

Energy, protein, zinc and copper status of twenty-one elderly inpatients: analysed dietary intake and biochemical indices

BY ANITA J. THOMAS^{1,2}, VALDA W. BUNKER², LESLIE J. HINKS²,
NIDISH SODHA², MARK A. MULLEE³ AND BARBARA E. CLAYTON²

¹ Geriatric Medicine, ² Chemical Pathology and Human Metabolism, and ³ Medical Statistics and Computing, Medical Faculty of the University of Southampton, Level D, South Laboratory Block, Southampton General Hospital, Tremona Road Southampton SO9 4XY

(Received 1 September 1987 – Accepted 11 November 1987)

1. Duplicate diet analysis for energy, protein, zinc and copper with estimates of biochemical status for Zn and Cu were undertaken in twenty-one elderly long-stay inpatients (mean age 82 (range 63–89) years) consuming their customary hospital diet and in a stable medical condition. Fourteen patients had a long-standing and significant healing problem, either a leg ulcer or pressure sore.

2. Mean daily intakes of energy (5.2 MJ), protein (45 g), Zn (85 μmol) and Cu (14 μmol) were low in comparison with both official recommendations and levels of intake at which metabolic equilibrium was observed in healthy elderly people studied by the same methods (Bunker *et al.* 1984a).

3. Mean leucocyte Zn (9 pmol/ 10^6 cells) and Cu (7.5 pmol/ 10^6 cells) were low in comparison with results from healthy elderly people (Bunker *et al.* 1984a), implying suboptimal status for these elements. Those patients with healing problems tended to have the lower values within the range.

4. Recommendations are made with respect to improving nutritional status in this disadvantaged group of people.

Experts doubt the capacity of the Health Service to meet the ever increasing demands of an ageing population, many of whom are chronically ill and disabled (Grundy, 1984; Andrews, 1985). In 1984 over one-third of all general medical and surgical beds were occupied by patients over 65 years of age (Department of Health and Social Security (DHSS), 1986). The elderly are at risk of poor nutrition for a number of reasons including economic pressures, poor dentition, reduced mobility, depression, loneliness and ageing tissues. Adequate intake of elements required in trace amounts may be particularly threatened. Clinical signs of frank nutritional deficiency are unusual, but fractured bones, pressure sores and leg ulcers are common and produce healing problems which are costly in both human and economic terms. The number of hospital admissions for fractured neck of femur alone (classically a disease of elderly women) has more than doubled in the last 20 years (Fenton Lewis, 1981). Elements other than calcium and phosphorus may be important in the pathogenesis of osteoporosis, as suggested by the finding that higher levels of intake in women are related to lower rates of loss of bone-mineral content as measured by single photon absorptiometry (Freudenheim *et al.* 1986). Zinc and copper have an important role in both healing and immunity, and Cu is vital to the integrity of connective tissues (Chandra, 1980; Solomons, 1985). There is no specific recommendation for the intake of trace elements in the elderly, either in the UK or the USA, and the only guidelines available are derived from studies of younger individuals (DHSS, 1979a; Food and Nutrition Board, 1980).

Our previous work on the trace-element content of hospital meals revealed an intake of Zn and Cu approximately half that recommended for younger individuals even if elderly patients were assumed to eat everything selected by them or on their behalf (Thomas *et al.* 1986). The present study reports findings of actual dietary intake by analysis from duplicate meals and uneaten food over a 5 d period for a group of elderly inpatients.

EXPERIMENTAL

Subjects

Twenty-one inpatients of a geriatric unit devoted to the care of patients whose rehabilitation is expected to take longer than 3 months (slow-stream rehabilitation) and those whose discharge from hospital is unlikely (long stay) (Hall, 1974) took part. They included seventeen women and four men with an overall mean age of 81.7 years (range 63–89 years). Fourteen patients had a long-standing (> 6 months) and significant healing problem, either a pressure sore or leg ulcer. None of them smoked. Eighteen were edentulous, and eleven reported problems with ill-fitting dentures. The patients had been resident on the ward and had consumed hospital food for more than 3 months. They were weighed weekly as part of normal ward routine, on the same scales. Patients were not included if they had gastrointestinal, hepatic, or renal disease or had been receiving vitamin and mineral supplements. None of them were taking prostaglandin-synthetase inhibitors or chelating agents and note was taken of all other medication including thiazide diuretics. The 'intervention effect' was minimal as the collections were supervised by investigators who were familiar to staff and patients as they had worked on the wards for over 2 years in other capacities.

