. When mechanisms are explored by which such "best of all possible brains" design is attained, significant instances turn out to emerge "for free, directly from physics": In such cases, generation of optimal brain structure appears to arise simply by exploiting basic physical processes, without need for intervention of genes. An idea that physics suffices here--of some complex biological structure as self-organizing, generated without genomic activity--turns attention to limning the role of the genome in morphogenesis. The familiar "nature / nurture" alternatives for origins of basic internal mental structure are that it arises either from the genome or from invariants of the external environment. A third alternative is explored for the neural cases here, a non-genomic nativism.

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The study of minimization of neural connections reveals interrelations between the Innateness Hypothesis and theses associated with the Central Dogma of genetics. The discussion shifts from the usual focus upon abstract cognitive structure instead to underlying brain hardware structure, to hardwired neuroanatomy.

1. Nativism

The familiar, prototypical Innateness Hypothesis is: Abstract mental structure—e.g., relating to knowledge of language (Stich, 1975)—is intrinsic to an organism, for instance genetically determined. But similarly also of course for underlying biological hardware, that is, brain structure, neuroanatomy. A brain is no more plausibly a blank slate with unlimited plasticity in response to its environment than is a mind.

As originally (narrowly) construed, the Central Dogma of genetics was: Genetic information flows one way, from the genome outward to its cellular milieu. That is, DNA \( \rightarrow \) RNA \( \rightarrow \) proteins (Crick, 1958; Watson, 1965). A recent example that may help to define by contrast is the challenge to the Central Dogma perceived in "prion" diseases (e.g., Kuru, Creuzfeld-Jacobsen Syndrome, Bovine Spongiform Encephalopathy (BSE)); prions are non-DNA containing particles that appear capable of replication—and infection—in mammalian central nervous systems (Keyes, 1999). To the Central Dogma we can also add a tacit "Pre-Central Dogma": The genome information encodes an
instruction-set, blueprint, representation, for construction of the organism (e.g., Watson, 1987; DePomera, 1985).

How complete can such a blueprint be? Can it approximate for biological structure-generation a kind of Maxwell's demon micromanaging busybody? The genome instructions cannot be total. A classical regress argument from philosophy of mind can be adapted here, for example, along lines of Ryle (1949). To stop the regression, some basic level of hardware capacities must be assumed, -- a Leggo / Mechanno / Erector-set repertoire of fundamental operations, e.g., the basic physical laws of our universe.

2. Optimized Neuroanatomy

However, a more extensive agenomic domain emerges for some large-scale neuroanatomical structure. The framework is a "Best of all possible brains" hypothesis, in particular, with respect to generative principles to "Save wire". If the brain had an unbounded supply of connections, there would be no pressure to optimize employment of wiring. But in fact connection resources are limited. And since the essential function of a brain is to connect, to make connections, connections seem to have an extremely high value, and so their use has been perfected to a very fine-grained degree. Optimized brain anatomy is detectible in minimized wiring at both micro and macro scale. In some cases, the refinement is discernible down to a best-in-a-billion level (Cherniak, 2000; Cherniak et al, 2002b). Such results begin to approach some of the most precise confirmed predictions in neuroanatomy; why such peculiarly finegrained optimization should
arise itself in turn needs explaining. These wiring optimization problems, typically encountered in computer microchip design, are believed to be intrinsically computationally intractable ("NP-complete"; cf. Garey & Johnson, 1979), unavoidably requiring a combinatorially exploding amount of computation to solve.

For instance, neuron arbor morphogenesis behaves like flowing water (Cherniak, 1992; Cherniak et al, 1999). (i) "Neural fluid mechanics": a fluid-dynamical model for minimized walldrag of pumped flow through a system of pipes will predict the geometry of a variety of dendrite and axon structures almost as well as it predicts configuration of river drainage networks. (ii) Waterflow in branching networks in turn acts like a tree with segments composed of weight-cords-and-pulleys, that is, vector-mechanically; so also do the neuron arbors. The result is that such axons and dendrites globally minimize their total volume to about 5% of optimum for interconnecting their terminal loci. (See Fig. 1.)

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Fig. 1 about here.

