Vitamin D deficiency in patients with intellectual disabilities: prevalence, risk factors and management strategies

Valeria Frighi, Alireza Morovat, Matthew T. Stephenson, Sarah J. White, Christina V. Hammond and Guy M. Goodwin

Background
People with intellectual disabilities have a high risk of osteoporosis and fractures, which could partly be as a result of vitamin D deficiency.

Aims
To compare the serum vitamin D (25(OH)D) levels of 155 patients with intellectual disabilities under psychiatric care and 192 controls, investigate potential risk factors for vitamin D deficiency in people with intellectual disabilities and assess available treatments.

Method
Cross-sectional observational study followed by treatment evaluation.

Results
Almost twice as many patients with intellectual disabilities had vitamin D deficiency (25(OH)D < 50 nmol/l) compared with controls (77.3% v. 39.6%, P < 0.0001). In the intellectual disabilities group, winter season (P < 0.0001), dark skin pigmentation (P < 0.0001), impaired mobility (P = 0.002) and obesity (P = 0.001) were independently associated with lower serum 25(OH)D. In most patients, 800 IU colecalciferol daily normalised 25(OH)D levels.

Conclusions
Vitamin D deficiency is highly prevalent in people with intellectual disabilities, partly because of insufficient exposure to sunlight. Screening and treatment strategies, aiming to reduce these patients’ high fracture risk, should be introduced. Similar strategies may be required in other psychiatric populations at risk for fractures and with a tendency to spend excessive time indoors.

Declaration of interest
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Vitamin D deficiency is a well-established precursor to osteopenia and osteoporosis and, in the most severe cases, to osteomalacia in adults and rickets in children. Proximal myopathy can accompany the bone changes, leading to a tendency to fall. The potential outcome of these abnormalities is an increased risk of fractures. Patients with intellectual disabilities in institutions are likely to have reduced exposure to sunlight, the major determinant of vitamin D production. Studies in a number of countries looking at institutionalised patients with intellectual disabilities have shown a prevalence of vitamin D deficiency as defined by serum 25 (OH)D < 50 nmol/l of between 57% and 83% and of 100% in a sample of patients in whom sunshine exposure was virtually non-existent. People with intellectual disabilities, particularly those with epilepsy and treated with anti-epileptics, are also known to have a high rate of osteoporosis and fractures. This increased risk will be partly related to epilepsy itself, but could also be as a result of vitamin D deficiency because anti-epileptic drugs increase 25-hydroxyvitamin D (25(OH)D) metabolism. However, to our knowledge, no information is available on the vitamin D status of community-living patients with intellectual disabilities and, whether institutionalised or not, no information is available for patients with intellectual disabilities living in the UK. Furthermore, no comparative studies between intellectual disabilities and matched general population controls have been published.

In this paper we report the prevalence of vitamin D deficiency, defined as a serum vitamin D (25(OH)D) of < 50 nmol/l in a community living group of people with intellectual disabilities under psychiatric care in comparison with a healthy control sample drawn from the same geographical area (Oxfordshire, UK, latitude 51°N). We also examine the risk factors for hypovitaminosis D in people with intellectual disabilities, its potential consequences in terms of low bone mineral density and fractures and evaluate the results of its treatment. The term ‘intellectual disabilities’ refers to the condition of people with disabilities characterised by significant limitations both in cognitive functioning and adaptive behaviour (conceptual, social and practical adaptive skills) that originate before age 18 years.

Study population
The 155 patients with intellectual disabilities presented in this paper were part of the Oxford Learning Disabilities Study. This was a cross-sectional observational study, approved by the regional ethics committee: all patients aged 18–70 years under the care of the Oxfordshire-based psychiatrists of the Oxfordshire Learning Disability NHS Trust were eligible. Capacity was assessed as outlined in the Mental Capacity Act 2005 Code of Practice. Written informed consent was given by capacitous patients. Those assenting (but formally incapacitous) were only entered into the study after unanimous agreement was obtained from key people (including one or more carers and/or relatives) that inclusion in the study was in the patient’s best interest.