The present investigation was approved by the Joint Ethical Subcommittee of the Southampton and South West Hampshire Health Authority and the Faculty of Medicine of the University of Southampton.

MATERIALS AND METHODS

Diet collection. Each item of oral intake was exactly reproduced, including hospital meals and snacks, sweets, medicines, and any food brought by visitors. One of the duplicates was presented to the patient for consumption and the other reserved for analysis. Any portions remaining after the patients had finished were collected as 'rejects' for analysis. Care was taken at all stages to avoid contamination. Food was collected into trace-metal-free, acid-washed white plastic containers.

Blood collection. Venous blood was collected with minimum stasis from nineteen patients between 09.00 and 10.00 hours. Separation of leucocytes was undertaken immediately on 15 ml of the blood taken into an acid-washed siliconized glass tube containing heparin. Plasma Zn, Cu, caeruloplasmin, retinol-binding protein (RBP) and albumin concentrations were determined on blood taken into trace-metal-free tubes containing heparin, and the leucocyte count was estimated on samples taken into trace-metal-free tubes containing EDTA.

Analyses

Duplicated intakes and rejects for a 5 d period were homogenized as previously described (Bunker *et al.* 1984a) and stored at -20° before analysis. All samples were analysed in triplicate. Table 1 gives details of all methods used in the present study.

National Bureau of Standards bovine liver was used as reference material and mean values (n 6) obtained for Zn were $1.95 \mu\text{mol}$ dry weight (assigned value 1.88 (estimated degree of uncertainty (EDU) 0.12)) and for Cu $2.44 \mu\text{mol/g}$ dry weight (assigned value 2.49 (EDU 0.11)). Within-batch coefficients of variation for Zn and Cu analyses in diet were 3.8 and 4.0% respectively, and between-batch variation was 4.4 and 4.5% respectively.

Metabolizable energy (ME) was derived from the formula of Miller & Payne (1959) converted to SI units:

$$\text{ME (kJ)} = (\text{gross energy (kJ)} \times 0.95) - (\text{nitrogen (mmol)} \times 0.0044).$$

Table 1. Details of the methods used in the study

Material	Constituent	Method
Food	Gross energy Protein	Bomb calorimetry* Calculated from nitrogen determined by semi-automated Kjeldahl (Tecator, Bristol)
Plasma	Zinc and copper	Dry ashing, FAAS†
	Caeruloplasmin Retinol-binding protein } Alkaline phosphatase (EC 3.1.3.1)	Radial immunodiffusion (Hoechst, UK) SP 120 Analyser (Vickers, Basingstoke) using 4-nitrophenyl disodium orthophosphate substrate
	Albumin	SP 120 Analyser (Vickers), bromo- cresol green dye-binding
	Zn and Cu	FAAS‡
Leucocyte	Zn and Cu	Dextran sedimentation§, followed by FAAS (Zn) and ETA AAS (Cu)

Coefficients of variation for within-batch precision of all these analyses were less than 5%.

FAAS, flame atomic absorption spectrometry; ETA AAS, electrothermal atomization atomic absorption spectrophotometry.

* Miller & Payne (1959).

† Bunker *et al.* (1984*b*).

‡ Meret & Henkin (1971).

§ Hinks *et al.* (1982).

Data were assessed for normality of distribution and where positively-skewed, log transformation was undertaken and geometric means and confidence intervals calculated from transformed data. Where transformation was necessary, this is indicated in the Tables. Means, 95% confidence intervals, correlation coefficients and unpaired *t* tests, were calculated using the Minitab package (Minitab Inc., 1985) on an IBM 3090/150 mainframe computer. A probability value of $P < 0.05$ was accepted as significant.

RESULTS

The results for the four men did not differ greatly from those for the women and as the former were too small a number to be considered separately, the twenty-one subjects were considered as a whole.

Daily body-weights were stable over a 12 month period in eight patients (± 2 kg), but weight losses of 2–4 kg were observed in ten patients and weight losses of 5–6 kg in three patients. The meals served to the twenty-one patients offered a daily mean ME content of 6.23 MJ, protein content of 57 g, and Zn and Cu contents of 110 and 18 μmol respectively if completely consumed. Table 2 presents the means and 95% confidence intervals, for the actual intakes of ME, protein, Zn and Cu expressed as amount per day, per kg body-weight per day, and per 10 MJ energy. Table 3 presents the means and 95% confidence intervals for the biochemical indices of Zn and Cu status. Of the mean total ME intake 15% was derived from protein. There was a significant correlation between ME intake and protein intake ($r 0.63$, $P = 0.003$). This relation was strengthened by taking account of body-weight ($r 0.89$, $P < 0.001$). Daily ME intake and body-weight were not related ($r 0.22$).