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As another instance, vector mechanics suffices for optimization of placement of the ganglia (neural subnetworks) of the roundworm *Caenorhabditis elegans* (Cherniak, 1994a; Cherniak et al, 2002a). "The web that weaves itself": Some of the above self-organizing model of arbor optimization also can serve as a mechanism of ganglion placement in the roundworm nervous system. Our prior work (Cherniak 1995) had
found that the actual observed positioning of the ganglia in the worm was optimal, in that it required the least total wirelength for the animal's interconnections. We have now constructed a force-directed placement simulator, Tensarama, where each of the worm's connections behaves like a micro weights / cords / pulley system. This vector-mechanical net outputs the actual minimized ganglion layout by converging on energy-minimization equilibrium at the actual positioning of the ganglia--without much susceptibility to local-minima traps. (See Fig. 2.)

Fig. 2 about here.

So, the hypothesis is that (a) Optimization accounts for a significant extent of observed anatomical structure. An additional hypothesis is that (b) Simple physical processes are responsible for some of this optimization. Combined, then, the picture is:

Physics \rightarrow Optimization \rightarrow Neural structure

That is, self-assembling brain structure "for free, directly from physics", i.e., generated via simply exploiting basic physical processes, without need for intervention of genes. Physics suffices; complex biological structure as self-organizing. So, besides conserving connections, another possible explanation for the extreme level of optimization observed might be just that it is a side-benefit of complex biostructure hitching a free ride from physics.

-- In particular, for minimal-wiring neuroanatomy for dendrites,
axons, and roundworm ganglia (Cherniak et al, 2002a). Now also we have observed even finer wiring optimization of cerebral cortex layout of cat and macaque monkey (Cherniak, 2000; Cherniak et al, 2000b). Thus, a harmony of physics and neuroanatomy via fluid-mechanics and mesh of springs; a sort of plate tectonics of the brain. Some of the methodological significance here is that discrete-state processes (e.g., as in mutation of genomic sequences (Mitchell, 1996) are not required; continuous-process models can suffice for neural wiring optimization. This work falls in the tradition of seeking simple underlying mathematical form in complex aspects of Nature, ranging from Pythagoras through D'Arcy Wentworth Thompson (1917/1961).

3. Non-Genomic Nativism

Hence, contrary to a "nature / nurture" dichotomy, a third possibility-zone--"nature" here includes not only biology, but also mathematics and physics. A via media between two extremes: (a) Brains of course cannot grow like crystals, entirely non-genomically. (b) Yet life must still play by the rules of the game, subject to mathematical and physical law. So, there is a division of labor between the genome and simple physical processes. The organism's genome is written not upon a tabula rasa, but upon a specifically pre-formatted, pre-printed form or slate, already inscribed with a significant proportion of structural information.

Perhaps such a picture of interpenetrating domains clashes with some metaphysics hardwired in human beings, with a category structure that draws a bright line between the animate and the inanimate. Also,
the expression "non-genomic nativism" may sound like a bit of a solecism to sharper ears. (Piaget, "That which is inevitable does not have to be innate." (1970)) Of course, label choice is not of interest. The underlying point is that some complex biological structure--comparable in extent to genome-specified structure--is intrinsic, inborn, yet not genome-dependent.

One rationale for organisms to exploit such free anatomy is the vast mismatch of scale between brain structure and genome structure. The human brain is commonly characterized as the most complex physical structure known, yet the total information representation capacity of the human genome is comparatively small. After discounting noncoding introns (about 95% of the total), the amount of brain-specific DNA available might amount to as little information as is contained in a desk dictionary (about 50,000 entries, ~ 100 Mb total). Hence, information on brain structure must pass through a "genomic bottleneck" constraint on DNA information-representation capacity (Cherniak, 1988, 1992). Brain structure for free lowers this genome information-carrying load.

