

A population-based case–control study of diet and melanoma risk in northern Italy

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Submitted 20 October 2004; Accepted 10 May 2005

Abstract

Objective: We aimed at examining the association between dietary constituents and risk of cutaneous melanoma.

Design: In an area of northern Italy we recruited 59 newly diagnosed melanoma patients and 59 age- and sex-matched population controls, to whom we administered a validated semi-quantitative food-frequency questionnaire.

Results: We found an excess risk of melanoma in subjects with a higher energy-adjusted intake of total polyunsaturated fatty acids and, in particular, of linoleic acid (relative risk = 2.16 for intake in the highest tertile compared with the lowest tertile, *P* for linear trend = 0.061). Conversely, disease risk was inversely associated with the consumption of soluble carbohydrates (relative risk = 0.34 for intake in the upper vs. the lowest tertile adjusting for total energy intake, *P* for linear trend = 0.046). No other dietary factors, including alcohol, vitamins and trace elements, correlated with melanoma risk. The association of melanoma risk with linoleic acid and soluble carbohydrates intakes was further strengthened in multivariate analysis, and when analysis was limited to females.

Conclusions: Overall, these results indicate that an excess energy-adjusted intake of linoleic acid and a lower consumption of soluble carbohydrates may increase melanoma risk.

Keywords

Diet
Linoleic acid
Melanoma
Soluble carbohydrates
Case–control study

A recent report from England on trends in cancer occurrence between 1991 and 2000 classifies melanoma as having the highest increase in incidence among all site-specific neoplasms¹. The incidence of melanoma has increased generally in western European countries during the last few years, and early detection owing to screening does not appear as the only explanation for this fact². This points to the role of environmental risk factors in increasing melanoma incidence, but the exact nature of these factors, apart from ultraviolet exposure, is still unclear³. Diet has long been suspected of playing a role in favouring melanoma occurrence, and new interest in this issue has recently been raised by two large hospital-based case–control studies^{4,5}, although the results of the few investigations carried out so far have yielded inconsistent results^{3,6}. To further explore the possible relationship between dietary factors and melanoma risk, we carried out a population-based case–control study in an Italian region.

Methods

We carried out a population-based case–control study to investigate the possible relationship between

environmental factors and risk of cutaneous melanoma in the population from Modena Province, an area in northern Italy with approximately 650 000 inhabitants⁷. We recruited for a period of three consecutive years all the patients newly diagnosed with cutaneous melanoma attending the Dermatologic Clinic of Modena University Hospital, which is the only centre for diagnosis, therapy and follow-up of the disease within the province. Inclusion criteria were residence in Modena Province and a histologically based recent diagnosis of cutaneous melanoma without clinical evidence of metastasis. Fifty-nine out of the 82 patients (72.0%) eligible to participate in the study accepted to do so.

Subsequently, we randomly selected a resident of Modena Province, sex- and age-matched (± 5 years) to each patient. This was carried out by using the database for all Modena residents available at the Modena Local Health Authority. Eligible controls were contacted by telephone were enrolled in the study after having given their informed consent to the protocol and aims of the study. When a control subject could not be enrolled, another resident age- and sex-matched to the patient was contacted and invited to participate.

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Table 1 Crude and energy-adjusted relative risk of melanoma according to daily intake of dietary factors, case-control study on environmental determinants of cutaneous melanoma in Modena Province, northern Italy

