adjusted mortality from asthma in Italy increased by over fivefold (figure), and the rises were larger in middle and older age than in children and young adults.³ This discrepancy therefore indicates that the rises in asthma mortality cannot simply be related to increased incidence or prevalence of severe asthma requiring hospital admission. Assuming that data on asthma mortality are satisfactorily reliable, and that the determinants of hospital admission have not dramatically changed, this would suggest that



Trends in age standardised (on the European standard population) hospital admission rates for asthma in Lombardv and in mortality rates in Italy, 1976-1986.

Trends in age specific hospital admission rates from asthma in Lombardy, Italy, 1976-1986.

Calender period	Hospital admissions/1000 population									
	0-4 Males	years Females	5–1 Males	4 years Females	15⊸ Males	44 years Females	45– Males	64 years Females	> 7. Males	l years Females
1976-78	3.6	2.4	0.6	0.4	0.3	0.3	1.3	1.0	3.2	2.0
1979-81	3.7	2.6	0.8	0.4	0.3	0.3	1.0	1.0	2.6	1.8
1982-84	4 ·8	3.1	1.0	0.6	0.2	0.3	1.0	1.0	2.5	1.7
198586	5∙2	3.4	1.1	0.7	0.3	0.3	0.9	0.9	1.9	1.6
Percent change	+ 44	+ 42	+83	+ 75	_	_	- 31	- 10	-41	- 20

changes in the (pharmacological) management of the disease are also implicated in the recent unfavourable trends in mortality.^{4-7 10 11}

MONICA FASOLI CARLO LA VECCHIA Istituto di Ricerche Farmacologiche "Mario Negri", Via Eritrea 62, 20157 Milan

> MARINA FORMIGARO FRANCESCA REPETTO Assessorato alla Sanità Regione Lombardia, Via Stresa 24 20100 Milan, Italy

- Jackson RT, Beaglehole R, Rea HH, Sutherland DC. Mortality from asthma: a new epidemic in New Zealand. BM7 1982; 285: 771-4.
 Burney PGJ. Asthma mortality in England and Wales: evidence for a further increase, 1974-84. Lancet 1986; ii: 2020.
- 323-6
- 3 La Vechia C, Fasoli M, Negri E, Tognoni G. Fall and rise in asthma mortality in Italy, 1968–84. Int J Epidemiol 1989; 18: 998-9
- 4 Crane J, Pearce N, Flatt A, et al. Prescribed fenoterol and death from asthma in New Zealand, 1981–83: case-control study. Lancet 1989; i: 917–2.
- 5 Speizer FE, Doll R, Heaf P. Observations on recent increase
- in mortality from asthma. BMJ 1968; i: 335-9.
 Inman WH, Adelstein AM. Rise and fall of asthma mortality in England and Wales in relation to use of pressurised aerosols. Lancet 1969; ii: 279-85.
- 7 Keating G, Mitchell EA, Jackson R, Beaglehole R, Rea H. Trends in sales and drugs for asthma in New Zealand, Australia, and the United Kingdom, 1975-81. *BMJ* 1984; 289: 348-51.
- 8 Charlton JRH, Velez R. Some international comparisons of mortality amenable to medical intervention. *BMJ* 1986; 292: 295–301.
- 9 Robin ED. Death from bronchial asthma. Chest 1988; 93:
- 10 Lanes SF, Walker AM. Do pressurized bronchodilator aerosols cause death among asthmatics? Am J Epidemiol 1987; 125: 755-9.
 11 Burr ML. Is asthma increasing? J Epidemiol Community
- Burr ML. Is asthma increasing? J Epidemiol Community Health 1987; 41: 185–9.

Affluence, age, and motor neurone disease

Motor neurone disease (MND), of which sporadic adult onset amyotrophic lateral sclerosis (ALS), with or without bulbar involvement, is the most common clinical subtype, is a disease of unknown cause.¹ One possibility is the interaction of aging with, as yet, unidentified environmental factors.² Support for this suggestion is derived from the study of the Western Pacific forms of motor neurone disease,³ the low concordance observed in twin studies,⁴ and reports of motor neurone disease or MND like diseases in relation to a variety of exogenous factors.

Occupational groups or socioeconomic status might provide clues to possible environmental risks and it has been suggested that leather workers⁵ or agricultural workers⁶ may be at increased risk, but there is no consistently overrepresented group.⁷

Environmental factors may well be related not just with occupation but with social deprivation. Although there is some work relating motor neurone disease to social class, measurement of the latter is fraught with difficulty, particularly for women employed in the home. Large casecontrol studies have found no difference in socioeconomic status of men,⁸ or any relationship to home space.9 Two studies based on death certificates are conflicting; in Finland an increased risk in the lowest social class was noted

but with no clear gradient, while Martyn et al^{10} observed a stepwise trend of the proportional mortality ratio in England and Wales, which increased with higher social class.

