Associations between vitamin D status and measures of glycaemia in participants with normoglycaemia, impaired fasting glucose and type 2 diabetes

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#### **Introduction**

There is emerging evidence linking vitamin D deficiency to impaired  $\beta$ -cell function, insulin resistance, and glucose intolerance, all of which are central to the pathogenesis of type 2 diabetes<sup>1</sup>.

When participants were separated based on FPG (group one  $\leq 6 \text{ mmol/l} (n = 49) \text{ vs group } 2 \geq 6.1 \text{ mmol/l} (n = 14)) \text{ a 2-way}$  mixed ANOVA revealed a significant main effect of group on

### Aims:

- To identify if baseline vitamin D status is associated with future glycaemic control.
- To assess whether vitamin D status differs between two groups when classified by glucose control (above or below 6.1 mmol/l: cut off point for impaired fasting glucose<sup>2</sup>).

### Methods:

Following ethical approval 104 participants (female 53; male 51) were recruited from the local community (Mid-Wales). Participants with varying levels of glucose control attended the laboratory in a fasted state on three occasions, each separated by 6 months. During each visit blood was drawn from an antecubital vein and stored for later analyses.

Vitamin D status was determined (25 hydroxyvitamin  $D_3$ ) by HPLC-MS/MS. Remaining samples were analysed for fasting plasma glucose (FPG), fasting plasma insulin (FPI), and HbA1c at the Diabetes Research Network Wales Laboratory.

25(OH)D concentration (F(1,59) = 4.860, p < 0.05, partial n2 = 0.076) and a main effect of time point (F(2,118) = 75.751, p < 0.05, partial n2 = 0.562). There was no statistically significant group × time interaction (F(2,118) = 0.680, p = 0.51, partial n2 = 0.011)



**Figure 2:** 25(OH)D concentration at the different time points in normal and high FPG groups (FPG  $\leq$  6 mmol/l vs  $\geq$  6.1 mmol/l).

# **Results:**

Significant negative correlations were observed between  $25(OH)D_3$  at 0 months and HbA1c at 12 months (r = -0.241, n = 75, p = 0.04) and FPI at 12 months (r = -.225, n = 81, p = 0.04), but not FPG at 12 months (r = -.105, n = 80, p = 0.35).



## **Conclusions:**

Vitamin D status only accounted for a small proportion of the variance (~5%) in the measures of glycaemia at 12 months, however, the development of type 2 diabetes is multifactorial and any easily modifiable risk factors are noteworthy. A seasonal variation in vitamin D status was observed for both FPG groups, and participants in the low FPG group had higher vitamin D concentrations than those in the high FPG group across the seasons.

These findings may provide additional evidence of a protective effect of vitamin D in relation to glycaemic control and the development of type 2 diabetes, although, further studies are required to determine cause and effect.

#### **References**

1.Song, Y. & Manson, J. E. (2010) Vitamin D, Insulin Resistance, and

**Figure 1:**The relationship between 25(OH)D at 0 months (winter) and HbA1c at 12 months (winter).

Type 2 Diabetes. Current Cardiovascular Risk Reports, 4, 40–47. 2.World Health Organisation (WHO) (2006). Definition and Diagnosis of Diabetes Mellitus and Intermediate Hyperglycaemia: Report of a WHO/IDF Consultation.

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