Factor	Relative risk (95% confidence interval)			P for linear trend	
	I*	II	III		
Total proteins	Crude	1.00	1.33 (0.46–3.84)	0.50 (0.15–1.66)	0.514
	Energy-adjusted	1.00	0.54 (0.10–3.01)	0.83 (0.14–4.71)	0.979
	Tertile cut-off (g)	< 74.138	≥ 74.138–< 96.616	≥ 96.616	
Animal proteins	Crude	1.00	0.87 (0.32–2.41)	0.62 (0.20–1.91)	0.477
	Energy-adjusted	1.00	0.57 (0.17–1.9)	0.86 (0.21–3.55)	0.803
	Tertile cut-off (g)	< 49.383	≥ 49.383–< 70.429	≥ 70.429	
Vegetable proteins	Crude	1.00	1.83 (0.68–4.96)	0.60 (0.14–2.51)	0.793
	Energy-adjusted	1.00	1.60 (0.50–5.07)	0.89 (0.05–14.47)	0.549
	Tertile cut-off (g)	< 19.256	≥ 19.256–< 30.911	≥ 30.911	
Total fat	Crude	1.00	1.67 (0.61–4.59)	0.60 (0.14–2.51)	0.799
	Energy-adjusted	1.00	1.06 (0.18–6.20)	1.21 (0.09–15.28)	0.369
	Tertile cut-off (g)	< 68.722	≥ 68.722–< 89.503	≥ 89.503	
Animal fat	Crude	1.00	0.87 (0.32–2.41)	0.56 (0.19–1.66)	0.446
	Energy-adjusted	1.00	0.57 (0.17–1.89)	0.80 (0.17–3.72)	0.752
	Tertile cut-off (g)	< 39.099	≥ 39.099–< 54.773	≥ 54.773	
Vegetable fat	Crude	1.00	2.33 (0.60–9.02)	0.83 (0.25–2.73)	0.623
	Energy-adjusted	1.00	2.87 (0.63–13.18)	1.01 (0.20–5.04)	0.215
	Tertile cut-off (g)	< 24.586	≥ 24.586–< 37.775	≥ 37.775	
Total saturated fatty acids	Crude	1.00	1.25 (0.49–3.17)	0.57 (0.17–1.95)	0.488
	Energy-adjusted	1.00	0.66 (0.17–2.53)	0.92 (0.12–7.28)	0.890
	Tertile cut-off (g)	< 23.465	≥ 23.465–< 33.606	≥ 33.606	
Oleic acid	Crude	1.00	9.00 (1.14–71.04)	1.00 (0.25–3.40)	0.962
	Energy-adjusted	1.00	43.07 (1.96–944.69)	5.01 (0.44–57.13)	0.355
	Tertile cut-off (g)	< 27.095	≥ 27.095–< 40.143	≥ 40.143	
Total monounsaturated fatty acids	Crude	1.00	8.00 (1.00–63.96)	1.25 (0.34–4.65)	0.927
	Energy-adjusted	1.00	30.51 (1.45–643.90)	5.33 (0.56–50.73)	0.377
	Tertile cut-off (g)	< 29.338	≥ 29.338–< 42.587	≥ 42.587	
Linoleic acid	Crude	1.00	0.78 (0.29–2.09)	1.00 (0.32–3.10)	0.290
	Energy-adjusted	1.00	0.35 (0.08–1.54)	2.16 (0.40–11.72)	0.061
	Tertile cut-off (g)	< 6.976	≥ 6.976–< 8.811	≥ 8.811	
Linolenic acid	Crude	1.00	1.25 (0.49–3.17)	0.43 (0.11–1.66)	0.859
	Energy-adjusted	1.00	0.59 (0.16–2.24)	0.97 (0.14–6.53)	0.459
	Tertile cut-off (g)	< 1.001	≥ 1.001–< 1.296	≥ 1.296	
Other polyunsaturated fatty acids	Crude	1.00	4.33 (1.23–15.21)	0.67 (0.19–2.36)	0.288
	Energy-adjusted	1.00	7.13 (1.49–34.06)	0.73 (0.16–3.29)	0.408
	Tertile cut-off (g)	< 0.411	≥ 0.411–< 0.799	≥ 0.799	
Total polyunsaturated fatty acids	Crude	1.00	1.14 (0.41–3.15)	1.00 (0.32–3.10)	0.399
	Energy-adjusted	1.00	0.55 (0.13–2.33)	2.62 (0.45–15.35)	0.080
	Tertile cut-off (g)	< 8.407	≥ 8.407–< 10.830	≥ 10.830	
Cholesterol	Crude	1.00	0.29 (0.06–1.37)	0.50 (0.17–1.46)	0.210
	Energy-adjusted	1.00	0.18 (0.03–0.96)	0.50 (0.12–2.12)	0.252
	Tertile cut-off (mg)	< 289.751	≥ 289.751–< 402.895	≥ 402.895	
Available carbohydrates	Crude	1.00	1.80 (0.60–5.37)	0.50 (0.12–2.00)	0.346
	Energy-adjusted	1.00	1.82 (0.43–7.62)	0.42 (0.03–5.90)	0.468
	Tertile cut-off (g)	< 195.850	≥ 195.850–< 291.882	≥ 291.882	
Starch	Crude	1.00	1.67 (0.61–4.59)	0.83 (0.25–2.73)	0.962
	Energy-adjusted	1.00	1.55 (0.45–5.28)	6.76 (0.57–80.02)	0.367
	Tertile cut-off (g)	< 97.738	≥ 97.738–< 165.190	≥ 165.190	
Soluble carbohydrates	Crude	1.00	0.75 (0.26–2.16)	0.57 (0.17–1.95)	0.055
	Energy-adjusted	1.00	0.73 (0.23–2.27)	0.34 (0.07–1.73)	0.046
	Tertile cut-off (g)	< 86.256	≥ 86.256–< 114.814	≥ 114.814	
Dietary fibre	Crude	1.00	1.75 (0.51–5.98)	2.00 (0.50–8.00)	0.560
	Energy-adjusted	1.00	2.24 (0.55–9.07)	2.14 (0.31–14.77)	0.905
	Tertile cut-off (g)	< 14.457	≥ 14.457–< 20.276	≥ 20.276	
Alcohol	Crude	1.00	1.67 (0.61–4.59)	0.50 (0.12–2.00)	0.807
	Energy-adjusted	1.00	1.86 (0.64–5.42)	0.97 (0.17–5.50)	0.978
	Tertile cut-off (g)	< 1.625	≥ 1.625–< 23.248	≥ 23.248	
Energy	Crude	1.00	1.67 (0.61–4.59)	0.43 (0.11–1.66)	0.482
	Tertile cut-off (kcal)	< 1766.41	≥ 1766.41–< 2365.391	≥ 2365.391	
	Iron	Crude	1.00	0.67 (0.24–1.87)	0.83 (0.25–2.73)
Calcium	Energy-adjusted	1.00	0.16 (0.02–1.07)	1.45 (0.19–11.3)	0.548
	Tertile cut-off (mg)	< 11.666	≥ 11.666–< 14.934	≥ 14.934	
	Crude	1.00	1.60 (0.52–4.89)	1.00 (0.32–3.10)	0.384
	Energy-adjusted	1.00	1.89 (0.50–7.12)	0.81 (0.14–4.73)	0.604
	Tertile cut-off (mg)	< 757.927	≥ 754.927–< 1057.486	≥ 1057.486	

