



Celiac disease among women

Women are affected by celiac disease twice as much as men. Moreover, it is associated with many other health issues that clinicians must be vigilant for.

The articles published in this edition of the Forum on birth complications, infertility and osteoporosis in celiac sufferers all have one thing in common: the connection between celiac disease and women's health. There is much interest surrounding this topic, not least because dietary treatment of celiac disease makes it possible to prevent the occurrence of such complications. This has a positive effect on the sufferer from a medical, psychological and social point of view.

However, there are also many other connections between celiac disease and women's health, particularly since (a) this condition affects more women than men, with a ratio of around 2:1 between female and male sufferers; (b) certain autoimmune complications, especially Hashimoto's thyroiditis, even at a young age are already more common in girls than boys; (c) some variants of celiac disease, in particular iron-deficiency anemia, can complicate medi-

cal problems that more commonly affect women. As Khashan and McCarthy claim in their overview of obstetric complications, that there is "not yet sufficient evidence to recommend serological screening for celiac disease at the start of pregnancy". However, a simple blood test (to identify anti-transglutaminase antibodies) could without a doubt be carried out in addition to the many other routine tests performed for pregnant women, thereby preventing at an early stage many possible problems for both the expecting mother and the unborn baby.

Summary

The topics addressed in this edition of the Forum – birth complications and osteoporosis linked to celiac disease, combined with the more frequent occurrence of this condition in women – make celiac disease one of the topics which are of primary interest for women, especially those of childbearing age.



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Obstetric complications in women with celiac disease

Women suffering from celiac disease have a higher risk of complications during pregnancy, however these may be lowered by strict adherence to a gluten free diet.



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Celiac disease is a gluten sensitive enteropathy which has an estimated prevalence of 1% worldwide.¹ It is widely thought that this represents the tip of the iceberg and a large proportion of celiac disease remains undiagnosed. Celiac disease is most commonly identified in either early childhood or in the third or fourth decades of life.

Certain ethnic groups are thought to have an increased prevalence of celiac disease, in particular those of Celtic origin. Approximately 96% of patients with celiac disease express the HLA molecule DQ2, whereas the remainder mostly express the less com-

mon haplotype DQ8, reflecting the pivotal role of this molecule in the pathogenesis of celiac disease. Celiac disease is characterised by permanent intolerance to dietary gluten. Although traditionally considered a nutritional disorder of childhood presenting with varying degrees of malabsorption and diarrhea, it is now recognised as a systemic illness with multiple clinical presentations.² Celiac disease has been associated with an increased risk of many adverse pregnancy outcomes including unexplained infertility,³ miscarriage,³ congenital malformation,⁴ preterm birth,⁵ intrauterine growth restriction,^{5,6} postpartum hemorrhage,⁷ and assisted delivery.⁷ However, conflicting results exist.^{4,7,8}

Celiac disease, in particular untreated celiac disease, is thought to adversely affect pregnancy via antibodies which interact with the developing placental tissue resulting in adverse pregnancy outcomes. Anti-tissue transglutaminase (tTG) immunoglobulin G antibodies have been shown to bind to human trophoblast cells in vitro, resulting in

impaired trophoblast function in a dose and time dependent manner. In women with celiac disease gluten also induces a T-cell immune response which may also contribute to adverse pregnancy outcomes. In addition, gliadin itself can activate peripheral blood T-cells resulting in elevated cytokine secretion which may affect trophoblast development.

More recently, large cohort studies and systematic reviews have helped to clarify the degree of association between both treated and untreated celiac disease and adverse pregnancy outcomes, which has helped to provide clarity in the counselling and investigation of women with adverse pregnancy outcomes. The largest population-based studies on maternal celiac disease and adverse pregnancy outcomes used data from Sweden⁹ and Denmark⁵ and were published in the last decade. These studies benefited from the Medical Birth Register and Hospital Register in each country which allowed the identification of all births during the study period and whether the woman had ever been diagnosed with celiac disease. It was possible to determine whether the mother was diagnosed with celiac disease and whether the diagnosis took

Celiac-specific antibodies may interact with placental tissue to cause pregnancy complications.

place before or after pregnancy using dates of diagnosis and pregnancy. In the population-based Swedish study including more than 2 million babies, Ludvigsson and colleagues⁹ reported an association between undiagnosed maternal CD and low birthweight (OR=2.13), SGA (OR=1.62) and preterm birth (OR=1.71). Women with diagnosed