The study investigated the side-effects of antipsychotic treatment as captured primarily by laboratory measures. In all the study patients (the intellectual disabilities group) who agreed
to give a blood sample, 25(OH)D was measured because of the widespread use of anti-epileptic medication, which can decrease vitamin D levels, and because of the clinical history of non-healing or low trauma fractures in several patients with no obvious risk factors for fractures.

The control group was made up of 192 people from the Oxfordshire general population. These were blood donors from the NHS Blood and Transplant Service of the Oxford University Hospitals NHS Trust, who had given consent for their gender, age and serum 25(OH)D levels to be used for establishing reference laboratory data.

Participants in the intellectual disabilities group were seen in their homes by a study nurse, who took consent, measured height and weight, collected information from carers about their medical and drug history, and took a fasting blood sample for various analytes including 25(OH)D and calcium. An attempt was also made to quantify the amount of exposure to sunlight and the use of sun protector creams in each individual patient, but the answers obtained from carers did not lead to high-quality, analysable data. Patients with vitamin D deficiency who were deemed by their psychiatrists and carers as able to tolerate the procedure without undue distress were referred for measurement of bone mineral density by a dual-energy x-ray absorptiometry (DXA) scan (Hologic).

Patients with normal renal function who had been diagnosed with vitamin D deficiency in the study and had not yet been given supplements by their general practitioners (GPs) were entered in a good clinical practice initiative. This consisted of offering colecalciferol supplementation according to the clinical algorithm used for the patients with vitamin D deficiency treated at the Metabolic Bone Unit/Oxfordshire Osteoporosis Centre of the Nuffield Orthopaedic Centre (Oxford University Hospitals NHS Trust). The treatment was started following a clinical assessment at the Metabolic Bone Unit and consisted of an oral loading dose of 50 000 IU of colecalciferol daily for 3 or 6 days (according to whether baseline serum 25(OH)D levels were 25–49 nmol/l or <25 nmol/l respectively), followed by monthly 50 000 IU as maintenance. Daily calcium supplements were added according to individual patient's intake, assessed with the help of carers, by a nurse-administered questionnaire, and, when possible, by the use of 3-day food diaries. Specifically, patients with a very low calcium intake (less than 500 mg daily) were given the equivalent of 1 g daily elemental calcium supplements, those with an intermediate intake (500 to <1000 mg) were given 500 mg daily, and those taking 1 g or more daily were considered replete and not given any calcium supplements.

As a result of a change in prescription rules, the high-dose vitamin D preparation used by the specialist service became unavailable to new patients during the course of the good practice initiative. Patients who had been started on the high-dose preparation could continue on it, but patients who had not yet been started on supplementation were referred for treatment to their GPs. For the latter patients, supplements mostly consisted of 400 IU of colecalciferol together with 500 mg calcium given twice daily, although five patients were treated with 400 IU colecalciferol daily and another four with various other preparations. Safety of the high-dose intermittent treatment used by the hospital service was evaluated by calcium and 25(OH)D measurements 1 month after the loading dose. Efficacy of maintenance treatment was evaluated by a 25(OH)D measurement after a minimum of 3 months from starting either the high-dose intermittent or the low-dose daily treatment.

Available data from the controls were age, gender, 25(OH)D level and date of blood sampling.

Biochemical analyses
We measured 25(OH) D using a routine in-house liquid chromatography and tandem mass spectrometry method. In 2010, after the release of the National Institute of Standards and Technology’s reference material for 25(OH)D, a re-standardised assay was used accordingly. The method specifically measured 25(OH) D2 and D3, and had between-batch coefficients of variation (CV) of 19.0% and 13.5% at 25(OH) D concentrations of 14.5 and 42.8 nmol/l, respectively. The limit of quantification, based on an ability to measure 25(OH)D with a CV of <20%, was taken to be 15 nmol/l. However, in 29 patients the measured 25(OH)D levels were lower than the limit of quantification, and down to a measured 4.5 nmol/l, where the between-batch CV of the assay was 26.8%. Calcium was measured by arsenazo III method on an Advia 2400 analyser. The instrument was also used for measuring phosphate by the use of the phosphomolybdate method, as well as the kinetic assay of alkaline phosphatase by the modified diethanolamine-buffered enzymatic method.