Table 1. Continued

Factor		Relative risk (95% confidence interval)			P for linear trend
		I*	II	III	
Sodium	Crude	1.00	0.78 (0.29–2.09)	1.20 (0.37–3.93)	0.979
	Energy-adjusted	1.00	0.63 (0.17–2.36)	5.88 (0.48–72.74)	0.335
	Tertile cut-off (mg)	< 1796.629	≥ 1796.629–< 2518.013	≥ 2518.013	
Potassium	Crude	1.00	1.14 (0.41–3.15)	1.50 (0.42–5.31)	0.628
	Energy-adjusted	1.00	1.52 (0.40–5.83)	3.43 (0.42–27.96)	0.873
	Tertile cut-off (mg)	< 2642.938	≥ 2642.938–< 3370.434	≥ 3370.434	
Phosphorus	Crude	1.00	1.00 (0.37–2.66)	0.40 (0.12–1.27)	0.325
	Energy-adjusted	1.00	1.19 (0.27–1.28)	0.42 (0.71–2.44)	0.392
	Tertile cut-off (mg)	< 1192.072	≥ 1192.072–< 1601.103	≥ 1601.103	
Zinc	Crude	1.00	1.80 (0.60–5.37)	0.67 (0.19–2.36)	0.491
	Energy-adjusted	1.00	1.60 (0.28–9.10)	1.57 (0.21–11.71)	0.895
	Tertile cut-off (mg)	< 9.797	≥ 9.797–< 13.560	≥ 13.560	
Thiamine	Crude	1.00	0.78 (0.29–2.09)	0.71 (0.23–2.25)	0.253
	Energy-adjusted	1.00	0.46 (0.12–1.73)	0.62 (0.11–3.57)	0.308
	Tertile cut-off (mg)	< 0.916	≥ 0.916–< 1.158	≥ 1.158	
Riboflavin	Crude	1.00	1.43 (0.54–3.75)	0.50 (0.12–2.00)	0.424
	Energy-adjusted	1.00	1.17 (0.34–4.01)	0.84 (0.13–5.37)	0.681
	Tertile cut-off (mg)	< 1.231	≥ 1.231–< 1.590	≥ 1.590	
Niacin	Crude	1.00	1.00 (0.35–2.85)	0.50 (0.15–1.66)	0.469
	Energy-adjusted	1.00	0.45 (0.10–1.95)	0.31 (0.05–1.93)	0.799
	Tertile cut-off (mg)	< 15.516	≥ 15.516–< 20.024	≥ 20.024	
Vitamin C	Crude	1.00	1.75 (1.51–5.98)	0.60 (0.14–2.51)	0.401
	Energy-adjusted	1.00	1.77 (0.51–6.15)	0.71 (0.13–4.03)	0.553
	Tertile cut-off (mg)	< 85.739	≥ 85.739–< 105.794	≥ 105.794	
Vitamin B ₆	Crude	1.00	0.37 (0.10–1.41)	1.33 (0.46–3.84)	0.590
	Energy-adjusted	1.00	0.26 (0.05–1.35)	1.82 (0.42–7.93)	0.930
	Tertile cut-off (mg)	< 1.545	≥ 1.545–< 1.950	≥ 1.950	
Folic acid	Crude	1.00	1.33 (0.30–5.96)	1.25 (0.34–4.65)	0.578
	Energy-adjusted	1.00	1.84 (0.27–12.29)	1.25 (0.18–8.60)	0.992
	Tertile cut-off (μg)	< 198.658	≥ 198.658–< 250.708	≥ 250.708	
Retinol equivalents	Crude	1.00	0.86 (0.29–2.55)	0.60 (0.14–2.51)	0.261
	Energy-adjusted	1.00	0.76 (0.21–2.81)	0.64 (0.14–2.95)	0.349
	Tertile cut-off (μg)	< 612.444	≥ 612.444–< 943.917	≥ 943.917	
Retinol	Crude	1.00	0.67 (0.24–1.87)	1.00 (0.25–4.00)	0.229
	Energy-adjusted	1.00	0.38 (0.10–1.48)	1.94 (0.33–11.54)	0.288
	Tertile cut-off (μg)	< 246.16	≥ 246.16–< 428.833	≥ 428.833	
β-Carotene	Crude	1.00	2.00 (0.60–6.64)	1.50 (0.42–5.31)	0.844
	Energy-adjusted	1.00	2.50 (0.67–9.28)	1.60 (0.42–6.12)	0.981
	Tertile cut-off (μg)	< 1747.094	≥ 1747.094–< 3012.109	≥ 3012.109	
Vitamin E	Crude	1.00	1.37 (0.55–3.42)	1.20 (0.37–3.93)	0.484
	Energy-adjusted	1.00	1.57 (0.53–4.59)	1.16 (0.26–5.07)	0.125
	Tertile cut-off (mg)	< 6.410	≥ 6.410–< 8.174	≥ 8.174	
Vitamin D	Crude	1.00	0.86 (0.29–2.55)	0.78 (0.29–2.09)	0.247
	Energy-adjusted	1.00	0.61 (0.17–2.10)	0.76 (0.23–2.50)	0.350
	Tertile cut-off (μg)	< 2.039	≥ 2.039–< 2.940	≥ 2.940	

*Lowest tertile.