Table 2. Nutrient intake in twenty-one geriatric inpatients by duplicate meal analysis over a 5 d period

Nutrient	Units/d	Recommended intake*	Mean intake	95% Confidence interval	Mean intake of apparently healthy elderly†
Energy	MJ ME	10 (9) ♂ 8 (7) ♀	5.16	4.66–5.66	7.58
	kJ/kg body-wt	—	89.7	75.7–103.6	114
Protein	g	60 (54) ♂ 47 (42) ♀	44.6	39.7–49.6	64.1
	g/kg body-wt	—	0.79	0.66–0.92	0.97
	g/10 MJ ME	—	87.2	80.3–94.1	86.0
Zinc	μmol	229	85.1	75.7–94.5	137
	μmol/kg body-wt	—	1.47	1.24–1.70	2.09
	μmol/10 MJ ME	—	167	151–182	177
Copper	μmol	31.5–47.2	13.2‡ (14.1)	11.2–15.8	18.7‡ (20.1)
	mmol/kg body-wt	—	0.22‡ (0.24)	0.18–0.27	0.28‡ (0.32)
	μmol/10 MJ ME	—	26.2‡ (27.7)	22.4–30.8	24.1‡ (26.6)

ME, metabolizable energy.

* Department of Health and Social Security (1979*a*) recommended daily intakes of energy and protein, values for 65–74-year-olds, with values for 75-year-olds in parentheses; Food and Nutrition Board (1980) recommended daily dietary allowance for Zn, and estimated safe and adequate dietary intake for Cu for 51+ years.

† Energy and protein, Bunker *et al.* (1987*b*); Zn and Cu, Bunker *et al.* (1984*a*).

‡ Values are geometric means with arithmetic means in parentheses.

Table 3. Biochemical indices of zinc and copper status in twenty geriatric inpatients

(Mean values and 95% confidence intervals)

	Geriatric inpatients		Healthy elderly controls†	
	Mean	95% Confidence interval	Mean	95% Confidence interval
Plasma Zn (μmol/l)	10.7	10.0–11.5	11.0	10.5–11.5
Leucocyte Zn (pmol/10 ⁶ cells)	91	81–101	120	110–130
Plasma Cu (μmol/l)	25.5	23.2–27.8	19.4	18.0–20.8
Leucocyte Cu (pmol/10 ⁶ cells)	7.5	6.1–8.9	11.5	10.0–12.9
Plasma:				
Retinol-binding protein (mg/l)	39	34–47	46	42–50
Caeruloplasmin (mg/l)	408	361–455	330	297–363
Alkaline phosphatase (EC 3.1.3.1) (i.u./l)	244* (263)	200–297	194	174–214
Albumin (g/l)	35	33–37	45	44–46

* Geometric mean with arithmetic mean in parentheses.

† Bunker *et al.* (1984*a*).

There was a close relation between Zn and protein intakes (r 0.79, P < 0.001). Again, the relation was strengthened by allowing for body-weight (r 0.95, P < 0.001). Similarly, intakes of Cu and protein were significantly correlated (r 0.47, P = 0.05) and taking account of body-weight slightly strengthened the relation (r 0.50, P = 0.03). The correlation