Bone mineral density
Standard definitions of osteopenia and osteoporosis were used for interpretation of the DXA scans. Osteopenia was defined by a T-score of −1.1 to −2.5 (i.e. bone mineral density (BMD) 1.1–2.5 standard deviations below the mean BMD of the normal population at age 30 years) and osteoporosis by a T-score of less than −2.5 (i.e. BMD more than 2.5 standard deviations below the mean BMD of the normal population at age 30 years). Z-scores were also used as they allow comparison with an age- and gender-matched population.

Season
Normally, around 90% of the body's vitamin D is produced in the skin from exposure to solar ultraviolet B radiation but at latitudes above 42°, the skin does not make any vitamin D in the winter months. Therefore, vitamin D data were segregated according to two seasons with high (Summer: June to November) and low (Winter: December to May) concentrations of 25(OH)D.

Statistical analyses
Data were analysed using IBM SPSS (version 20) for Windows. As baseline serum 25(OH)D was not normally distributed, non-parametric tests were used. In the intellectual disabilities and the healthy control groups, baseline 25(OH) D levels were compared by Mann–Whitney test. Prevalence of vitamin D deficiency in the two groups using different 25(OH)D cut-off levels was compared by chi-square.

For analyses of the data for individuals in the intellectual disabilities group, the degree of disability was classified as mild or severe, a category which included the diagnostic groups of moderate, severe and profound. This decision was made to avoid analyses of groups with very small number of participants and was based on the rationale that only individuals with mild intellectual disabilities are capable of independent or partly independent living.

In the intellectual disabilities group, median 25(OH)D levels were compared by gender, skin pigmentation (dark v. White), mobility (impaired v. normal), body mass index (BMI ≥30 v. <30), use of anti-epileptics (on any anti-epileptic v. not on anti-epileptics), degree of intellectual disability (mild v. severe) using the Mann–Whitney test. A correlation with age was investigated by Spearman correlation coefficient.
A stepwise linear regression model was used to test for independent associations with 25(OH)D concentrations, where log transformed 25(OH)D was the dependent variable and the above factors were the independent variables. We also compared the 25(OH)D values of patients with a history of fractures v. those without known fractures as a whole group and after the exclusion of patients with obvious causes of fractures (such as highly traumatic falls, long-term untreated hypogonadism).

In the treatment analyses, the rates of normalisation of serum 25 (OH)D to 50 nmol/l were compared for the high-dose intermittent, the daily 800 IU and the daily 400 IU colecalciferol groups by the Freeman–Halton extension of Fisher’s exact test. Mean 25 (OH)D levels were compared for the three groups at baseline and after starting treatment by ANOVA, with addition of the Welch test for equality of means in view of the uneven size of the groups. Changes from baseline within each treatment group were assessed by paired t-tests. Differences in the change of serum 25 (OH)D levels between baseline and after starting treatment were compared across the groups by ANOVA, with addition of the Welch test. Five of the 155 participants with intellectual disabilities (3.2%) were already on treatment with vitamin D preparations because of osteoporosis or osteoporotic fractures, when first assessed in the study. These patients were excluded from baseline analyses but their data contributed to treatment evaluation.

**Diagnostic thresholds for vitamin D deficiency**

We used a serum concentration of 25(OH)D of <50 nmol/l, which is the most widely accepted threshold, because this is the level at which parathyroid hormone secretion starts to increase, promoting bone demineralisation. We also investigated the prevalence of severe deficiency, defined as serum 25(OH)D >30 nmol/l, as this is the level at which osteomalacia can appear.

**Results**

**Demographics**

The 155 participants in the intellectual disabilities group were aged 42.8 years (s.d. = 12.2), 55% were males and 94% were of White ethnicity. The 192 participants in the control group were aged 42.8 years (s.d. = 13.0), 51% were males and 94% were of White ethnicity. The 192 participants in the control group were similar to the intellectual disabilities group in age and gender (43.0 years (s.d. = 12.2), 55% male). Ethnicity was not formally recorded but it was assumed to reflect that of the general population of Oxfordshire, which is 92% White.