The study protocol included a dermatological examination, administration of questionnaires on anamnestic data, diet and lifestyle habits, as well as the sampling of blood and toenails. Among the anamnestic data we collected, we paid particular attention to the overall history of sun exposure by asking information about estimated periods of the day and of the year spent in sunshine during the subject's lifetime, visits to tropical countries, history of any sunburn and of solar lentigines, and use of sunscreen. We also investigated the possible use of sunlamps and the phototype of the subject.

The dietary questionnaire was developed by epidemiologists from the Milan National Cancer Institute within the framework of the European Prospective Investigation on

Cancer and Nutrition (EPIC)⁸ and specifically designed for different Italian areas, such as Nord-Central Italy for the version adopted in the present study⁹. It was a self-administered, semi-quantitative food-frequency questionnaire divided into 14 sections: pasta and rice, soup, meat excluding cured meats, fish, raw vegetables, cooked vegetables, eggs, sandwiches, salami and other cured meat, cheese, fruit, bread and wine, milk/coffee/cakes, herbs and spices. The questionnaire included 248 questions on frequency (per day, month or year) of consumption of 188 food items. The quantity of each food consumed was assessed by selecting an image of a food portion using the 17 sets of pictures indicating small, medium and large portion sizes or slight modification of

these quantities. Images of glasses and cups of different sizes were also used to estimate wine and milk consumption, and the type of fats used as condiments for some foods or if added after cooking was assessed. Cooking procedures were taken into consideration.

After compilation and collection, the questionnaires were read by expert personnel and computerised using an optical reader and software developed at the National Cancer Institute. In this process possible inconsistencies between 'linked' replies were checked (for example, details on cooking methods for salmon were not acceptable if no consumption of salmon was acknowledged). The software allowed for the calculation of consumption frequency and daily quantities of each food item, as well as the intake of dietary factors using the *Italian Food Composition Tables for Epidemiologic Studies* as the reference database¹⁰.

We computed odds ratios of melanoma as an estimate of relative risk (RR) according to categories of dietary factor intake, using a matched analysis based on bivariate and multivariate conditional logistic regression models. We divided the subjects according to tertile cut-offs of dietary factors, based on distribution in referents; for gender-specific analyses, we used the median as cut-off for OR calculation. To compute *P*-values for linear trends we entered into the model intakes of dietary factors as continuous variables. Adjustment for energy in multivariate analysis was performed by entering overall energy intake as a covariate in the model; for selected nutrients showing in this analysis an abnormally high RR or a tendency towards an association with melanoma risk, we also repeated the analyses by adjusting for energy intake using the residual method¹¹. We fitted a cubic natural spline regression to evaluate the continuous relationship with melanoma risk of the two dietary factors more strongly associated with disease itself in the conditional logistic regression model, simultaneously adjusting for intake of the other dietary factor. To examine the relationship between age and intake of selected factors and total energy, we calculated the Spearman rank correlation coefficient.

Results

In total, 59 patients (28 males and 31 females, mean age 58.1 and 53.9 years, respectively) and 59 age- and sex-matched referents were included in the study. In Table 1 we report the RR of melanoma in the whole study population according to tertile distribution of the dietary factors, with and without adjustment for total energy intake. We found little evidence of an association with the majority of dietary constituents, with three exceptions: an excess risk associated with higher energy-adjusted intake of polyunsaturated fatty acids (PUFA) (*P* for linear trend = 0.080) and linoleic acid alone (*P* trend = 0.061),

Table 2 Multivariate analysis of melanoma risk* according to intake of linoleic acid and soluble carbohydrates, case-control study on environmental determinants of cutaneous melanoma, Modena Province, northern Italy

Dietary factor	Category		<i>P</i> for linear trend
	I†	II	
Linoleic acid	1.00	1.59 (0.61–4.13)	0.054
Soluble carbohydrates	1.00	0.48 (0.20–1.12)	0.017

*Relative risk of melanoma (95% confidence interval) using conditional logistic regression and adjusting for the remaining dietary factor.

†Referent category (average nutrient intake below the median as calculated in controls).

and, in both the energy-adjusted and unadjusted analyses, a lower risk in subjects characterised by a higher intake of soluble carbohydrates (*P* trend = 0.046). Total energy intake was not associated with risk of the disease. We also found a high risk of melanoma in subjects belonging to the middle tertile of intake for oleic acid and for total monounsaturated fatty acids, but the risks in the highest tertile and the trend analysis did not suggest an association between high intake of these dietary factors and disease risk.

