Association between sun exposure and serum 25-hydroxyvitamin D₃ levels

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Abstract. Low vitamin D levels are associated with increased risk of many diseases. However, there is uncertainty about the amount of sun exposure required to increase vitamin D. Personal UV dosimeters were used to quantify the association between sun exposure over 8-weeks, corrected for the proportion of skin not covered by clothing, and change in serum 25-hydroxyvitamin D₃ (25OHD₃) concentration. There was a non-linear association between UV exposure and 25OHD₃. This finding suggests that vitamin D status can be increased by regular small sun exposures and that greater exposures result in only small additional increases in 25OHD₃.

Background

There is increasing evidence from prospective epidemiological studies that low vitamin D status, as measured by blood 25-hydroxyvitamin D (25OHD) concentration, predicts a wide range of diseases (Scragg 2011). About 90% of vitamin D in humans is synthesised through sun exposure. A better understanding of the relationship between sun exposure and vitamin D status is required so that sun exposure policies can be developed that result in increased vitamin D levels without also increasing risk of skin cancer.

Two approaches have been used to quantify the association between sun exposure and vitamin D status. These have been either: a) solaria studies of change in 25OHD level from measured doses of artificial UV-B; or b) observational studies comparing 25OHD levels with recent UV exposure. Most of the latter have measured UV exposure with questionnaires, complemented by measures of ambient UV exposure. Few have used objective measures of UV exposure (such as polysulphone badges or electronic dosimeters) and only two have quantified the association between UV exposure and 25OHD status (Datta et al, 2012; Kimlin et al, 2014). The aim of the current study is to examine, in a large multi-ethnic community sample, the association between natural outdoor UV exposure and change in serum 25OHD levels over 4-weeks using electronic UV dosimeters.

Methods

The participants in this report comprise 503 volunteers recruited from the community in Auckland (n=332) and Dunedin (n=171), equally distributed across the adult ages of 18-85 years and the four main ethnic groups (European 126, Maori 126, Pacific 123 and Asian 128). Baseline interviews were carried out during February-October in 2008 and 2009, in which the following information was collected: self-defined ethnicity, self-reported reaction to sun exposure (Fitzpatrick scale), weight and height to estimate body mass index and body surface area, and skin colour measured by a portable spectrophotometer. At this visit, participants were given a personal electronic dosimeter to wear on the outer wrist when outdoors and a daily diary to record their main clothing during following three day-time periods: before 11am, 11am-4pm, and after 4pm. The diary was used to estimate the proportion of their skin exposed to UV radiation in each time period. Participants returned weekly for 8-weeks for the dosimeter data to be downloaded and clothing diary to be checked and replaced. Blood samples were collected at the end of week 4 and week 8, to measure serum 25(OH)D₃ concentrations using Liquid Chromatography-Tandem Mass Spectrometry.

The dosimeters sampled erythemally-weighted UV irradiance at 8-second intervals (Allen & McKenzie 2005). These data were aggregated for each of the three daily time periods (above) and converted to standard erythemal doses (SED), which were corrected for clothing to generate equivalent full-body exposures, SEDₑFB. Data were analysed using a multivariate linear mixed model with an autoregressive structure for repeated measures to estimate simultaneously both between-person and within-person effects (Neuhaus & Kalbfleisch 1998). Full details of methods and results have been reported (Scragg et al, 2014).

Results

Median SED of UV radiation received was 7.3 in weeks 1-4 and 6.5 in weeks 5-8, less than 2% of ambient UV radiation. When converted to SEDₑFB per week, this was 0.33 in weeks 1-4 and 0.34 in weeks 5-8. Mean (SD) 25OHD₃ concentration was 48.2 (24.2) nmol/L at the end of week 4 and 47.4 (22.4) nmol/L at the end of week 8.

Participants were ranked into nine groups by increasing cumulative SEDₑFB per week during each 4-week period, and mean 25(OH)D₃ concentrations at the end of each 4-week period were calculated for each group (Figures 1 & 2). The variation in mean 25(OH)D₃ from the lowest to the highest SED group for each 4-week period was substantial, being from 34.9 up to 64.7 nmol/L for week-4 and 38.4 to 57.3 nmol/L for week-8. This represents the between-person variation in 25(OH)D₃ concentration.

The change in 25(OH)D₃ from week-4 to week-8 for each participant was calculated, and then ranked by increasing SEDₑFB per week received during weeks 5-8 (Figure 3). This change, which represents the within-person variation in vitamin D status, was much smaller than the between-person variation, and varied from a mean change of -5.4 nmol/L up to 2.1 nmol/L across the range of grouped SEDₑFB exposures in weeks 5-8. UV exposures < 0.4 SEDₑFB per week (groups 1-5) were associated with decreasing 25OHD₃ concentrations in weeks 5-8, from weeks 1-4.
Statistical analyses identified a non-linear association between SED_{EFB} exposure and 25(OH)D_3 concentration over the 8-week period. Using the multivariate mixed model described above, the coefficient for the between-person variation showed that average exposures of 0.1, 1.0 and 10 SED_{EFB} per week were associated with 25(OH)D_3 levels that were greater by 10.9, 21.9 and 32.8 nmol/L, respectively, than an average exposure of 0.01 SED_{EFB} per week (ie. almost zero), adjusting for age, sex, ethnicity, body mass index, skin colour, number of days of UV exposure and within-person variation in UV exposure. A graph of this coefficient showed that most of the increase in 25(OH)D_3 associated with between-person variation in UV exposure occurred at exposures of < 2 SED_{EFB} per week, and that greater exposures were associated with only small additional increases in 25(OH)D_3 concentration.

Conclusions
Our results indicate a non-linear association between sun exposure and serum vitamin D concentration which suggest that vitamin D status can be increased by regular small sun exposures (< 2 SED_{EFB} per week), and that greater exposures result in only small additional increases in 25(OH)D_3.

References