**Characteristics of the intellectual disabilities group**

The degree of intellectual disabilities was mild, moderate, severe and profound in 49.7%, 32.3%, 14.2% and 3.9%, respectively. A total of 90 patients (58%) had a psychiatric diagnosis with or without challenging behaviour whereas 42% were under psychiatric care because of challenging behaviour. In total, 40.6% of the group were on anti-epileptics. The great majority lived in small group homes of two to six patients with their carers, the remaining living independently or with their families.

**Vitamin D levels in both groups**

The overall prevalence of vitamin D deficiency was 77.3% in the intellectual disabilities group and 39.6% in the control group (P < 0.0001), with a wide seasonal variation that was more marked in the control than the intellectual disabilities group. Different cut-offs according to group and season of blood sampling are shown in Table 1.

Median 25(OH)D levels were 28.7 nmol/l (IQR, 16.9–49.1) for the intellectual disabilities group and 57.9 nmol/l (IQR, 39.9–77.9) for the control group (P < 0.0001). In the participants with vitamin D deficiency, median (IQR) 25(OH)D levels were 22.4 nmol/l (IQR, 15.2–34.5) and 35.8 nmol/l (IQR 28.3–49.1) in the intellectual disabilities and the control groups, respectively (P < 0.0001). Phosphate and adjusted calcium levels were normal in all patients.

During the summer season, median serum 25(OH)D concentrations for the control and the intellectual disabilities populations were 65.8 nmol/l (IQR, 51.8–85.1) and 36.7 nmol/l (IQR, 22.6–53.2), respectively (P < 0.0001). During winter, the control and intellectual disabilities groups had median 25(OH)D levels that were 32.6 nmol/l (IQR, 26.2–41.7) and 20.3 nmol/l (IQR, 13.3–31.2), respectively (P < 0.0001), Fig. 1.

Table 1 shows 25(OH)D levels in the intellectual disabilities group according to their demographic and clinical characteristics, as well as to season of blood sampling. There was no correlation between serum 25(OH)D and age and no difference in 25(OH)D levels between those patients with or without a psychiatric diagnosis.

In linear regression analysis, winter season (β = 0.488, P < 0.0001), dark skin pigmentation (β = 0.717, P < 0.0001),

<table>
<thead>
<tr>
<th>25(OH)D</th>
<th>Intellectual disabilities group (n = 155)</th>
<th>Control group (n = 192)</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt;50 nmol/l</td>
<td>77.3</td>
<td>39.6</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Summer</td>
<td>69.0</td>
<td>19.9</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Winter</td>
<td>89.4</td>
<td>87.5</td>
<td>NS</td>
</tr>
<tr>
<td>&lt;30 nmol/l</td>
<td>50.0</td>
<td>10.9</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Summer</td>
<td>34.5</td>
<td>0.7</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Winter</td>
<td>69.7</td>
<td>35.7</td>
<td>&lt;0.0001</td>
</tr>
</tbody>
</table>

The overall percentages refer to the whole group irrespective of season. The percentages by season represent the participants within the relevant 25(OH)D cut-off as a percentage of the participants tested in that season.
Impaired mobility (β = 0.530, P = 0.002) and obesity (β = 0.292, P = 0.001) were independently associated with lower 25(OH)D levels. Together, these factors explained 32% of the variance.

Answers obtained from carers about patients’ exposure to sunlight were too subjective to allow the collection of quantifiable data. On the basis of direct questioning as well as observation, it seemed that the use of sun protector creams was widespread, if not universal, when it was warm and not too cloudy and patients were taken outdoors. Nevertheless, the data collected was anecdotal.

**Clinical symptoms of vitamin D deficiency**

Although osteoporosis is asymptomatic until a fracture occurs, an attempt was made to investigate key symptoms of osteomalacia, namely generalised bone pain and proximal muscle weakness. No patient seemed in pain but it was not possible to assess muscle power except in a very small minority of individuals, because of difficulties with cooperation.