When we entered in a multivariate model the two dietary factors most strongly associated with melanoma risk, linoleic acid and soluble carbohydrates, a stronger effect of these variables on melanoma risk than that detected in the bivariate analyses emerged (Table 2). In cubic spline multivariate regression analysis, risk increased smoothly at higher linoleic acid intake, particularly for low and high levels of this dietary factor, whilst there was little evidence of an association at intermediate levels of intake (Fig. 1). Concerning consumption of soluble carbohydrates, we detected a strong inverse linear relationship with disease risk only at the highest levels of intake (Fig. 2).

We repeated data analysis after stratification for gender, calculating the unadjusted and adjusted RR of melanoma

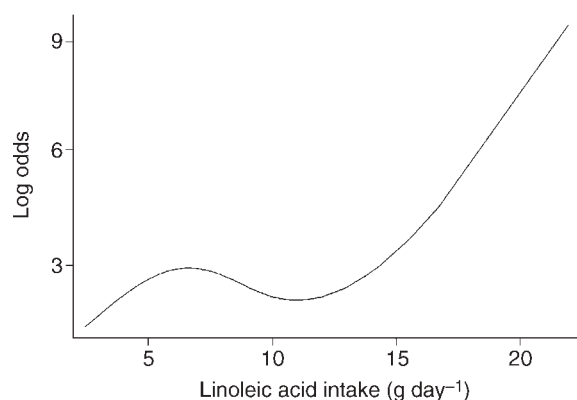


Fig. 1 Log odds of being a case at increasing intake of linoleic acid among population referents and patients with cutaneous melanoma, Modena, Italy (natural cubic spline regression adjusting for intake of soluble carbohydrates)

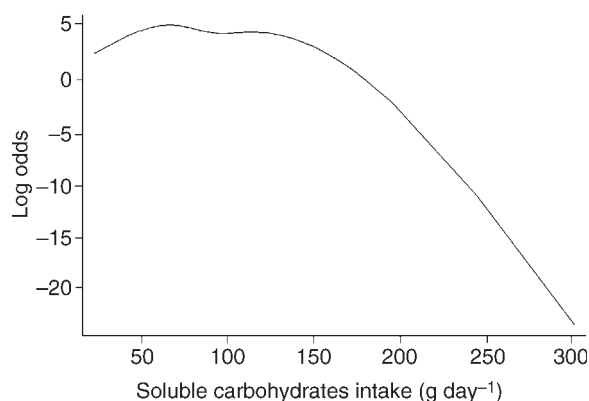


Fig. 2 Log odds of being a case at increasing intake of soluble carbohydrates among population referents and patients with cutaneous melanoma, Modena, Italy (natural cubic spline regression adjusting for linoleic acid intake)

associated with intake of linoleic acid and soluble carbohydrates equal or above the median value (Table 3). In multivariate analysis adjusting for energy and the other dietary factor, we observed in females a tendency towards excess risk in subjects having a higher intake of linoleic acid and a lower consumption of soluble carbohydrates. In males, despite RRs being generally

comparable to estimates obtained in females, estimates were statistically very unstable. Inclusion in the analysis of factors such as body mass index, smoking, education, family history, indicators of overall sun exposure and history of sunburn, phototype and number of atypical nevi, or using the residual method for total energy intake adjustment, or repeating the analyses after stratification for age (≤ 54 years and > 54 years), had no substantial effects on these risk estimates. However, using the residual method for total energy intake adjustment strongly reduced the very high RR observed in the second tertile of intake of total monounsaturated fatty acids and of oleic acid.

In Table 4 we report overall total energy intake and the intake of factors found to be associated with disease risk: PUFA, linoleic acid and soluble carbohydrates. We detected in patients a tendency towards a lower intake of soluble carbohydrates and a higher intake of PUFA and linoleic acid, despite the lower intake of energy particularly in females. When we analysed the possible association of age with these factors, inverse relationships emerged in referents, whilst in patients the only association we detected was the inverse one between age and soluble carbohydrates intake (Table 5).

Table 3 Gender-specific risk* of melanoma according to daily intake of selected dietary factors, case-control study on environmental determinants of cutaneous melanoma, Modena Province, northern Italy

Factor	Category		P for linear trend
	I†	II	
<i>Males</i>			
Linoleic acid			
Crude	1.00	1.67 (0.40–6.97)	0.391
Adjusted for energy	1.00	2.24 (0.42–12.05)	0.300
Adjusted for energy and soluble carbohydrates	1.00	2.01 (0.36–11.21)	0.421
Adjusted for soluble carbohydrates	1.00	1.89 (0.43–8.27)	0.388
Cut-off (g)	< 7.47	≥ 7.47	
Soluble carbohydrates			
Crude	1.00	0.67 (0.24–1.87)	0.242
Adjusted for energy	1.00	0.64 (0.20–2.06)	0.231
Adjusted for energy and linoleic acid	1.00	0.70 (0.22–2.30)	0.314
Adjusted for linoleic acid	1.00	0.65 (0.22–1.85)	0.243
Cut-off (g)	< 95.53	≥ 95.53	
<i>Females</i>			
Linoleic acid			
Crude	1.00	0.88 (0.32–2.41)	0.506
Adjusted for energy	1.00	1.24 (0.31–4.93)	0.077
Adjusted for energy and soluble carbohydrates	1.00	1.34 (0.31–5.76)	0.071
Adjusted for soluble carbohydrates	1.00	1.73 (0.47–6.39)	0.046
Cut-off (g)	< 7.62	≥ 7.62	
Soluble carbohydrates			
Crude	1.00	0.30 (0.08–1.09)	0.132
Adjusted for energy	1.00	0.19 (0.03–1.10)	0.096
Adjusted for energy and linoleic acid	1.00	0.14 (0.02–1.02)	0.077
Adjusted for linoleic acid	1.00	0.10 (0.02–0.65)	0.023
Cut-off (g)	< 103.91	≥ 103.91	

* Relative risk of melanoma (95% confidence interval) comparing subjects with average intake greater than the cut-off value versus remaining subjects.