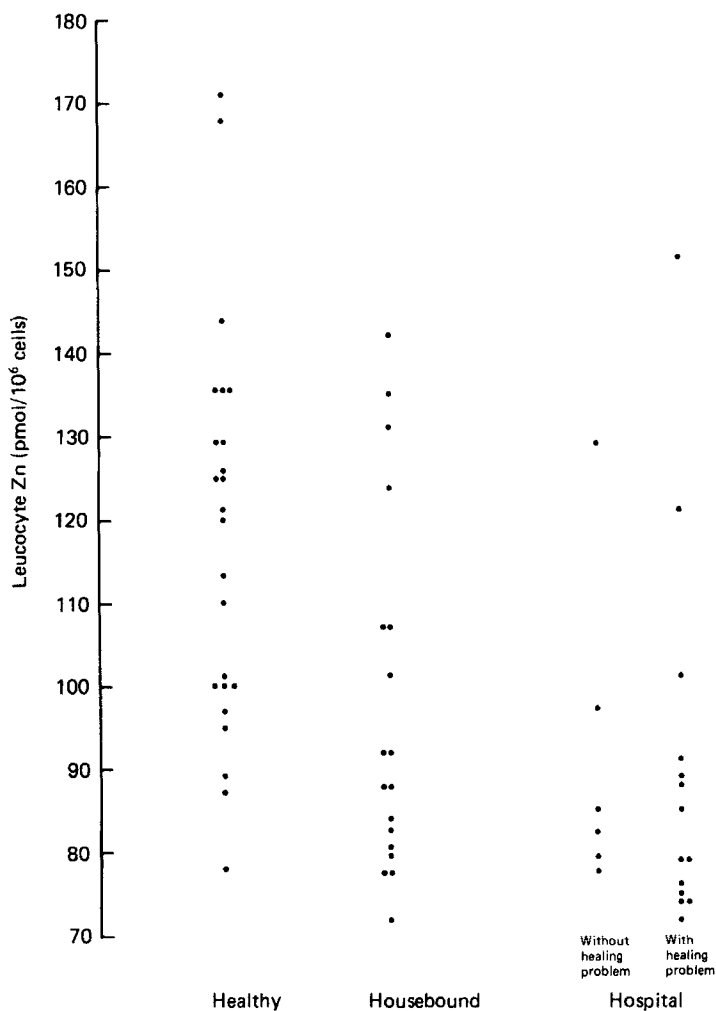


Fig. 1. Leucocyte zinc levels in healthy (Bunker *et al.* 1984*a*), housebound (Bunker *et al.* 1987*b*) and hospitalized old people (present study).

between Zn and Cu intakes was only significant when expressed in terms of body-weight (r 0.50, P = 0.03).

There was no significant correlation between age and intakes of ME, protein, Zn or Cu. Neither was any significant relation demonstrated between intakes of Zn and Cu and any of the biochemical indices of Zn and Cu status measured.

There was no significant difference between patients with or without healing problems in relation to intakes of ME, protein, Zn and Cu. Neither were there any significant differences between these two groups in terms of plasma Zn, leucocyte Zn, RBP, plasma albumin, alkaline phosphatase (*EC* 3.1.3.1), plasma Cu or caeruloplasmin. However, those patients with healing problems had leucocyte Zn levels that clustered at the lower end of the range, as illustrated in Fig. 1. Fig. 1 includes results for healthy (Bunker *et al.* 1984*a*) and housebound (Bunker *et al.* 1987*b*) elderly subjects for comparison.

DISCUSSION

The proper conduct of the duplicate-diet technique and trace-element analysis demand meticulous methodology in collection and preparation of samples, and high analytical standards. Such studies are time-consuming and labour intensive. The number of patients in our study is necessarily small but to the best of our knowledge it is the only study in the UK to date using analysed, rather than calculated, values in this population. Values calculated from food tables can be misleading (Lawson *et al.* 1982).

We have demonstrated an ME intake of less than two-thirds the recommended level (DHSS, 1979*a*). Protein intake was marginally lower than that advised, but Zn and Cu intakes were only 40% of the available recommendation (Food and Nutrition Board, 1980). Recently, in a published abstract, low intakes for Zn and Cu were recorded in a similar population in Edinburgh but these values were calculated from food tables (Henery & Smith, 1987).

Recommended allowances are intended as guidelines for an adequate intake and include a safety margin. Informed application of these ideal values is of great importance. Schneider *et al.* (1986) have drawn attention to the lack of specific recommendations for the elderly and the many factors which will complicate the task, and there are as yet no guidelines regarding trace element intake in this group.

The present results are best compared with metabolic balance information on healthy and housebound elderly people studied in their own homes in Southampton using the same methodology (Bunker *et al.* 1984*a*, 1987*b*). The mean daily ME intake of 5.2 MJ in the hospitalized elderly was 30% lower than that of the healthy elderly (7.2 MJ). We are unable to compare energy expenditure in these two groups. The healthy elderly may be thought to be more physically active on first impressions, but considerable energy may be expended in the achievement of very limited mobility by a disabled person (Isakov *et al.* 1985).

The mean daily protein intake of 45 g for the hospital population represents three-quarters that of the healthy elderly (64 g) (Bunker *et al.* 1987*b*). This relatively small difference may be important when tissue repair and muscular development are significant anabolic functions. The percentage of energy derived from protein was identical in both groups (15%).