To avoid some of these difficulties a measure of socioeconomic deprivation in Scotland has been devised¹¹ using information available from the last (1981) census which allocates a numerical score to individual postcode sectors (which contain an average of 6000 persons). Using the Z score technique to give a single score, the percentage values are combined for each post code sector based on the four variables of car ownership, degree of overcrowding, unemployment, and social class. The scores are distributed into seven categories within 0.3, 0.8, 1.5, and > 1.5 standard deviations (SD) either side of the mean (+0.3 SD to -0.3 SD as the middle)category). One important advantage of this method is that it can be applied to males and females, regardless of occupation, whether working or not. When standardised mortality ratios based on this scale are calculated, a number of common causes of death (eg, cancer of the lung) show a highly significant trend (figure, A), with higher ratios from deprived areas.

In contrast the figure (A) also shows the standardised mortality ratios based on the 533 deaths from motor neurone disease (ICD 335),



в MND by age group



Age standardised mortality ratios in Scotland by deprivation category (1 = affluent, 7 = deprived)

> 1980-85, male and female. This demonstrates the reverse trend, that is, higher standardised mortality ratios in individuals who live in areas of greater affluence. Confidence limits overlap with unity, and the χ^2 value for the Poisson regression is 2.21 (NS), but the graph does follow a "dose-response' relationship. Because of concern about accurate diagnosis of the cause of death in the elderly, particularly for a rare neurological disease, the total group has also been divided into those aged 20-74 years and aged >75 years (figure, B). It is then apparent that elderly people largely account for the overall trend in standardised mortality ratio with the χ^2 value for the regression being 3.2 (p<0.1) for those >74 years. This observation may reflect a failure to diagnose motor neurone disease in elderly patients from deprived areas which probably are not so well supplied with medical services as affluent areas. By inference perhaps the decline in incidence rates for motor neurone disease in the very elderly, as reported in all but one¹² of 40 population based mortality and incidence studies since 1950, is due to a similar diagnostic artefact. Alternatively, if environmental factors associated with affluence do predispose to motor neurone disease, the reason for the difference in the two age groups may be that the

- Williams DB, Windebank AJ. Motor neurone disease (amytrophic lateral sclerosis). Mayo Clin Proc 1991; 66: 54-82.
 Calne DB, Eisen A, McGeer E, Spencer P. Alzheimer's disease, Parkinson's disease and motorneurone disease: a biotropic interaction between ageing and environment? Lancet 1986; ii: 1067-70.
 Steele JC, Guaman T. Observations about amyotrophic lateral sclerosis and the parkinsonism-dementia complex of Guam with regard to epidemiology and etiology. Can J Neurol Sci 1987; 14: 358-62.
 Hawkes CH, Graham AJ. What causes motorneuron disease? [letter]. Lancet 1991; 337: 180.
 Buckley J, Warlow C, Smith P, Hilton-Jones D, Irvine S, Tew JR. Motor neuron disease in England and Wales 1959-1979. J Neurol Neurosurg Psychiatry 1983; 43: 197-205.
 Rosati G, Pinna L, Granieri E, et al. Studies on epidemiological, clinical, and etiological aspects of ALS disease in Sardinia, Southern Italy. Acta Neurol Scand 1977; 55: 231-44.
 Gawel M, Zaiwalla Z, Rose FC. Antecedent events in motor neurone disease. J Neurol Neurosurg Psychiatry 1983; 46: 1041-3.

- neurone 1041-3.





elderly have benefited less than younger people from improvements in the standard of living since the second world war. Clearly however our observation argues strongly against factors associated with deprivation increasing the risk for subsequent motor neurone disease and our prospective register of patients with the disease in Scotland will allow us to explore this issue further.¹³

> A M CHANCELLOR C P WARLOW Bramwell Dott Building Department of Clinical Neurosciences Western General Hospital Edinburgh EH4 2XU

V CARSTAIRS Health Services Research Network

> R A ELTON Medical Statistics Unit Edinburgh University Edinburgh

R J SWINGLER Department of Neurology Dundee Royal Infirmary

- 8 Kurtze JF, Beebe GW. Epidemiology of amyotrophic lateral
- Kurtze Jr., Beebe Gw. Epidemiology of amyotrophic lateral sclerosis: 1. A case-control comparison based on ALS deaths. Neurology 1980; 30: 453-62.
 Kondo K, Tsubaki T. Case-control studies of motor neurone disease. Association with mechanical injuries. Arch Neurol 1981; 38: 220-6.
 Martyn CN, Barker DJP, Osmond C. Motoneuron disease and past poliomyelitis in England and Wales. Lancet 1988; i: 1310-22

- and past poliomyentis in Eligibility and where a second second
- 13 Chancellor AM, Swingler RJ, Fraser H, Warlow CP. The Scottish motor neuron disease register: a prospective study of adult onset motor neuron disease in Scotland. of Methodology, demography and clinical features of incident cases in 1989. J Neurol Nurosurg Psychiatry (in press).