† Referent category (average nutrient intake below the median as calculated in controls).

Table 4 Mean daily intake (\pm standard deviation) of selected dietary factors in population referents and in patients with cutaneous melanoma, Modena Province, northern Italy

Dietary factor	Females ($n = 62$)	Males ($n = 56$)
Total polyunsaturated fatty acids (g day^{-1})		
Referents	9.43 \pm 3.30	9.84 \pm 3.39
Patients	9.94 \pm 3.84	10.52 \pm 3.68
Linoleic acid (g day^{-1})		
Referents	7.59 \pm 2.66	8.00 \pm 2.91
Patients	8.20 \pm 3.52	8.73 \pm 3.36
Soluble carbohydrates (g day^{-1})		
Referents	112.19 \pm 56.29	97.54 \pm 40.27
Patients	95.84 \pm 24.73	86.25 \pm 27.26
Total energy intake (kcal day^{-1})		
Referents	2014.4 \pm 758.6	2228.8 \pm 720.4
Patients	1893.4 \pm 501.2	2195.6 \pm 518.7

Discussion

Mackie originally reported in 1974 an unusually high incidence of melanoma (of the leg) in subjects who had partially replaced their intake of saturated fat with PUFA¹². In subsequent studies, this investigator and co-workers reported higher dietary intakes and tissue levels of PUFA – and, in particular, of linoleic acid – in melanoma patients compared with referents^{13–15}. These observations fuelled a strong debate about the possible aetiological role of unsaturated fats^{16,17} and have also prompted a few epidemiological investigations. The first studies yielded no evidence of association between PUFA intake and melanoma risk^{18–21} or even an inverse relationship²². However, a subsequent prospective investigation by Veierod *et al.* from Norway yielded strong evidence of a direct association between PUFA intake and subsequent melanoma occurrence²³, although such a relationship was detected in females only, consistent with our results. A recent large case–control study from the USA also found a direct relationship between energy-adjusted linoleic acid intake and melanoma risk⁴, but that investigation differed from the present one in being hospital-based and using a so-called ‘brief’ food-frequency questionnaire, based on 60 items²⁴.

The different results obtained in the epidemiological studies published so far may be ascribed to methodological reasons, such as methodological limitations of some of the food-frequency questionnaires adopted, or to actual

differences in the role of dietary risk factors across different populations. In the present study, we used a validated semi-quantitative food-frequency questionnaire specifically developed for a northern Italy population, designed to analyse consumption of a very large number of foods and important details such as cooking methods and type of fat used as condiment before and after cooking.

In this investigation, the intake of linoleic acid (and, in part, of soluble carbohydrates) was associated with melanoma risk in the multivariate analysis, i.e. when adjusting for energy and/or for the other dietary factor, but not in the bivariate one. This seems to suggest that relative excess or deficiency of these dietary variables is of considerable importance, more than their absolute intakes, in enhancing melanoma risk.

There is suggestive biological and some epidemiological evidence indicating that PUFA and more specifically linoleic acid may increase cancer occurrence at some sites, such as the breast, colon and prostate^{25–29}, thus yielding concern about potential adverse effects of ‘high’ linoleic acid intake^{30,31}. Animal studies have also indicated that unsaturated lipids enhance ultraviolet-induced carcinogenesis^{32–34}, adding biological plausibility to a possible relationship between PUFA intake and melanoma aetiology. A recent laboratory study in epidermal reconstructs suggests that, following ultraviolet irradiation, omega-6 PUFA such as linoleic acid and arachidonic acid increase oxidative damage in melanocytes without inducing apoptosis, thus possibly increasing melanoma risk³⁵.

An unanticipated finding of the present study was the indication that a higher intake of soluble carbohydrates may decrease melanoma risk. This possible effect emerged independently from adjustment for overall energy intake, and it appeared to occur in both males and females, although the statistical association was considerably stronger in females. As far as we know, there are no prior studies which have analysed in detail the association between intake of specific carbohydrates and melanoma risk, thus hampering the evaluation of our results. In a recent hospital-based case–referent study, however, energy-adjusted intake of total carbohydrates correlated inversely with melanoma risk⁴. In our study population, intake of soluble carbohydrates is expected to be due

Table 5 Correlation between age and selected dietary factors in population referents and in patients with cutaneous melanoma, Modena Province, northern Italy. Data expressed as Spearman correlation coefficient (P -value)

	Age	
	Referents ($n = 59$)	Patients ($n = 59$)
Total polyunsaturated fatty acids (g day^{-1})	–0.299 (0.022)	–0.023 (0.866)
Linoleic acid (g day^{-1})	–0.253 (0.053)	0.010 (0.938)
Soluble carbohydrates (g day^{-1})	–0.233 (0.076)	–0.254 (0.053)
Total energy intake (kcal day^{-1})	–0.292 (0.025)	–0.144 (0.275)

mainly to the consumption of fruit, milk, sugar and sweets, suggesting the opportunity to further investigate a potential beneficial effect of fruit and milk intake, in particular. Recall bias does not appear to explain this association, as is also true for linoleic acid, since there is no awareness in the general population of a potential relationship of these (and other) dietary factors with melanoma risk.