The hospitalized elderly had a mean daily Zn intake of less than two-thirds that of the healthy elderly (85 μmol compared with 137 μmol) (Bunker *et al.* 1984*a*). The situation was similar for Cu, mean daily intakes being 13 and 20 μmol for the hospital and healthy populations respectively.

ME, protein, Zn and Cu intakes, whether expressed per day, per kg body-weight per day, or per 10 MJ are comparable in the housebound (Bunker *et al.* 1987*a*, *b*) and hospitalized groups. Indeed, the housebound elderly of Bunker *et al.* (1987*a*, *b*) and the present hospitalized subjects were similar in terms of ill-health and dependency and distinguishable primarily by their differing environments, the disabilities of the former being supported in the community, the latter in hospital. It has been well-recognized since Sheldon's (1948) original observations on the elderly in Wolverhampton that great disability and dependence can be managed at home.

The results of eight reports concerned with the intakes of ME and protein in elderly subjects in the UK are summarized in Table 4. The low mean ME intake of our subjects is similar to findings in other studies of the elderly at home, in sheltered housing and in hospitals in Britain using dietary recall or weighed inventory. When comparing these values with the recommended daily amounts (DHSS, 1979*a*) we should bear in mind that studies using calorimetry have suggested that we are overestimating energy requirements (Daly *et al.* 1985) and that adaptation may occur at low levels of energy intake (Waterlow, 1986).

Nevertheless, low ME intakes have been linked to high mortality from stroke (Lapidus *et al.* 1986).

Protein intakes were probably adequate in our study, at a level commensurate with the World Health Organization's recommendation for the elderly of not less than 0.75 g good-quality protein/kg body-weight per d (World Health Organization, 1985), although overall mean intakes were somewhat lower than others have found (Table 4). Protein synthesis and degradation are linked with both protein intake *per se*, and non-protein energy intake acting as summative stimulating factors for these processes (Reeds, 1983). However, this linkage is not inextricable and the effects of chronic subnutrition and illness are ill-defined. Indeed, protein utilization may be less efficient in the elderly than in the young (World Health Organization, 1985).

The apparently healthy elderly people studied by Bunker *et al.* (1984*a*) were in metabolic balance for Zn and Cu and displayed no clinical or biochemical signs of deficiency despite an intake of Zn and Cu which was 60% of that recommended for younger people (Food and Nutrition Board, 1980). Analysed intakes for these essential trace elements in this group of ill elderly people in hospital were well below half the recommended daily amounts (DHSS, 1979*a*; Food and Nutrition Board, 1980) at a time when good nutrition was of paramount importance. This situation obtains in a Health District where the hospital kitchens have won awards for excellence and every care is taken by the staff to cater for individual preference. The nutrient density offered by the hospital diet was similar to that found for the diet eaten by apparently healthy elderly people in the community (Bunker *et al.* 1984*a*). This suggests a reduced intake of food by patients rather than the provision of an inadequate diet.

In contrast to the healthy elderly, although displaying no clinical evidence of deficiency (and chronic marginal deficiency may present quite differently to an acute severe deficit) the inpatients had low levels of leucocyte Zn. Most of the evidence linking improvement of healing with Zn supplementation is circumstantial (Pories *et al.* 1967; Husain, 1969; Haeger & Lanner, 1974). Many studies on healing and Zn supplementation failed to make any assessment of Zn status, and those that did relied on plasma Zn levels which are affected by many non-specific factors including stress and infection (Delves, 1985). The value of leucocyte Zn as an index of tissue stores has been well-reviewed elsewhere (Patrick & Dervish, 1984; Delves, 1985). Our own patients had a lower range of leucocyte Zn levels than the group of healthy elderly (Bunker *et al.*, 1984*a*) and the leucocyte Zn levels of those with healing problems clustered at the lower end of this range. Leucocyte Cu levels were lower in these elderly inpatients than in the apparently healthy elderly (Bunker *et al.* 1984*a*), perhaps suggesting tissue deficit and suboptimal body Cu status. Low levels of leucocyte Zn have also been found in another study of long-stay geriatric patients (Stafford *et al.* 1987).

