

Weather forecasts still tricky

SIR—The recent paper on 'Fractal characterization of inhomogeneous geophysical measuring networks' by Lovejoy *et al.*¹ and a discussion in *News and Views*² address the important problem of detectability of sparse atmospheric phenomena and, therefore, of the very feasibility of useful weather forecasting. Now, whatever the difficulty of its detection might be, a sparse phenomenon will exert an influence on the *time course* of a multitude of more systematic phenomena occurring over a wide range of space scales. The main reason for this is the fundamental instability of the atmospheric system³: a perturbation arising from an uncontrollable localized event and acting on a larger scale phenomenon pertaining to atmospheric circulation may give rise to an (exponentially) divergent history and to an altogether different regime, or to an aperiodic sequence of regimes perceived by an unaware observer as 'noise'.

Let γ be the precision with which the initial state of the system can be determined in phase space. Assuming that the atmosphere behaves as an unstable dynamical system we obtain the following estimate for the error growth after time t :

$$\gamma(t) = \gamma e^{Kt} \quad (1)$$

where K is the largest positive Lyapounov exponent (or more generally the Kolmogorov entropy).

Since our variables are distributed in physical space, an inadequate spatial resolution l will clearly induce an error in the estimate of the variables. Conversely, if for some reason their values are subject to uncertainty, it will be difficult to locate with precision the region of space from which these variables had initially emanated. We may therefore substitute, in our crude estimate, the phase space uncertainty γ by the precision l with which the initial state of the atmosphere can be determined in the physical space and write, instead of (1)

$$l(t) = l e^{Kt} \quad (2)$$

When the size $l(t)$ will reach a value of the order of the dimensions of the entire system, L , predictions concerning individual histories will become meaningless and should be replaced by statistical considerations. The time τ_m for which this will happen is⁴:

$$\tau_m \approx \frac{1}{K} \ln \frac{L}{l} \quad (3)$$

This gives a quantitative measure of the limits of predictability of the system. A significant feature of τ_m is its logarithmic dependence on the parameter l , and thus its relative insensitivity to space resolution. Going from a quasi-uniform to a fractal distribution of l or vice versa is not going to change this picture drastically.

So although detection of sparse phenomena is important in its own right, it is

worth stressing that even with a considerable improvement of the world detection network the problem of forecasting of the global state of the atmosphere is likely to remain as acute as ever.

C. NICOLIS

*Institut d'Aéronomie Spatiale de Belgique,
1180 Bruxelles, Belgium*

G. NICOLIS

*Faculté des Sciences,
Université Libre de Bruxelles,
1050 Bruxelles, Belgium*

1. Lovejoy, S. *et al.* *Nature* **319**, 43–44 (1986).
2. Hollingsworth, S. *Nature* **319**, 11–12 (1986).
3. Lorenz, E.N. *Tellus* **36A**, 98–110 (1984).
4. Schuster, H.G. *Deterministic Chaos* (Physik, Weinheim, 1984).

Activation of cellular calcium entry

SIR—A recent *News and Views* piece by Baker¹ discusses the mechanisms of intracellular calcium ion (Ca^{2+}) release from the endoplasmic reticulum (ER) but refers briefly to cellular calcium entry from the extracellular fluid as being mediated by Ca^{2+} channels gated either chemically or by voltage. Voltage-activated Ca^{2+} channels that can be modulated by intracellular messengers have been characterized in many systems², but are unlikely to be present in a number of different epithelial tissues such as exocrine glands and liver simply because they are unable to fire action potentials and many activating hormones or neurotransmitters evoke membrane hyperpolarization rather than depolarization in these cells³. Nevertheless there is clear evidence that hormones and neurotransmitters can increase unidirectional Ca^{2+} flux into, for example, pancreatic acinar cells⁴.