**Bone mineral density**

In total, 47 patients with vitamin D deficiency (median 25(OH)D of 22.2 nmol/l (IQR, 12.6–31.4)), and no evidence of premature hypogonadism, underwent a DXA scan. This showed low bone mineral density at the spine, hip or both in 70.2% (n = 33) of individuals: 31.9% (n = 15) with T-scores less than −2.5, consistent with osteoporosis and 38.3% (n = 18) with T-scores of −1.1 to −2.5, consistent with osteopenia. Additionally, the z-scores were consistent with the T-scores in 85% (n = 40) of individuals. This signifies that the majority of patients with abnormal DXA scans had decreased bone mineral density not only compared with the normal population at age 30 but also a reference population of their same age and gender. Moreover, although a DXA scan cannot differentiate between osteoporosis and osteomalacia, it is likely that patients with very low T-score may have also had osteomalacic changes. This is corroborated by the finding of raised alkaline phosphatase in 14.3% (n = 10) of the patients with severe vitamin D deficiency (25(OH)D < 30 nmol/l), and by the subsequent normalisation of the enzyme following colecalciferol treatment.

**Fractures**

Twenty participants in the intellectual disabilities group (12.9% of the whole group) had a history of one or more fractures, as reported by their carers and whenever possible cross-checked on the hospital radiology database. This represented 7.5% of men and 18.7% of women (χ² = 4.3, P = 0.038), 13.1% of those of White and 10% of those of non-White ethnicity (not significant). The mean age at fracture was 48.6 years (s.d. = 13.6). A history of fractures was more prevalent in patients on anti-epileptics (20.6% (n = 13) vs. 7.6% (n = 7) in patients not on anti-epileptics, χ² = 6.06, P = 0.014), but there was no difference in the frequency of fractures between patients with normal 25(OH)D levels or patients with vitamin D deficiency (12% of patients in each group).

A history of fractures was also analysed, after the exclusion of patients with obvious causes such as catastrophic falls or long-term untreated hypogonadism. This left 13 patients with a positive fracture history. In this analysis, fractures were apparently more frequent in patients with vitamin D deficiency (13 patients had a fracture) than in those with normal serum 25(OH)D levels (no patients had a fracture) but the difference was not statistically significant (χ² = 3.6, P = 0.059).

### Table 2 Median (IQR) serum 25(OH)D levels in the intellectual disabilities group not on supplements

<table>
<thead>
<tr>
<th>Variable</th>
<th>n</th>
<th>25(OH)D, nmol/L: median (IQR)</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gender</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Males</td>
<td>77</td>
<td>27.6 (17.7–50.1)</td>
<td>0.348</td>
</tr>
<tr>
<td>Females</td>
<td>73</td>
<td>28.9 (16.4–47.9)</td>
<td></td>
</tr>
<tr>
<td>Season</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Summer</td>
<td>84</td>
<td>36.7 (22.6–53.3)</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Winter</td>
<td>66</td>
<td>20.3 (13.2–31.3)</td>
<td></td>
</tr>
<tr>
<td>Skin colour</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Dark skin</td>
<td>9</td>
<td>16.3 (11.5–17.3)</td>
<td>0.002</td>
</tr>
<tr>
<td>White skin</td>
<td>141</td>
<td>30.8 (18.1–49.9)</td>
<td></td>
</tr>
<tr>
<td>Level of disabilities</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mild intellectual disabilities</td>
<td>77</td>
<td>34.1 (19.4–51.6)</td>
<td>0.039</td>
</tr>
<tr>
<td>Severe intellectual disabilities</td>
<td>73</td>
<td>22.7 (16.7–38.3)</td>
<td></td>
</tr>
<tr>
<td>Body mass index³</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>≥ 30</td>
<td>59</td>
<td>25.9 (15.8–41.2)</td>
<td>0.065</td>
</tr>
<tr>
<td>&lt; 30</td>
<td>73</td>
<td>31.7 (19.2–53.7)</td>
<td></td>
</tr>
<tr>
<td>Mobility</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Impaired</td>
<td>23</td>
<td>17.2 (10.7–28.9)</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Normal</td>
<td>127</td>
<td>32.2 (18.7–50.3)</td>
<td></td>
</tr>
<tr>
<td>Medication</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>On anti-epileptics</td>
<td>59</td>
<td>24.3 (16.5–41.8)</td>
<td>0.092</td>
</tr>
<tr>
<td>Not on anti-epileptics</td>
<td>91</td>
<td>33.0 (17.3–52.8)</td>
<td></td>
</tr>
</tbody>
</table>

³ Body mass index data are missing for patients in whom height and weight measurements with standard equipment were not possible (abnormal body posture, wheelchair bound or unable to stand still).