Values for plasma RBP concentrations were similar in housebound and hospitalized people and lower than those in healthy elderly in Southampton. Plasma albumin levels were lower in the hospital group, but patients were recumbent for venepuncture and this may have affected the value obtained (Freeman & Cox, 1984). Plasma Zn levels were comparable in all three groups and leucocyte Zn showed a downwards trend through the three groups, from healthy to housebound to hospitalized. Plasma Cu and caeruloplasmin levels were similar in the housebound and hospitalized patients and higher than those in the healthy. This was not unexpected as most Cu in plasma is bound to caeruloplasmin which is an acute-phase protein.

The low mean intake of Zn and low concentrations of leucocyte Zn suggest a state of chronic marginal deficiency. Foods rich in protein are usually rich in Zn (Osis *et al.* 1972). Although protein intakes were just adequate in our group, Zn intakes were suboptimal.

Table 4. Energy and protein intake in elderly women: a review of some studies in the UK
(Mean values and standard deviations)

Location of study	Reference	Method	No. of subjects	Age (years)	Residence of subjects	Total metabolizable energy intake (MJ/d)		Protein intake (g/d)	
						Mean	SD	Mean	SD
Multicentre	Department of Health and Social Security (1979 <i>b</i>)	7 d weighed record	196	65+	Community	6.9	1.7	55.7	13.3
Edinburgh	Loneragan <i>et al.</i> (1975)	7 d record	73	62-90	Community	6.9	1.4	54.4	12.3
Glasgow	Macleod <i>et al.</i> (1974)	7 d record	187	65+	Community	7.3	1.8	60	13
Belfast	Vir & Love (1979)	3 d weighed record	27	65-95	Community	7.4	2.3	51.9	15.5
Southampton	Bunker <i>et al.</i> (1984 <i>a</i>)	5 d duplicate-diet analysis	13	70-86	Healthy in community	6.6	1.2	59.7	13.8
Southampton	Bunker <i>et al.</i> (1987 <i>a</i>)	5 d duplicate-diet analysis	13	71-83	Housebound in community	4.8	0.9	39.1	7.0
Belfast	Vir & Love (1979)	3 d weighed record	30	65-95	Long-stay hospital	6.2	0.9	50.5	9.2
Southampton	MacLellan <i>et al.</i> (1975)	7 d record	66	65+	Ashurst long-stay hospital	5.7	1.5	—	—
Southampton	Present study	5 d duplicate-diet analysis	21	72-89	Moorgreen long-stay hospital	5.3	1.1	42.8	10.4
	RDA for women > 75 years	—	—	—	—	7	—	42	—

RDA, recommended daily amount (Department of Health and Social Security, 1979*a*).

Some hospitals use textured vegetable protein in the form of 'extended mince', but the hospital in which the present study was conducted does not. The foods from which the hospital meals were prepared were fresh or frozen, so losses during processing or preservation were unlikely (Shroeder, 1971).

It is a common observation that old people prefer food that requires little chewing and bland rather than strong flavours. This may explain the apparent incongruity between the Zn and protein intakes in our patients. Many otherwise excellent studies of nutrition in the elderly fail to assess the state of dentition which is of obvious importance in the selection and ingestion of food. Smith (1977), in her study of the oral health and dental status of 254 elderly people living at home, found that almost one-third had difficulty chewing and more than one in ten had needed to change the type of food eaten, or method of cooking, to enable them to chew effectively. Over three-quarters of our patients were edentulous and half reported problems with dentures that interfered with eating. The nutritional effects of tooth loss have been well-reviewed by Geissler & Bates (1984). Protein requirements may be met by milk, eggs, cheese and perhaps the white meats that are palatable to the elderly and prepared as dishes that are easy to eat. Coincidentally these foods contain less Zn per g protein than red meat (Murphy *et al.* 1975). Indeed, roast beef has been removed from the menu in the hospital studied as so much was wasted, although minced ham and lamb are still presented. It may be, therefore, that at a critical level of intake self-selection results in lower-than-adequate daily Zn intake, but sufficient protein.

The present vogue for adding bran to many dishes, such as porridge, soups and pies (as occurs in this hospital) may be counterproductive as it is well-known that phytate may decrease the amount of Zn and Cu available for absorption in the intestinal lumen (Rheinhold *et al.* 1976; Davies & Olpin, 1979; Turnlund *et al.* 1984). Similarly, the higher than recommended daily intake of Ca and P in this group, revealed by our own unpublished results, may render more of the dietary Zn unavailable (Sandstead, 1982). An increase in the level of dietary protein results in increased dietary provision of Zn, by virtue of its intrinsic content and also increased retention of Zn (Greger & Snedeker, 1980).