One caveat in interpreting genome information limitations concerns data compression. The key idea, from algorithmic information theory (Li & Vitanyi, 1997; Chaitin, 1987), is: For a given symbol sequence, what is the smallest program of a given format that will generate the sequence (and then halt)? A related concept (intimately connected with unsolvability of the Halting Problem) is that of, e.g., a 5-state "Busy Beaver" Turing machine--a 10-line Turing machine that
generates the largest number of 1's for its size on an initially all-
blank tape and halts; the most productive such program presently known
outputs 4,098 1's (Dewdney, 1993). (See Fig. 3) 6-state TM's are now
known with much vaster output. Thus, one can ask, what is the minimum
such program that emits the text of Anna Karenina, or the bitstream
soundtrack of a Beethoven quartet performance, or that generates a
given genome sequence? One could imagine such information compression
occuring on the genome to help pack in extensive brain structure
specification. However, it seems a mistake to go on to conclude that
this data compression is in practice virtually unlimited; for
compression / decompression itself entails computation costs. For
instance, the above 5-state busy beaver candidate requires over 11
million steps to generate its string of 1's. So, no free lunch here:
storage space in effect is traded off for encode / decode time.

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Fig. 3 about here.

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4. Misgivings

An uneasy "biology from physics" coda may help in articulating
nongenomic nativism. Proceeding from static structure to behavior,
physiology: That is, let us explore instances of complex biological
functions similarly originating directly from simple physical
processes. For example, as explained above, Tensarama yields the
neuroanatomical layout of C. elegans by simulating each of the worm's
thousand connections as approximately a microspring; as the system
proceeds to vector-mechanical equilibrium, it weaves the ganglia into the actually observed layout--a one-in-ten-million achievement. Of course, such complex anatomy-generation can in itself be pictured instead as a slow-motion intelligent behavior.

A corresponding behavioral instance would be to derive realistic locomotion behavior similarly from a mesh of springs. In this connection, the "Sodaconstructor" package yields strikingly lifelike crawling motion for a simulated "worm" device, walking for a generic quadruped, etc. (Sodaplay.com). Yet all this animation is generated from what amounts to again just a mesh of springs (with masses), driven by a single simple sine wave. Another case is a random-path searcher device: a simple smooth sphere containing a motor-driven counterweight will robustly outperform robots with conventional obstacle detection and evasion chip software. Simulated annealing is an example of a system using only basic physical processes (thermodynamic temperature-schedule models) to solve search problems that have local minima traps (Kirkpatrick et al, 1983). Also suggestive is the idea of mechanisms that exploit chaos-theoretic (nonlinear dynamical) phenomena--e.g., intricate behavior from a simple device, such as a jointed pendulum (Thompson & Stewart, 1986).

One question is how far such a for-free-from-physics approach can proceed; what is the most complex behavior so derivable? (For examples of intricate behavior from simple rotational dynamics, see Walker, 1985.)

So, the obvious leap to a conclusion would be a sunny picture, of
complex neuroanatomical structure "for free, directly from physics". And similarly also of complex behavior derived directly from basic physical processes, the latter again partaking of the Pythagorean tradition of D'Arcy Thompson (1917/1961), of simple mathematical form in Nature. However, a darker picture also crystallizes, of a retrogression back to a familiar type of neoreductionism: rather than progress, a recantation of a generation of philosophy of mind threatens.

Let us contemplate the development of mind / brain sciences in the post-behaviorist era, since c. 1970. Functionalist / computationalist theories of mind emerged, opening the conceptual possibility of mental states not having to be identified just with physical hardware states, but instead with abstract software states (Putnam, 1964). The corresponding idea in genetics was of course where we began the paper, the picture of genome as program. However, by now some of us may wonder whether AI has perhaps not fulfilled its early promise. For instance, compare the 1966 "Eliza" conversationalist program (an anti-AI project, really) (Weizenbaum, 1976) with the remarkably unimpressive "Alice"--recent two-time winner of the Loebner Turing Test prize (Alice, 2001). AI and von Neumann-architecture machines both emerged a half century ago: if hardware had developed as has AI, we would still be using abacuses and sliderules--computers would merely be exotic laboratory confections. Hardware and software engineering differ profoundly (Cherniak, 1988). Perhaps faith here in the future of progress may begin to waver.
The above 5-state Busy Beaver candidate, although of course itself a program, illustrates another potential Pyrrhic victory for computationalism. This TM table was discovered by a huge constrained brute-force search of many possible TM tables by a program written by Heiner Marxen, of the Technical University of Berlin. When one contemplates the TM's operation, it indeed displays a kind of inhuman, unintelligible elegance—it seems repeatedly to re-embroider segments of its tape, to re-use parts of its own code palindromically, etc. Rather than incomprehensibly huge, it is incomprehensibly compact (a "nano-kluge"). And so an idea emerges of unintelligible intelligence, of programs that might generate intelligent behavior, but without identifiable representations of their world, rules for using such models, etc. This corresponds to the above picture of intelligence for free from physics. What turns out to work efficiently may not be—indeed, may be antagonistic to—what is humanly comprehensible (Cherniak, 1988). (The satirist Emo Phillips proposed, in another context, a universal non-denominational prayer, (approximately) Lord, Please arrange the Universe for my convenience.)

