

tasks, providing another window to prevention through psychomotor training (2). Finally, with the use of methods that build on those of Manuca and Savit and the team at the University of Michigan, we have found that more than half of epileptic crises can be anticipated by a brief window and are amenable to electrical intervention of the kind described in Glanz's article (3).

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References

1. M. Le Van Quyen, J. Martinerie, C. Adam, H. Schuster, F. Varela, *Physica E* **56**, 3401 (1997).
2. M. Le Van Quyen *et al.*, *Neuroreport* **8**, 1703 (1997).
3. C. Adam *et al.*, *Epilepsia* **38** (suppl. 3), 217 (1977); J. Martinerie *et al.*, in preparation.

Glanz quotes one investigator to the effect that neural noise does not have clinical application as yet. But three decades ago, it was discovered that the pupil motor response has a neural noise component (1).

We uncovered a signal processing method to isolate this noise from the rest of the pupil response and found that the noise variance had a significant clinical application in that it could accurately (90% confidence) distinguish individuals with attention deficit disorder (narcoleptics) from normal (control) individuals (2). In memory experiments, we also observed that neural noise has a dramatic impact on cognitive performance. We believe that the reticular activating system controls the noise we measure and, if so, there should be more clinical applications for attention deficit disorders in the future.

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References

1. S. F. Stanton and L. Stark, *IEEE Trans. Biol. Eng.* **13**, 140 (1966).
2. W. D. O'Neill *et al.*, *J. Sleep Res.* **5**, 265 (1997).

Haeckel's Embryos

A recent paper that we authored with several colleagues challenges the idea that there is a

single, highly conserved embryonic stage in the vertebrates (1). Principally because of our conclusions regarding the inaccuracies of drawings made by the 19th-century naturalist-philosopher Ernst Haeckel, our paper received considerable coverage in the popular and scientific press (2), including an article by Elizabeth Pennisi ("Haeckel's embryos: Fraud rediscovered," *Research News*, 5 Sept., p. 1435). Regrettably, in the resulting debate over Haeckel and the reality of the vertebrate "phylotypic" stage, what we regard as one of the main implications of our results for contemporary studies of the developmental basis of evolutionary change has been largely overlooked.

The idea that there is an identical embryonic stage (the phylotypic stage) common to all vertebrates implies that changes in development that underlie the considerable variation in adult morphology of these animals appear only later in ontogeny. In this sense, the concept of the phylotypic stage is an explicit statement, or hypothesis, about the temporal deployment of evolutionary changes in development. Yet we show that at least some significant differences in adult morphology, involving characters as fundamental to the vertebrate body plan as limbs and somites, begin to appear before, and are apparent at, the putative phylotypic stage. These and similar observations (3) seriously diminish the validity and applicability of the phylotypic stage concept for the vertebrates. More important, they remind us of the potential significance of earlier developmental events to the determination of animal form, and that these too are frequent targets of evolutionary perturbation.

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References

1. M. K. Richardson *et al.*, *Anat. Embryol.* **196**, 91 (1997).
2. N. Hawkes, *The Times (London)*, 12 Aug. 1997, p. 14; B. K. Hall, *Trends Ecol. Evol.* **12**, 461 (1997).
3. M. K. Richardson *et al.*, *Development*, in press.

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