ger RNA from that carrying the mdx4cv mutation. As shown in Fig. 1b, c, a 5' primer specific to the wild-type sequence allowed amplification of a 240-base-pair fragment encompassing exons 53 and 54 of mRNA extracted from muscle tissue of con-C57Bl/6 and all transplanted animals, but not from mRNA from mock-transplanted mdx4cv mice. All PCR reactions were internally normalized by simultaneous amplification of a 378-basepair fragment encompassing exons 33–36.

Our results indicate that repair by bonemarrow-derived cells never exceeds 1% of total muscle fibres during the lifespan of transplanted mdx mice. Transplantation of total bone marrow, or even of highly enriched stem-cell fractions from bone marrow², therefore has a very limited impact on muscle-cell replacement, and no ameliorating effect on murine muscular dystrophy. This might be explained by poor recruitment of bone-marrow cells to the dystrophic muscle, or by bone-marrow-derived cells having no selective advantage over resident myogenic precursors, which regenerate muscle throughout life in *mdx* mice.

We do not address strategies for overcoming these limitations here, although they are likely to be the focus of future research. It should be stressed that the mdx mouse model does not reproduce the clinical picture of Duchenne's muscular dystrophy in humans, in which muscle degeneration gives rise to fibrosis rather than to repair⁵, so it remains possible that dystrophin-producing cells derived from bone marrow might be subject to positive selection in a human context.

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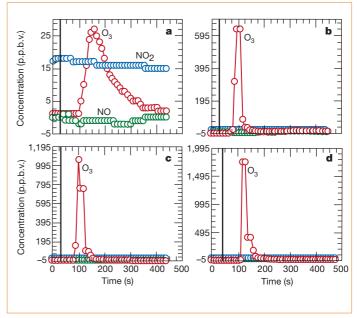
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- Ferrari, G. et al. Science 279, 1528–1530 (1998).
- 2. Gussoni, E. et al. Nature 401, 390-394 (1999).
- 3. Bittner, R. E. et al. Anat. Embryol. 199, 391-396 (1999).
- 4. Danko, I. et al. Pediatr. Res. 32, 128-131 (1992).
- 5. Emery, A. E. H. Duchenne Muscular Dystrophy 2nd edn (Oxford Univ. Press, Oxford, 1993).

Formation of ozone by fireworks

zone is a secondary pollutant and greenhouse gas that is formed from molecular oxygen in the presence of sunlight and nitrogen oxides. The extent of production also depends on the presence of volatile hydrocarbons, carbon monoxide and methane1-6. But we have discovered a surprising source of ozone which is generated in spontaneous bursts even in the

Figure 1 Ambient concentration (parts per billion by volume) of O₃ (red circles), NO (green) and NO₂ (blue) in the environment as a function of time after lighting colour-emitting 'pencil' sparklers. a-d, Values correspond to the simultaneous burning of one, three, four and six sparklers. respectively: vertical black lines indicate the duration of burning for the sparklers (starting at 0 s). O₃ begins to accumulate in each case at around 100 s; this time lag corresponds to the volume of air in the assembly of the Teflon tube and manifold attached to the O3 and NO2 analysers. During the experiments, the open end of the



Teflon tube leading to the analysers was kept 900 cm away from the sparklers. The average weight of inflammable material (initial weight minus weight after burning) was 9.15 ± 0.85 g per sparkler. The burn time varied from 26 s for one sparkler to a maximum of 42 s for six; this variation was due to differences in the weight of the burning mixture. O_3 formation was independent of microclimatic factors such as temperature and humidity of the ambient air. The concentrations of NO and NO2 did not change during the O3 burst. Further details are available from the authors.

absence of sunlight and nitrogen oxides namely, the exuberant mass of colouremitting sparklers that are lit during the Diwali festivities, which take place every year during October and November in Delhi, India. The underlying process of ozone formation resembles that induced by ultraviolet radiation in the stratosphere^{7,8}.

We undertook a routine, real-time monitoring of the concentrations of NO, (NO and NO₂), O₃ and other microclimatic factors at a known pollutant-receptor site9 in Delhi in order to determine the effects of burning unprecedented numbers of fireworks on the local environment during the festive period in November 1999. We found that during the festival period the ozone concentration peaked at around noon and fell to negligible levels after sunset.

On Diwali night (7 November), a small build-up of O_3 (9 ± 1 parts per billion by volume) was detected between 20:40 and 02:30 hours (results not shown). During this period, no correlation was found between NO_x concentration and O₃ formation, indicating that the ozone was unlikely to have been generated in reactions involving ambient NO_x. This observation was surprisingly different from night-time O₃ measurements obtained on the other dates, when no O₂ formation was detected.

Further experiments carried out under different climatic conditions showed that there is a linear regression between the total amount of inflammable material present in sparklers and the cumulative O3 formed (correlation, 0.993). There was no change in ambient NO_x concentration before, during or after these experiments (Fig. 1).

Sparklers depend on a combination of different metal salts to generate their colour and sparkle¹⁰ — these include potassium perchlorate, sulphur, strontium nitrate, barium nitrate, sodium oxalate, calomel, aluminium and manganese. When burnt, a significant proportion of the light emitted by these constituents has a wavelength below 240 nm (ref. 11). The radiative energy of these emissions is sufficient to dissociate atmospheric molecular oxygen into atomic oxygen, enabling the reaction $O_2 + O \rightarrow O_3$ to take place. This proposed mechanism could explain the formation of bursts of O₃ without the participation of NO_v, and is therefore similar to the process of ultraviolet-radiation-induced formation of O_3 in the stratosphere^{7,8}.

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- 1. Ramanathan, V., Singh, H. B., Ciceron, R. J. & Kiehl, J. T. J. Geophys. Res. 90, 5547-5566 (1985).
- 2. Lacis, A. A., Weubbles, D. J. & Logan, J. A. J. Geophys. Res. 95, 9971-9981 (1990).
- Wang, W. C. & Sze, N. D. Nature 286, 589-590 (1980).
- 4. Crutzen, P. J. & Zimmermann, P. H. Tellus 43A-B, 136-151(1991).
- Walcek, C. J. & Yuan, H.-H. J. Appl. Meteorol. 34, 1056-1069 (1995).
- 6. Johnson, D. W. et al. Nature 353, 617-621 (1991).
- Chapman, S. R. Meteorol. Soc. Mem. 3, 103–125 (1930).
- 8. Brausser, G. P. et al. J. Geophys. Res. 95, 5639-5655 (1990).
- 9. Singh, A. et al. Atmos. Environ. 31, 3421-3427 (1997).
- 10. Holmes, H. H. in The Encyclopedia Americana Int. edn, Vol. 11, 263-265 (American Corp., New York, 1983).
- 11. Reader, J. & Corliss, C. H. in CRC Handbook of Chemistry and Physics 63rd edn (eds Weast, R. C. & Astle, M. J.) E205-E368 (CRC, Florida, 1982-83).