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Vitamin D deficiency in patients with intellectual disabilities
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Prevalence of vitamin D deficiency (25(OH)D in winter. Thus, whereas in winter there was an almost 90% prevalence in the general population. Moreover, the difference in prevalence is more severe in such patients than in individuals from the control group. In over a third of the intellectual disabilities group but in less than twice as common in the intellectual disabilities group compared with the control group. In summer the prevalence was 3.5 times higher in both groups, in the intellectual disabilities group compared with the general population and also the first study on community living rather than institutionalised patients with intellectual disabilities.

Our study has shown that the prevalence of vitamin D deficiency, defined as percentage of patients who reached 25(OH)D levels ≥ 50 nmol/l while on treatment, and in the intellectual disabilities group achieved the rate of normalisation, was 100% (n = 28) for the high-dose intermittent colecalciferol group, 80.7% (n = 46) for the 800 IU daily and 20% (n = 1) for the 400 IU daily colecalciferol group (P < 0.0001). There was no difference in age, gender or skin pigmentation between the treatment groups. Table 3 shows the levels at baseline and after 3 months of treatment according to treatment group. There were no cases of treatment emergent hypercalcaemia in any of the groups.

### Discussion

Our study has shown that the prevalence of vitamin D deficiency, as defined by a serum 25(OH)D level < 50 nmol/l, is markedly higher in the group with intellectual disabilities than in the control group from the general population with similar demographic characteristics living in the same geographical area (Oxfordshire, UK; latitude 51°N). It has also shown that vitamin D deficiency is more severe in such patients than in individuals from the general population. Moreover, the difference in prevalence between the two groups was far more prominent in summer than in winter. Thus, whereas in winter there was an almost 90% prevalence of vitamin D deficiency (25(OH)D < 50 nmol/l) in both groups, in summer the prevalence was 3.5 times higher in the intellectual disabilities group compared with the control group. Severe vitamin D deficiency (25(OH)D < 30 nmol/l), was twice as common in the intellectual disabilities group compared with the control group in winter whereas in summer it was found in over a third of the intellectual disabilities group but in less than 1% of the control group.

The standard diet contains only minimal quantities of vitamin D,20,21 so by far the greatest contribution to the body’s vitamin D levels comes from skin irradiation. Accordingly, insufficient exposure to sunlight is the most likely cause of the difference in the vitamin D levels between the intellectual disabilities group and the control group. Moreover, patients with impaired mobility, who generally need help to go out and may have to cover themselves more than others to maintain body temperature, had vitamin D levels almost 50% lower than those of patients with normal mobility. This further supports our interpretation of the data.

High prevalence of vitamin D deficiency has been reported in other studies in people with intellectual disabilities.3,4 However, to our knowledge this is the first one comparing such patients with the general population and also the first study on community living rather than institutionalised patients with intellectual disabilities.

Our analysis of predictors for vitamin D deficiency in people with intellectual disabilities confirmed well-known risk factors such as winter season, dark skin pigmentation and obesity, as well as impaired mobility.3 However, these predictors explained only 32% of the variance, and other unmeasured factors such as sun exposure, genetic variation22 and, to a limited extent, nutrition may play a role.

In particular, the extent of the use of ultraviolet-protection skin creams among patients with intellectual disabilities living in supervised care may be questioned. Such creams decrease the synthesis of vitamin D,23 so an excessive use may partly explain the 69% prevalence of vitamin D deficiency in the intellectual disabilities population in the summer months. We have no quantitative data to support our hypothesis, but carers regarded protection from sun burn as an important priority for their patients. Realisation of a need to educate carers in the use of sunscreens to allow sufficient production of vitamin D while avoiding sunburn could be a useful outcome from this study.

