Fusion in 1947?

Sir-Fleischmann and Pons' claim to have achieved fusion at room temperature in a glass container. I was struck by the similarity to a much earlier experiment by Rayleigh², which demonstrated that a nitrogen (or oxygen or hydrogen) atom impinging on a metal wire in a glass container releases energies ranging up to \sim 100 eV. At the time, no explanation of the results was advanced, although physicists found no fault with the experiment. In 1957, Burgess and Robb³ showed that in the presence of traces of oxygen (0.22-0.99 mm Hg) hydrogen atoms will cause the temperature of a metal wire to rise many times above that expected to result from the heat of recombination of H atoms on the wire; this does not, however, explain Rayleigh's results with nitrogen or oxygen. Have they been explained or can we count Rayleigh's experiment as an earlier observation of 'fusion' in a bottle?

KEITH TAYLER

11 Kemp Street. Brighton. East Sussex BN1 4EF, UK

- 1. Fleischmann, M. J. & Pons, S. J. electroanalyt. Chem. 261 301-308 (1989).
- Lord Rayleigh *Proc. R. Soc.* A **189**, 296–299 (1947). Burgess, R.H. & Robb, J.C. *Trans. Faraday Soc.* **54**, 1008–

Sympatric origins of R. pomonella

SIR—Carson raises two important questions concerning the possible recent sympatric origin of the apple infesting form of the maggot fly Rhagoletis pomonella¹. First, he points out that R. pomonella may have shifted to domestic apples (Malus pumila) from a native host other than hawthorns, such as endemic crabapples. Second, he postulates that different races of R. pomonella may exist on different species of hawthorns and that the apple race could have been derived from just one of these 'hawthorn races'. We have addressed these issues in some detail in manuscripts submitted to the journals Evolution and Heredity and we will therefore highlight only a few of the more crucial reasons why the alternative explanations are unlikely.

The most likely potential native North American hosts for R. pomonella other than hawthorns are several species of crabapple (M. angustifolia, M. diversifolia, M. glabrata, M. ioensis and M. coronaria), none of which supports populations of the fly²⁻⁴. The reason for this is not clear, but O'Kane² has suggested that fruits of native crabapple species are too acidic and ripen too late for R. pomonella to complete development before winter. Pree has found a correlation between larval mortality and the total phenol content of fruit from several introduced species and varieties of crabapples, which suggests that phenolics also contribute to the unsuitability of native crabs as hosts.

Furthermore, in the highlands of Mexico, an isolated population of R. pomonella has independently shifted from hawthorns to domestic apples within historical times (apples were introduced into Mexico from Spain in 15226). Both the apple and hawthorn fly populations in Mexico are morphologically distinct from flies in the north-east⁴. Mexican R. pomonella could not, therefore, have given rise to the apple race in New England. However, Mexico does not have any endemic species of crabapple⁶ (J. personal communication). Therefore, in Mexico at least, R. pomonella seems to have shifted directly from hawthorns to domestic apples.

It is still possible, of course, that different races of R. pomonella exist on different species of hawthorns. Almost 100 different species of Crataegus, comprising 19 different species groups, may be endemic to North America7-9. But the taxonomic status of most of these Crataegus species is questionable as hybridization, polyploidy and apomixis are common in the genus⁹⁻¹². Nevertheless, R. pomonella has been reported to attack only a restricted set of Crataegus 'species', with infestations confirmed for only 14 endemic hawthorns^{4,13-16}. In the northeastern United States, only hawthorn species that produce relatively large, soft fruits ripening in early autumn are heavily parasitized by the fly2.4.13. It therefore seems that R. pomonella uses only a few potential hawthorn hosts in the northeast, a trend that is not conducive to the formation of different hawthorn races.

In addition, electrophoretic analysis of R. pomonella populations infesting C. punctata, C. brachyacantha, C. douglasii, C. mollis and C. monogyna from across the United States provides no evidence for differentiated genetically hawthorn races¹⁴⁻¹⁶. Instead, we find that latitudinal allele-frequency clines exist in the eastern United States among both apple and hawthorn populations (mainly C. mollis) for five of the six allozymes showing host associated differentiation^{15,16}. The slopes of these allele-frequency clines differ, however, between the two races with the hawthorn race having steeper clines16. Inter-host divergence is, therefore, superimposed on north-south clinal patterns of intra-host variation such that the magnitude of genetic divergence between hawthorn and apple flies is a function of latitude. Consequently, different apple populations do not cluster together as a discrete genetic subdivision from populations attacking C. mollis, as would be expected if the apple race was formed from a distinct race infesting a hawthorn species other than C. mollis.

Instead, the electrophoretic results

suggest that differential selection (affecting development rates for the fly and related to ambient temperature conditions) and differences in host use are key factors responsible for the pattern of allozyme variation for R. pomonella. Although further sampling of R. pomonella at field sites with sympatric hawthorn species is needed to discount completely the possibility of host-specific hawthorn races, all available data indicate that hawthorn populations represent a single race displaying extensive latitudinal variation.

Guy L. Bush

Department of Zoology, Michigan State University. E. Lansing, Michigan 48824, USA

JEFFREY L. FEDER*

Department of Biology, Princeton University. Princeton. New Jersey 08544, USA

STEWART H. BERLOCHER

Department of Entomology, University of Illinois. Urbana, Illinois 61801, USA

BRUCE A. McPHERON

Department of Entomology, Pennsylvania State University, University Park. Pennsylvania 16802, USA

D. COURTNEY SMITH

Department of Biology, University of Utah. Salt Lake City, Utah 84112, USA

CHARLEY A. CHILCOTE

Department of Natural Resources, University of Michigan, Ann Arbor, Michigan 48109, USA

- Carson, H.L. *Nature* **338**, 304 (1989).
 O'Kane, W.C. *The Apple Maggot* (New Hampshire Exp. Stat. Bull. 171, 1914).
- Porter, B.A. The Apple Maggot (US Dept. Agric. Tech. Bull. 66, 1928).
- Bush, G.L. The Taxonomy, Cytology, and Evolution of the Genus Rhagoletis in North America (Diptera: Tephritidae) (MCZ, Cambridge, Massachusetts, 1966).
- Pree, D.J. J. econ. Ent. 70, 611-614 (1977)
- Standley, P.C. Trees and Shrubs of Mexico (Smithsonian, Washington, DC, 1922).
- 7. Fernald, M.L. Gray's Manual of Botany 8th edn. (American Book Co, New York, 1950). Correll, D.S. & Johnston, M.C. *Manual of the Vascular*
- Plants of Texas (Texas Res. Fdn, Renner, 1970).
 9. Muniyamma, M. & Phipps, J.B. Can. J. Bot. **63**, 1319–
- 1324 (1985). Muniyamma, M. & Phipps, J.B. Can. J. Bot. 62, 2316– 2324 (1984).
- Phipps, J.B. Ann. Mo. Bot. Gdn 70, 667-700 (1983).
- 12. Phipps, J.B. in *Plant Biosystematics* 417–438 (Academic Press, Toronto, 1984).
- Wasbauer, M.S. An Annotated Host Catalog of the Fruit Flies of America North of Mexico (Diptera: Tephritidae) (Dept. Agric., Sacramento, California, 1972).
- Berlocher, S.H. thesis, Univ. of Texas at Austin (1976). McPheron, B.A. thesis, Univ. of Illinois at Urbana (1987).
- Feder, J.L. thesis, Michigan State Univ. (1989).