CD, who were presumably treated before giving birth, had no increased risk of adverse fetal outcomes compared with non-celiac women. These findings were later replicated in a population-based Danish study including more than 1.5 million babies.⁵ Women with undiagnosed celiac disease at the time of pregnancy had a higher risk of SGA (OR=1.3), preterm birth (OR=1.33) and smaller babies with birthweight reduced by 100 grams on average compared to women with diagnosed celiac disease. Similar to the Swedish study women with diagnosed, and presumably treated, celiac disease had no increased risk of adverse pregnancy outcome.

A recent systematic review included data from ten cohort studies which included data from over four and a half million women.¹⁰ This systematic review demonstrated that women with celiac disease (both treated and untreated) had a significantly higher risk of the development of preterm birth (adjusted OR 1.35), intrauterine growth restriction (OR 2.48), stillbirth (OR 4.84), low birthweight (OR 1.63), and small for gestational age defined as individualised birth centile under the tenth centile (OR 4.52). No significant differences were observed in the incidence of pre-eclampsia.

Subgroup analysis was then performed to examine the association between diagnosed (and assumed treated) celiac disease. The risk of preterm birth remained significantly higher both in the subgroup analysis of only women with diagnosed and treated celiac disease (OR 1.26) and in the subgroup analysis of only women with undiagnosed and untreated celiac disease (OR 2.50). Women with diagnosed and assumed treated celiac disease had a significantly lower risk of having a pregnancy complicated by preterm birth, compared with women with undiagnosed and untreated celiac disease (OR 0.80).

Undiagnosed maternal celiac disease is associated with low birthweight and preterm birth



Adherence to a gluten-free diet significantly reduces the risk of pregnancy complications.

Data regarding the prenatal and perinatal risk factors for the development of celiac disease in the offspring are also conflicting. However, it appears the biggest determinant of development of celiac disease in the offspring is the presence of maternal celiac disease.¹¹ The odds ratio from one mother and baby cohort of approximately one hundred thousand mother and baby pairs was approximately twelve for the development of offspring celiac disease.

What are the risks relevant to the general obstetric population and how can these be minimized?

Overall, women with celiac disease have an increased risk of adverse pregnancy outcomes. Treatment by means of a gluten free diet ameliorates this risk. As a result, women

with celiac disease should be advised to adhere to a strict gluten free diet pre-conceptually and during pregnancy to minimize any risks which may occur as a result of celiac disease. There remains insufficient evidence, both clinically and cost-effectively, to support a policy of screening healthy pregnant women for undiagnosed celiac disease at the start of pregnancy with the aim of improving pregnancy outcomes. Similarly, there is insufficient evidence to support the screening of women with adverse pregnancy outcomes for undiagnosed celiac disease. However, increasingly high-risk groups such as those with recurrent pregnancy loss are being screened for undiagnosed celiac disease. Non celiac pregnant women can be reassured that maternal consumption of gluten in pregnancy does not appear to be associated with an increased risk of development of celiac disease in the offspring.