A previous study in patients with intellectual disabilities had shown no association between vitamin D deficiency and fractures.5 However, hypogonadism, a powerful risk factor for fractures, which is highly prevalent in the intellectual disabilities population (details available from the author on request and also see Jawli et al22), was not investigated. Our results, although based on a non-significant trend in a small number of patients, suggest that once other causes, such as catastrophic falls or long-term untreated hypogonadism, have been excluded vitamin D deficiency may play a role in the occurrence of fractures in people with intellectual disabilities. This interpretation is in line with the known causal role of vitamin D deficiency in osteomalacia and osteoporosis. It also accords with the decreased efficacy of bisphosphonates to improve bone mineral density and reduce fracture incidence unless vitamin D deficiency is corrected25 and with the clinical requirement that patients be vitamin D and calcium replete when starting fracture prevention treatment with any of the osteoporosis-specific drugs, which would otherwise be ineffective.26 Decreased bone mineral density is a predictor of fractures, so our demonstration of low bone mineral density in 70% of the vitamin D deficient patients in whom a DXA was performed, also supports a causal role of vitamin D deficiency in the pathogenesis of fractures in people with intellectual disabilities.

### Treatment

A high-dose intermittent regime, which included a maintenance dose of 50 000 IU colecalciferol monthly, achieved the optimal serum 25(OH)D concentration of ≥ 50 nmol/l more frequently (100% of the patients) than a daily dose of 800 IU colecalciferol.

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### Table 3  Mean serum 25(OH)D concentrations at baseline and after ≥ 3 months of treatment[^1^]

<table>
<thead>
<tr>
<th>D3 supplements</th>
<th>n</th>
<th>Baseline</th>
<th>≥ 3 months</th>
<th>Delta</th>
<th>P (≥ 3 months v. baseline)</th>
</tr>
</thead>
<tbody>
<tr>
<td>High-dose intermittent</td>
<td>28</td>
<td>20.0 (10.2)</td>
<td>86.2 (18.8)</td>
<td>68.0 (22.4)</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>800 IU daily</td>
<td>57</td>
<td>20.9 (9.8)</td>
<td>70.5 (20.7)</td>
<td>49.7 (21.0)</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>400 IU daily</td>
<td>5</td>
<td>26.9 (17.2)</td>
<td>41.8 (8.7)</td>
<td>14.9 (22.1)</td>
<td>0.2 (NS)</td>
</tr>
</tbody>
</table>

[^1^]: NS, not significant.

a. Delta 25(OH)D gives the mean (s.d.) increase in values between the two time points and is significantly different across the treatment groups (P = 0.001).
daily (80% of the patients). However, in the 20% of patients treated with 800 IU colecalciferol daily who did not achieve optimal serum vitamin D levels, the mean serum concentration obtained was 43 nmol/l, which is a level for which treatment is not considered mandatory according to UK recommendations.\(^3\)

Given that oral high-dose intermittent colecalciferol treatment regimens in the UK are only available via secondary care and that a standard preparation of 800 IU daily, available through primary care, achieves optimal serum 25(OH)D concentrations in 80% of patients, it seems reasonable to opt for the latter preparation in the vast majority of patients. It could also be argued that, as we showed the difference between serum 25(OH)D levels before and after supplementation with colecalciferol to be related to the dose of the supplements, a slightly higher dose than 800 IU daily might be able to normalise 25(OH)D levels in all patients. Therefore, colecalciferol preparations of 1000 IU, which are available over the counter, could also be considered.

A dose of 400 IU colecalciferol daily is the standard dose recommended by the Department of Health for population groups thought to be at risk of vitamin D deficiency, including those who have low or no exposure to the sun.\(^27\) Our findings, limited admittedly to only five patients, suggest this dose is unlikely to normalise 25(OH)D levels in people with intellectual disabilities, and that this recommendation requires further scrutiny and should be treated with caution as practical guidance.