The associations of linoleic acid and soluble carbohydrates with disease risk we detected were statistically more stable among females than in males. This might be due to a gender-specific effect of dietary factors in influencing melanoma risk, to an effect of unmeasured confounders or, we speculate, to a more accurate compilation of the questionnaire by the female group, with consequent beneficial effects on exposure classification and statistical precision of the point estimates.

Our results do not support sparse observations from epidemiology suggesting a relationship between melanoma and a higher intake of alcohol^{4,18,20,22} and vitamin C³⁶ or with a lower intake of vitamins A⁵, D⁴ and E^{21,22}, zinc^{21,22}, retinol³⁶, β -carotene^{4,22} and iron^{20,22}. However, the epidemiology literature about these associations has generally been inconsistent, such as in the case of alcohol consumption, which has also been inversely related to melanoma risk^{19,37}, and most studies have been unable to identify associations between the above-mentioned dietary factors and melanoma risk³.

Two important limitations of the present study must be outlined. First, the point estimates we calculated have reduced statistical precision, mainly due to the small study size: such instability of the risk estimates suggests caution in evaluating the findings of the present investigation, which need to be confirmed in larger populations. Furthermore, we cannot entirely rule out that the associations we detected might only be simple correlates of other true aetiological factors, such as dietary constituents that could not be analysed in the present study. However, we consider this hypothesis unlikely, due to the completeness of the food-frequency questionnaire and its consequent suitability to report extensively on the usual intake of dietary factors.

Acknowledgements

The authors are grateful to all the patients and referents who participated in the study, and to Giuliano Carrozzini, epidemiologist of the National Health Service Modena Unit, for providing the database of Modena residents. The authors acknowledge the co-operation in data analysis and manuscript review of Kenneth J Rothman of Boston University School of Public Health and of Annibale Biggeri of University of Florence School of Medicine. This study was supported by the 'Angela Serra Association for Cancer Research' of Modena and by the Italian Ministry of the

University and of the Scientific and Technological Research (grant no. 2002063519_001).

Conflicts of interest: None declared.

References

- Ferriman A. Large increases occur in cancers of skin and prostate. *British Medical Journal* 2003; **327**: 1306.
- de Vries E, Bray FI, Coebergh JW, Parkin DM. Changing epidemiology of malignant cutaneous melanoma in Europe 1953–1997: rising trends in incidence and mortality but recent stabilizations in western Europe and decreases in Scandinavia. *International Journal of Cancer* 2003; **107**: 119–26.
- Desmond RA, Soong SJ. Epidemiology of malignant melanoma. *Surgical Clinics of North America* 2003; **83**: 1–29.
- Millen AE, Tucker MA, Hartge P, Halpern A, Elder DE, Guerry D 4th, *et al.* Diet and melanoma in a case–control study. *Cancer Epidemiology, Biomarkers & Prevention* 2004; **13**: 1042–51.
- Naldi L, Gallus S, Tavani A, Imberti GL, La Vecchia C. Oncology Study Group of the Italian Group for Epidemiologic Research in Dermatology. Risk of melanoma and vitamin A, coffee and alcohol: a case–control study from Italy. *European Journal of Cancer Prevention* 2004; **13**: 503–8.
- Le Marchand L. Dietary factors in the etiology of melanoma. *Clinics in Dermatology* 1992; **10**: 79–82.
- Vinceti M, Bassissi S, Malagoli C, Pellicani G, Alber D, Bergomi M, *et al.* Environmental exposure to trace elements and risk of cutaneous melanoma. *Journal of Exposure Analysis and Environmental Epidemiology* 2005; **15**: 458–62.
- Pisani P, Faggiano F, Krogh V, Palli D, Vineis P, Berrino F. Relative validity and reproducibility of a food frequency dietary questionnaire for use in the Italian EPIC centres. *International Journal of Epidemiology* 1997; **26**(Suppl. 1): S152–60.
- Pala V, Sieri C, Palli D, Salvini S, Berrino F, Bellegotti M, *et al.* Diet in the Italian EPIC cohorts: presentation of data and methodological issues. *Tumori* 2003; **89**: 594–607.
- Salvini S, Parpinel M, Gnagnarella P, Maisonneuve P, Turrini A. *Banca dati di composizione degli alimenti per studi epidemiologici in Italia*. Milan: Istituto Europeo di Oncologia, 1998.
- Willett WC, Howe GR, Kushi LH. Adjustment for total energy intake in epidemiologic studies. *American Journal of Clinical Nutrition* 1997; **65**: 1220S–8S.
- Mackie BS. Malignant melanoma and diet. *Medical Journal of Australia* 1974; **1**: 810.
- Mackie BS, Johnson AR, Mackie LE, Fogerty AC, Ferris M, Baxter RI. Dietary polyunsaturated fats and malignant melanoma. *Medical Journal of Australia* 1980; **1**: 159–63.
- Mackie BS, Mackie LE, Curtin LD, Bourne DJ. Melanoma and dietary lipids. *Nutrition and Cancer* 1987; **9**: 219–26.
- Mackie BS, Mackie LE. Cancer and dietary lipids. *New Zealand Medical Journal* 1991; **104**: 322.
- Simpson LO. Melanoma and fatty acids. *New Zealand Medical Journal* 1990; **103**: 569.
- Holborow P. Melanoma and polyunsaturated fat; cancer and diet. *New Zealand Medical Journal* 1990; **103**: 515–6.
- Holman CD, Armstrong BK, Heenan PJ, Blackwell JB, Cumming FJ, English DR, *et al.* The causes of malignant melanoma: results from the West Australian Lions Melanoma Research Project. *Recent Results in Cancer Research* 1986; **102**: 18–37.
- Osterlind A, Tucker MA, Stone BJ, Jensen OM. The Danish case–control study of cutaneous malignant melanoma. IV.