Yet I suspect anyone who actually experienced the aridity of the empty-organism era could not regard its revival in any form with equanimity. I conclude, or fail to conclude, with this anomie, this uneasy equivocation. More is to be done; I do not now know how. A variation upon Matthew Arnold's lines that Henry Adams (1918/1973) took as motto comes to mind: to find oneself caught between
worldviews, one declining, one struggling to bring itself into existence.
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REFERENCES


ALICE (2001) [2000 & 2001 Loebner "Turing Test" Prize Winner]
www.alicebot.org


www.glue.umd.edu/~cherniak/philcomp/


Sodaconstructor locomotor simulator package: www.sodaplay.com


**Fig. 1.** Complex biological structure arising directly from basic physics: Nerve cell anatomy behaves like flowing water, and waterflow in turn acts like a tree composed of springs. -- "Instant arbors, just add water." The dendrite (input) and axon (output) branchings are portions of mammalian neurons, the river is an experimentally generated drainage network. In each of the three cases, the actual structure is within a few percent of the optimal, minimum-volume configuration shown; this is evidence of goodness of fit of the physical optimization model (Cherniak et al, 1999). Diagram: Mark Changizi.
TENSARAMA

Head
0 0 1 1 2 2 3 3 4 4 5
0 5 0 5 0 5 0 5 0 5 0 Tetrons

PH (100.000000)
AN (300.000000)
RNG (440.000000)
DO (506.000000)
LA (564.000000)
VN (744.000000)
RV (948.000000)
VCA (1856.000000)

VCP (3856.000000)
PA (4726.000000)
DR (4810.000000)
LU (4884.000000)

Final layout popped out after: 100,000 iterations  [@final 100]
Tension Constant: 0.010000
Total Wirecost: 87802.750000 um

Fig. 2. Runscren for "Tensarama", a force-directed placement algorithm for optimizing layout of ganglia of the nematode Caenorhabditis elegans: that is, minimizing total length of interconnections. This vector mechanical simulation represents each of the roundworm's ~ 1,000 interconnections as a sort of micro-spring acting upon the horizontally movable ganglia (nervous system clusters) "PH", "AN", etc. (Connections themselves do not appear on runscreen, nor fixed components such as sensors and muscles.) The above screen dump shows the final configuration of the system after 100,000 iterations (re-update cycles for forces and locations): The system has found the global minimum-cost positioning of the ganglia (with about 8.7 cm total of wire) -- which is also the actual layout. In this way, physics suffices to generate this neuroanatomical structure, out of ~ 40 million alternative possible configurations (Cherniak, 1995; Cherniak et al, 2002a).
\begin{center}
\begin{tabular}{lllll}
  CST & INP -> & OUT & MOV & NST \\
  \hline
1    & 0       & 1    & L   & 2 \\
1    & 1       & L    & 1   & \\
2    & 0       & 1    & R   & 3 \\
1    & 1       & R    & 2   & \\
3    & 0       & 1    & L   & 1 \\
1    & 1       & R    & 4   & \\
4    & 0       & 1    & L   & 1 \\
1    & 1       & R    & 5   & \\
5    & 0       & 1    & R   & 0 \\
1    & 0       & R    & 3   & \\
\end{tabular}
\end{center}

\textbf{Fig. 3.} Turing machine program that has been the contender for title of 5-state "busy-beaver" -- maximally productive TM program -- without challenge for over a decade (Dewdney, 1993). It takes 11,798,826 steps to generate 4,098 "1"s on an initially blank tape before it halts. Thus, the program illustrates ~ 1:100 data-compression, which genome encoding might exploit. However, the program also illustrates the high computation cost of message "decompression" (a human being with pencil and paper would take about a year to complete the computation; a TM simulator (Cherniak, 1990) takes a few hours).