REFERENCES

- 1 Green PH, Jabri B. Coeliac disease. *Lancet*. 2003;362(9381):383-91.
- 2 van Heel DA, West J. Recent advances in coeliac disease. *Gut*. 2006;55(7):1037-46.
- 3 Tersigni C, Castellani R, de Waure C, Fattorosi A, De Spirito M, Gasbarrini A, et al. Celiac disease and reproductive disorders: meta-analysis of epidemiologic associations and potential pathogenic mechanisms. *Hum Reprod Update*. 2014;20(4):582-93.
- 4 Ban L, West J, Abdul Sultan A, Dhalwani NN, Ludvigsson JF, Tata LJ. Limited risks of major congenital anomalies in children of mothers with coeliac disease: a population-based cohort study. *BJOG*. 2015;122(13):1833-41.
- 5 Khashan AS, Henriksen TB, Mortensen PB, McNamee R, McCarthy FP, Pedersen MG, et al. The impact of maternal celiac disease on birthweight and preterm birth: a Danish population-based cohort study. *Hum Reprod*. 2010;25(2):528-34.
- 6 McCarthy FP, Khashan AS, Quigley E, Shanahan F, O'Regan P, Cronin C, et al. Undiagnosed maternal celiac disease in pregnancy and an increased risk of fetal growth restriction. *J Clin Gastroenterol*. 2009;43(8):792-3.
- 7 Abdul Sultan A, Tata LJ, Fleming KM, Crooks CJ, Ludvigsson JF, Dhalwani NN, et al. Pregnancy complications and adverse birth outcomes among women with celiac disease: a population-based study from England. *Am J Gastroenterol*. 2014;109(10):1653-61.
- 8 Dhalwani NN, West J, Sultan AA, Ban L, Tata LJ. Women with celiac disease present with fertility problems no more often than women in the general population. *Gastroenterology*. 2014;147(6):1267-74 e1; quiz e13-4.
- 9 Ludvigsson J, Montgomery S, Ekbohm A. Celiac disease and risk of adverse fetal outcome: a population-based cohort study. *Gastroenterology* 2005; 129:454 – 463.
- 10 Saccone G, Berghella V, Sarno L, Maruotti GM, Cetin I, Greco L, et al. Celiac disease and obstetric complications: a systematic review and metaanalysis. *Am J Obstet Gynecol*. 2016;214(2):225-34.
- 11 Emilsson L, Magnus MC, Stordal K. Perinatal risk factors for development of celiac disease in children, based on the prospective Norwegian Mother and Child Cohort Study. *Clin Gastroenterol Hepatol*. 2015;13(5):921-7.

Gluten and female infertility



Celiac disease is known to affect fertility in women, however less is known regarding the relationship between gluten sensitivity and fertility issues. Even in the case of negative serology, there is evidence to suggest a possible role for the gluten free diet as an adjunct therapy in some patients.

Current guidelines do not recommend routine screening of women diagnosed with most types of infertility for celiac disease (CD), even though the research studies that CD can affect fertility. In women CD can delay puberty¹, cause malabsorption and many nutritional deficiencies such as zinc, B12, iron and folate². These nutrients are important for conception/pregnancy and low status has been implicated in both fertility and pregnancy problems. CD is also linked to amenorrhea, premature ovarian failure and obstetric complications such as pre-term birth and low birth weight.³ There are, however, reports of successful pregnancy outcomes

with unexplained fertility appear to have higher rates of CD compared to the general population.⁶ In one study women with unexplained infertility had a six times higher odds of having CD than controls.⁶ Given the financial cost and emotional impact of fertility treatment, health professionals working with such patients should be encouraged to screen for CD particularly in women diagnosed with infertility, especially unexplained infertility.

While it is recognized that there is an association between CD and fertility problems there is little information about non-celiac gluten



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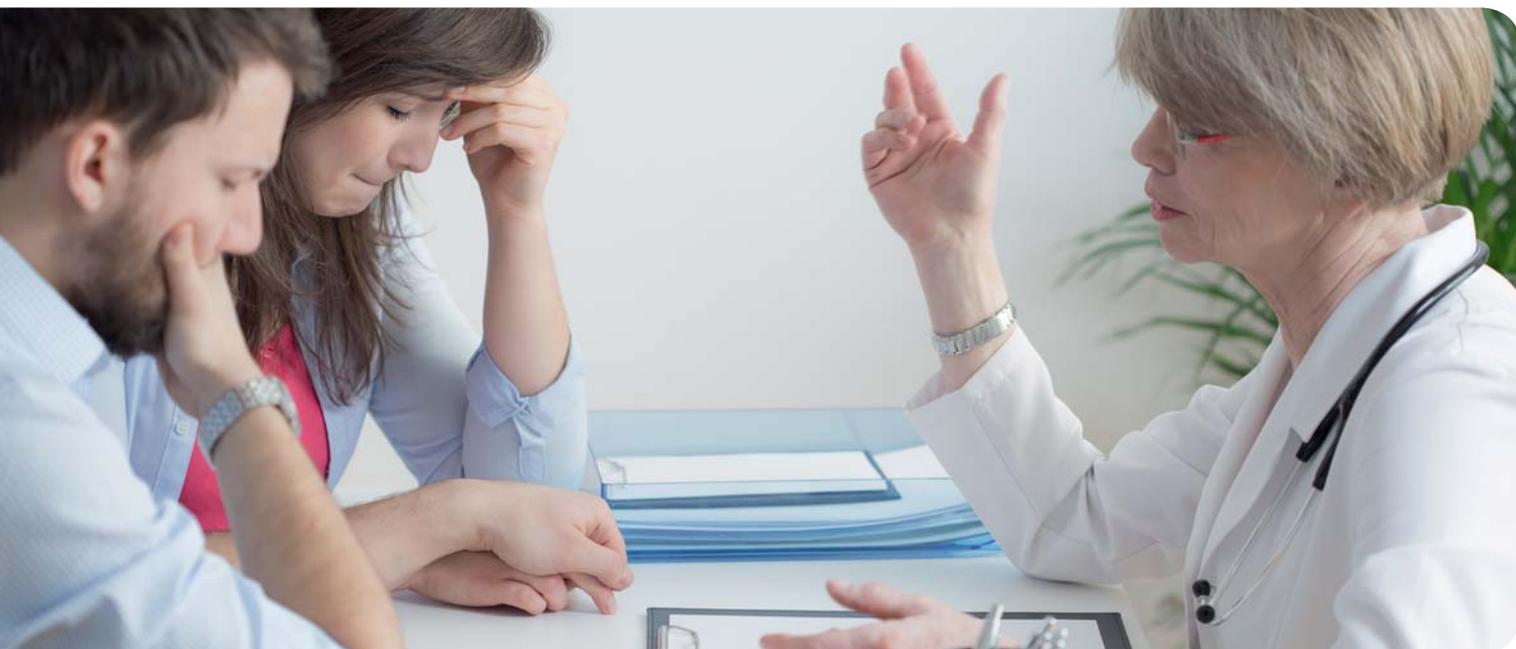
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The prevalence of celiac disease is estimated to be 4 to 8 times higher in women with fertility problems than in the general population.