- No association with nutritional factors, alcohol, smoking or hair dyes. *International Journal of Cancer* 1988; **42**: 825–8.
- 20 Stryker WS, Stampfer MJ, Stein EA, Kaplan L, Louis TA, Sober A, *et al.* Diet, plasma levels of β -carotene and α -tocopherol, and risk of malignant melanoma. *American Journal of Epidemiology* 1990; **131**: 597–611.
 - 21 Kirkpatrick CS, White E, Lee JA. Case-control study of malignant melanoma in Washington State. II. Diet, alcohol, and obesity. *American Journal of Epidemiology* 1994; **139**: 869–80.
 - 22 Bain C, Green A, Siskind V, Alexander J, Harvey P. Diet and melanoma. An exploratory case-control study. *Annals of Epidemiology* 1993; **3**: 235–8.
 - 23 Veierod MB, Thelle DS, Laake P. Diet and risk of cutaneous malignant melanoma: a prospective study of 50,757 Norwegian men and women. *International Journal of Cancer* 1997; **71**: 600–4.
 - 24 Block G, Hartman AM, Naughton D. A reduced dietary questionnaire: development and validation. *Epidemiology* 1990; **1**: 58–64.
 - 25 Rose DP. Dietary fat, fatty acids and breast cancer. *Breast Cancer* 1997; **4**: 7–16.
 - 26 Zock PL, Katan MB. Linoleic acid intake and cancer risk: a review and meta-analysis. *American Journal of Clinical Nutrition* 1998; **68**: 142–53.
 - 27 Bartsch H, Nair J, Owen RW. Dietary polyunsaturated fatty acids and cancers of the breast and colorectum: emerging evidence for their role as risk modifiers. *Carcinogenesis* 1999; **20**: 2209–18.
 - 28 Newcomer LM, King IB, Wicklund KG, Stanford JL. The association of fatty acids with prostate cancer risk. *Prostate* 2001; **47**: 262–8.
 - 29 Astorg P. Dietary *n*-6 and *n*-3 polyunsaturated fatty acids and prostate cancer risk: a review of epidemiological and experimental evidence. *Cancer Causes & Control* 2004; **15**: 367–86.
 - 30 Erickson KL. Is there a relation between dietary linoleic acid and cancer of the breast, colon, or prostate? *American Journal of Clinical Nutrition* 1998; **68**: 5–7.
 - 31 Hamazaki T, Okuyama H. The Japan Society for Lipid Nutrition recommends to reduce the intake of linoleic acid. A review and critique of the scientific evidence. *World Review of Nutrition and Dietetics* 2003; **92**: 109–32.
 - 32 Black HS, Lenger W, Phelps AW, Thornby JI. Influence of dietary lipid upon ultraviolet-light carcinogenesis. *Nutrition and Cancer* 1983; **5**: 59–68.
 - 33 Mathews-Roth MM, Krinsky NI. Effect of dietary fat level on UV-B induced skin tumors, and anti-tumor action of β -carotene. *Photochemistry and Photobiology* 1984; **40**: 671–3.
 - 34 Black HS, Thornby JI, Gerguis J, Lenger W. Influence of dietary omega-6, -3 fatty acid sources on the initiation and promotion stages of photocarcinogenesis. *Photochemistry and Photobiology* 1992; **56**: 195–9.
 - 35 Cario-Andre M, Briganti S, Picardo M, Nikaido O, de Verneuil H, Taieb A. Polyunsaturated fatty acids partially reproduce the role of melanocytes in the epidermal melanin unit. *Experimental Dermatology* 2005; **14**: 194–201.
 - 36 Feskanich D, Willett WC, Hunter DJ, Colditz GA. Dietary intakes of vitamins A, C, and E and risk of melanoma in two cohorts of women. *British Journal of Cancer* 2003; **88**: 1381–7.
 - 37 Sigvardsson S, Hardell L, Przybeck TR, Cloninger R. Increased cancer risk among Swedish female alcoholics. *Epidemiology* 1996; **7**: 140–3.