**Limitations of the study**

Recruitment of patients with intellectual disabilities is challenging, so the sample size remained too small to allow a conclusive analysis of the relationship between vitamin D deficiency and fractures in the intellectual disabilities population. Choice of appropriate controls that reflect population norms is also difficult because volunteers to clinical studies tend to be drawn from more healthy subgroups. Nevertheless, 25(OH)D levels in our patients were low compared with laboratory values that do reflect the whole population. Finally, the findings apply mostly to people with intellectual disabilities of White ethnic background, as they represented 94% of our population. Given the association between skin colour and vitamin D deficiency and the limited data we obtained, the problem in many Black and minority ethnic people with intellectual disabilities is likely to be more marked.

**Implications of the study**

The results of our study show that vitamin D deficiency is highly prevalent in community-living patients with intellectual disabilities, and that it can be effectively, safely and inexpensively treated. This group should be actively screened for vitamin D deficiency because, although the condition is widespread, treatment of populations with normal vitamin D levels is unnecessary as it does not improve bone mineral density.\(^28\)

Concomitantly to correction of vitamin D deficiency, calcium intake should also be optimised by dietary improvements or addition of supplements as necessary, which is what we did in our good practice initiative and is recommended by current guidelines.\(^14\) The benefits that could in theory be gained by a simple intervention such as management of vitamin D deficiency are large, given the human cost of osteoporotic fractures, particularly in patients who are already disabled, and the direct cost of treating these fractures. These have been estimated, for hip fractures, at an average of £12,000 for hospital costs\(^29\) and £13,000 in the first 2 years for health and social after care following discharge for those who make a poor recovery.\(^30\)

In populations at high risk of osteoporosis, supplementation with colecalciferol at a median dose of 800 IU daily has been shown to reduce the risk of hip fracture by 30%.\(^31\) Our study was not designed, and would have been too small, to investigate an effect of treatment of vitamin D deficiency on fractures. A large study specifically on fracture prevention with vitamin D normalisation in people with intellectual disabilities is needed, particularly in view of the other factors for bone loss, for example physical inactivity, poor nutrition or hypogonadism, that are common in the intellectual disabilities population.

Our results relate to a special population but they may be relevant to other populations subject to poor health, with clinical or lifestyle characteristics similar to patients with intellectual disabilities. A substantial number of people with severe mental illness, particularly those with psychotic features, may be admitted to hospital or, because of negative symptoms, tend to live indoors. Evidence of vitamin D deficiency in these patients has already been published,\(^32–36\) and patients with schizophrenia have an increased risk of osteoporosis and fractures.\(^37\) Whether vitamin D deficiency should be added to the risk factors for osteoporosis, such as smoking,\(^38\) alcohol excess,\(^39\) sedentary lifestyle or hyperprolactinaemic hypogonadism,\(^40,41\) which are common in people for schizophrenia, seems very likely. As attention switches to the poor general medical outcomes of people with severe mental illness,\(^42\) a greater awareness of vitamin D deficiency as a measureable and treatable risk factor would be timely.

**Valeria Frighi, MD, Department of Psychiatry, University of Oxford and Oxfordshire Learning Disability NHS Trust (now Southern Health NHS Foundation Trust), Oxford; Allireza Morovat, PhD, FRCPPath, Department of Clinical Biochemistry, Oxford University Hospitals NHS Trust, Oxford; Matthew T. Stephenson, FRCPsych, MScPsych, Oxfordshire Learning Disability NHS Trust (now Southern Health NHS Foundation Trust), Oxford; Sarah J. White, RN, Oxford Centre for Diabetes, Endocrinology and Metabolism, University of Oxford, Oxford; Christina V. Hammond, BSc, Guy M. Goodwin, OBE, FRCPsych, FMedSci, Department of Psychiatry, University of Oxford, Oxford, UK**

**Correspondence:** Valeria Frighi, University of Oxford, Department of Psychiatry, Neurosciences Building, Warneford Hospital, Warneford Lane, Oxford OX3 7JX, UK. Email: valeria.frighi@psy.ox.ac.uk

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**References**

7. Srikanth R, Cassidy G, Joiner C, Tidshukuddy H. Osteoporosis in people with intellectual disabilities: a review and a brief study of risk factors for
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