for women with CD who have a history of miscarriage once a gluten free diet (GFD) has been introduced.⁴ Undiagnosed and untreated celiac disease in particular can have far reaching effects on both female fertility and pregnancy outcomes.

Reports on the prevalence of CD in women with infertility are between 4 and 8% in Europe; this is higher than the general population where it is estimated at 1%.⁵ Women





A gluten free diet may alleviate the painful symptoms of endometriosis.

sensitivity (NCGS) and infertility, though a case report on a possible association was published in 2015.⁷ Interestingly, iron, folic acid, vitamin D and B12 deficiency have also been documented in NCGS.^{8,9} Therefore the possibility of malabsorption of nutrients key to fertility remains a possibility in NCGS along with immunological abnormalities, which have also been reported.⁷

An association between CD and endometriosis has also been documented.¹⁰ Endometriosis affects around two million women in the UK¹¹ and is one of the leading causes of female infertility. It is also interesting that endometriosis has been reported as the primary symptom of CD when classical symptoms

were absent,¹² and that in many patients gastro-intestinal symptoms and irritable bowel syndrome (IBS) are frequently seen alongside endometriosis¹³. It is also important to remember some patients with IBS are gluten sensitive, as the literature documents both associations between IBS and CD¹⁴ and IBS and NCGS¹⁵, though this area and underpinning pathogenesis are still being investigated.

A postgraduate student research project currently in progress at the University of Worcester exploring motivations of people without CD for following a GFD has ascertained many do so in an attempt to self manage symptoms. This is interesting given the widely held view that GFD is increasing in popularity among the non-celiac patients. Could it be that some are self-managing women's health issues with a GFD?

There are few studies exploring the potential therapeutic effect of a GFD in supporting female health conditions such as endometriosis. However, a study undertaken in Italy in 2012¹⁶ reported that the painful symp-



toms of endometriosis reduced after a year on a GFD. Two-hundred-and-seven patients were studied, all were diagnosed with endometriosis and after a year, 75 per cent of the patients reported a statistically-significant reduction in symptoms. However, 25 per cent reported no improvement of symptoms though no patients experienced a worsening of pain. Improved scores were also reported in all patients for general health perception, vitality, mental health and social and physical functioning.

Professionals working with women diagnosed with unexplained fertility and other women's health issues such as endometriosis should assess clinical symptoms, co-morbidities and consider screening for CD, whilst being mindful that there may be no digestive symp-

toms present. Health professionals should, therefore, also consider NCGS when CD serology is negative and there are no indications for biopsy. Even though further research in this area is undoubtedly needed, given the emotional and financial impacts of infertility, a gluten free diet could be discussed with patients and perhaps considered as an adjunct to other treatments to address infertility or support conditions such as endometriosis.