We have studied a small number of elderly people and find that, for Zn and Cu, institutional malnutrition (Bender, 1984) is a real possibility. Recommendations on the advisability of increasing the ME intakes cannot be made on the information presented here, but these low levels of intake should not be accepted with equanimity merely on the basis of apparent reduction in physical activity, especially with evidence of weight loss in over half the patients studied. Intakes of Zn and Cu should be improved commensurate with those found in healthy elderly people in metabolic balance for these elements. Measures that may reduce the availability of trace elements, such as the addition of extra fibre, should not be implemented before careful consideration. Nutritional intake generally could be improved if patients were to eat in small groups, more attention were paid to oral health, and foods were presented so that it could be eaten without utensils if necessary. Strategies that may improve Zn and Cu intakes without resort to supplementation include increasing the proportion of Zn-rich protein sources in the diet, as well as increasing the protein density of the diet. If supplementation is considered this should be at physiological rather than pharmacological levels as side-effects of Zn supplementation have been reported (Moore, 1978).

The authors are very grateful to the patients, nursing staff and catering staff at Moorgreen Hospital for all their help during this study. Professors M. R. P. Halls and R. S. J. Briggs and Dr I. R. Reid kindly gave permission for patients under their care to be studied. The authors would particularly like to thank Professor Hall for his help and interest. The study was supported financially by the BMA Charlotte Eyck Award (held by A. J. T.), the

Foundation for Age Research, The Wessex Medical School Trust, and Associated Octol Ltd. They are also indebted to Dr M. Campbell of Community Medicine, Medical Statistics and Computing, Faculty of Medicine, University of Southampton for statistical advice.

REFERENCES

- Andrews, K. (1985). *British Medical Journal* **290**, 1023–1024.
- Bender, A. E. (1984). *British Medical Journal* **288**, 92–93.
- Bunker, V. W., Hinks, L. J., Lawson, M. S. & Clayton, B. E. (1984a). *American Journal of Clinical Nutrition* **40**, 1096–1102.
- Bunker, V. W., Hinks, L. J., Stansfield, M. F., Lawson, M. S., Clayton, B. E. (1987a). *American Journal of Clinical Nutrition* **46**, 353–359.
- Bunker, V. W., Lawson, M. S., Delves, H. T. & Clayton, B. E. (1984b). *American Journal of Clinical Nutrition* **39**, 797–802.
- Bunker, V. W., Lawson, M. S., Stansfield, M. F., Clayton, B. E. (1987b). *British Journal of Nutrition* **57**, 211–221.
- Chandra, R. K. (1980). *American Journal of Clinical Nutrition* **33**, 736–738.
- Daly, J. M., Heymsfield, S. B., Head, C. A., Harvey, L. P., Nixon, D. W., Katzef, H. & Grossman, G. D. (1985). *American Journal of Clinical Nutrition* **42**, 1170–1174.
- Davies, N. T. & Olpin, S. E. (1979). *British Journal of Nutrition* **41**, 590–603.
- Delves, H. T. (1985). *Clinical Endocrinology and Metabolism* **14**, 725–760.
- Department of Health and Social Security (1979a). *Recommended Daily Amounts of Food Energy and Nutrients for Groups of People in the United Kingdom. Report on Health and Social Subjects no. 15*. London: H.M.S.O.
- Department of Health and Social Security (1979b). *Nutrition and Health in Old Age. Report on Health Subjects no. 16*. London: H.M.S.O.
- Department of Health and Social Security (1986). *Hospital Inpatient Enquiry 1984. Office of Population Censuses and Surveys Series MB4 no. 24*. London: H.M.S.O.
- Fenton Lewis, A. (1981). *British Medical Journal* **283**, 1217–1219.
- Food and Nutrition Board (1980). *Recommended Dietary Allowances*, 9th revised ed. Washington DC: National Academy of Sciences.
- Freeman, H. & Cox, M. C. (1984). In *Clinical Biochemistry of the Elderly*, pp. 48–74 [H. M. Hodkinson, editor]. Edinburgh: Churchill Livingstone.
- Freudenheim, J. L., Johnson, N. E. & Smith, E. L. (1986). *American Journal of Clinical Nutrition* **44**, 863–876.
- Geissler, C. A. & Bates, J. F. (1984). *American Journal of Clinical Nutrition* **39**, 478–489.
- Greger, J. L. & Snedeker, S. M. (1980). *Journal of Nutrition* **110**, 2243–2253.
- Grundy, E. (1984). *British Medical Journal* **288**, 663–664.
- Haeger, K. & Lanner, E. (1974). *VASA Journal for Vascular Diseases* **3**, 77–81.
- Hall, M. R. P. (1974). *Medicine* **25**, 1465–1480.
- Henery, E. C. & Smith, R. G. (1987). *Proceedings of the Nutrition Society* **46**, 63A.
- Hinks, L. J., Colmsee, M. & Delves, H. T. (1982). *Analyst* **107**, 815–823.
- Husain, L. (1969). *Lancet* **i**, 1069–1071.
- Isakov, E., Susak, Z. & Becker, E. (1985). *Scandinavian Journal of Rehabilitation and Medical Suppl.* **12**, 108–111.
- Lapidus, L., Anderson, H., Bengtsson, C. & Bosaeus, I. (1986). *American Journal of Clinical Nutrition* **44**, 444–448.
- Lawson, M., Bunker, V., Delves, H. & Clayton, B. E. (1982). *Proceedings of the Nutrition Society* **41**, 91A.
- Lonergan, M. E., Milne, J. S., Maule, M. M. & Williamson, J. (1975). *British Journal of Nutrition* **34**, 517–527.
- MacLellan, W. J., Martin, P. & Mason, B. J. (1975). *Gerontologica Clinica* **17**, 173–180.
- MacLeod, C. C., Judge, T. G. & Caird, F. I. (1974). *Age and Ageing* **3**, 158–166.
- Meret, S. & Henkin, R. I. (1971). *Clinical Chemistry* **17**, 369–376.
- Miller, D. S. & Payne, P. R. (1959). *British Journal of Nutrition* **13**, 501–508.
- Minitab Inc. (1985). *Minitab Reference Manual*. Birmingham: CLE. COM.
- Moore, R. (1978). *British Medical Journal* **i**, 754.
- Murphy, E. W., Wells Willis, B. & Watt, B. (1975). *Journal of the American Dietetic Association* **66**, 345–355.
- Osis, D., Kramer, L., Wiatrowski, E. & Spencer, H. (1972). *American Journal of Clinical Nutrition* **25**, 582–588.
- Patrick, J. & Dervish, C. (1984). *CRC Critical Reviews in Clinical Laboratory Sciences* **20**, 95–114.
- Pories, W. J., Henzel, J. H., Rob, C. G. & Strain, W. H. (1967). *Lancet* **i**, 121–124.
- Reeds, P. J. (1983). *Proceedings of the Nutrition Society* **43**, 463–471.
- Rheinhold, J. G., Faradj, B., Abadi, P. & Ismail-Beigi, F. (1976). *Journal of Nutrition* **106**, 493–503.