The mechanisms underlying this receptor operated Ca^{2+} entry are at present obscure although a link with the increased turnover of phospholipids after stimulation has been suspected for more than ten years⁵ with evidence for such a link being particularly clear in relation to Ca^{2+} entry in blowfly salivary glands⁶. A recently formulated hypothesis⁷ suggests that receptor operated Ca^{2+} entry is secondary to the emptying of an intracellular Ca^{2+} pool mediated by the action of inositol trisphosphate (IP_3) on an electrogenic Ca^{2+} pathway (channel) in the ER^{8,9}. The well-established biphasic nature of agonist-activated Ca^{2+} mobilization is, according to the new hypothesis, explained by an initial emptying of the intracellular (ER) Ca^{2+} pool by IP_3 followed by entry of extracellular Ca^{2+} into the pool and, in the continued presence of IP_3 , into the cytosol. On removal of the stimulant, IP_3 is rapidly broken down closing the Ca^{2+} channel in the ER and Ca^{2+} entry from the outside continues only until the content of the ER pool has reached a level that prevents further entry.

In this model⁷ intracellular Ca^{2+} release and Ca^{2+} entry are both controlled by the same messenger, IP_3 . Ca^{2+} entry is therefore not directly linked to the hormone receptor, but is messenger-mediated. A recent patch-clamp study on pancreatic acinar cells¹⁰ demonstrating an external Ca^{2+} requirement for sustained K^+ channel (Ca^{2+} -activated) opening evoked by a peptide belonging to the cholecystokinin-gastrin group may be relevant in this context since the Ca^{2+} -requirement could be localized at a site separated from the hormone-receptor interaction indicating messenger-mediated rather than receptor-mediated Ca^{2+} uptake. These recent findings¹⁰ appear to be compatible with the hypothesis forwarded by Putney⁷.

OLE H. PETERSEN

*MRC Secretory Control Research Group,
University of Liverpool,
Liverpool L69 3BX, UK*

1. Baker, P.F. *Nature* **320**, 395 (1986).
2. Reuter, H. *Nature* **301**, 569–574 (1983).
3. Petersen, O.H. & Maruyama, Y. *Nature* **307**, 693–696 (1984).
4. Kondo, S. & Schulz, I. *Biochim. biophys. Acta* **419**, 76–92 (1976).
5. Michell, R.H. *Biochim. biophys. Acta* **415**, 81–147 (1975).
6. Berridge, M.J. & Fain, J.N. *Biochem. J.* **178**, 59–69 (1979).
7. Putney, J.W. *Cell Calcium* **7**, 1–12 (1986).
8. Streb, H., Irvine, R.F., Berridge, M.J. & Schulz, I. *Nature* **306**, 67–69 (1983).
9. Mualem, S., Schoeffield, M., Pandolf, S. & Sachs, G. *Proc. natn. Acad. Sci. U.S.A.* **82**, 4433–4437 (1985).
10. Suzuki, K., Petersen, C.C.H. & Petersen, O.H. *FEBS Lett.* **192**, 307–312 (1985).

Beam balance test of weak equivalence principle

SIR—In the recent article by Fischbach *et al.*¹, commenting on a *News and Views* piece², a new hypercharge force is postulated as being responsible for various apparently anomalous effects in physics. Although there may not be unanimity in the interpretation of the Eötvös data in terms of this new force³, its existence would have such far reaching consequences in physics that further experiments are clearly required. As suggested by Fischbach *et al.*, further Eötvös-type or galilean experiments come to mind. Here we propose an experiment using a common balance which would allow the force, if it exists, to be measured with good precision. The experiment would be as follows.

The weights of the two test masses made of materials having different values of (B/μ) (see ref. 1 for notation) are compared using a beam balance at two different sites. The first site is in a tunnel deep inside a mountain where it can be assured that there are some kilometres of solid rock in all directions. Here, in the absence of any net hypercharge force, the test masses are adjusted to have apparently equal masses as measured using the balance. The balance and test masses are then moved out of the tunnel to a second site on the surface of the Earth, preferably