- Sandstead, H. H. (1982). In *Clinical, Biochemical and Nutritional Aspects of Trace Elements* pp. 83–101 [A. Prasad, editor]. New York: Alan Liss.
- Schneider, E. L., Vining, E. M., Hadley, E. C. & Farnham, S. A. (1986). *New England Journal of Medicine* **314**, 157–160.
- Sheldon, J. H. (1948). *The Social Medicine of Old Age*. London: Nuffield Foundation, Oxford University Press.
- Shroeder, H. (1971). *American Journal of Clinical Nutrition* **24**, 562–573.
- Smith, J. M. (1977). A study of the oral handicaps, dental status and dental needs of an elderly population living at home. Dissertation submitted for Master of Medical Science in Community Medicine, University of Nottingham.
- Solomons, N. (1985). *Journal of the American College of Nutrition* **4**, 83–105.
- Stafford, W., Smith, R. G., Henery, E. C., Lewis, S. & O'Rorke, K. (1987). *Proceedings of the Nutrition Society* **46**, 62A.
- Thomas, A. J., Bunker, V. W., Brennan, E. & Clayton, B. E. (1986). *Human Nutrition: Applied Nutrition* **40A**, 440–446.
- Turnlund, J., King, J., Keyes, W., Gong, B. & Michel, M. (1984). *American Journal of Clinical Nutrition* **40**, 1071–1077.
- Vir, S. C. & Love, A. M. G. (1979). *American Journal of Clinical Nutrition* **32**, 1934–1947.
- Waterlow, J. C. (1986). *Annual Review of Nutrition* **6**, 495–526.
- World Health Organization (1985). *Energy and Protein Requirements. Technical Report Series no. 724*. Geneva: WHO.