REFERENCES

- 1 Leffler DA, Green PH, Fasano A. Extraintestinal manifestations of coeliac disease. *Nat Rev Gastroenterol Hepatol*. 2015 Oct;12(10):561-71.
- 2 Vici G, Belli L, Biondi M, Polzonetti V. Gluten free diet and nutrient deficiencies: A review. *Clin Nutr*. 2016 May 7.
- 3 Bykova SV, Sabel'nikova EA, Parfenov AI, Gudkova RB, Krums LM, Chikunova BZ. Reproductive disorders in women with celiac disease. Effect of the etiotropic therapy. *Eksp Klin Gastroenterol*. 2011;(3):12-8.
- 4 Tursi A, Giorgetti G, Brandimarte G, Elisei W. Effect of gluten-free diet on pregnancy outcome in celiac disease patients with recurrent miscarriages. *Dig Dis Sci*. 2008 Nov; 53(11):2925-8.
- 5 Fortunato F, Martinelli D, Prato R, Pedalino B. Results from Ad Hoc and Routinely Collected Data among Celiac Women with Infertility or Pregnancy Related Disorders: Italy, 2001–2011. *The Scientific World Journal*. Volume 2014.
- 6 Singh P, Arora S, Lal S, Strand TA, Makharia GK. Celiac Disease in Women With Infertility: A MetaAnalysis. *J Clin Gastroenterol*. 2016 Jan; 50(1):339.
- 7 Bold J, Rostami K. Non-coeliac gluten sensitivity and reproductive disorders. *Gastroenterol Hepatol Bed Bench* 2015;8(4):294-297.
- 8 Volta U, Bardella MT, Calabrò A, Troncone R, Corazza GR; Study Group for Non-Celiac Gluten Sensitivity. An Italian prospective multicenter survey on patients suspected of having non-celiac gluten sensitivity. *BMC Med* 2014;12:85.
- 9 Molina-Infante J, Santolaria S, Sanders DS, Fernández-Bañares F. Systematic review: non-coeliac gluten sensitivity. *Aliment Pharmacol Ther* 2015;41:807-20.
- 10 Stephansson O, Falconer H, Ludvigsson J. Risk of endometriosis in 11,000 women with celiac disease. *Hum Reprod*. 2011 Oct;26(10):2896-901.
- 11 Adamson G, Kennedy S, Hummelshoj L. (2010) Creating solutions in endometriosis: global collaboration through World Endometriosis Research Foundation. *Journal of Endometriosis* 2, 13–16.
- 12 Caserta D, Matteucci E, Ralli E, Bordi G, Moscarini M. Celiac disease and endometriosis: an insidious and worrisome association hard to diagnose: a case report. *Clin Exp Obstet Gynecol*. 2014;41(3):3468.
- 13 Ek M, Roth B, Ekström P, Valentin L, Bengtsson M, Ohlsson B. Gastrointestinal symptoms among endometriosis patients-A case-cohort study. *BMC Womens Health*. 2015 Aug 13;15:59.
- 14 SánchezVargas LA, ThomasDupont P, TorresAguilera M, AzamarJacome AA, RamírezCeervanes KL, AedoGarcés MR, MeixueiroDaza A, RoeschDietlen F, GrubePagola P, VivancoCid H, Remes Troche JM. Prevalence of celiac disease and related antibodies in patients diagnosed with irritable bowel syndrome according to the Rome III criteria. A case control study. *Neurogastroenterol Motil*. 2016 Jul;28(7):9941000.
- 15 Makharia A, Catassi C, Makharia GK. The Overlap between Irritable Bowel Syndrome and Non-Celiac Gluten Sensitivity: A Clinical Dilemma. *Nutrients*. 2015 Dec 10;7(12):1041726.
- 16 Marziali M, Venza M, Lazzaro S, Lazzaro A, Micossi C, Stolfi VM. Gluten-free diet: a new strategy for management of painful endometriosis related symptoms? *Minerva Chir*. 2012 Dec;67(6):763499504.

Osteoporosis in the celiac population

Celiac disease increases the risk of osteoporosis due to calcium malabsorption, secondary to villous atrophy. Conversely, a calcium-enriched gluten free diet can significantly increase bone mineral density in celiac patients.



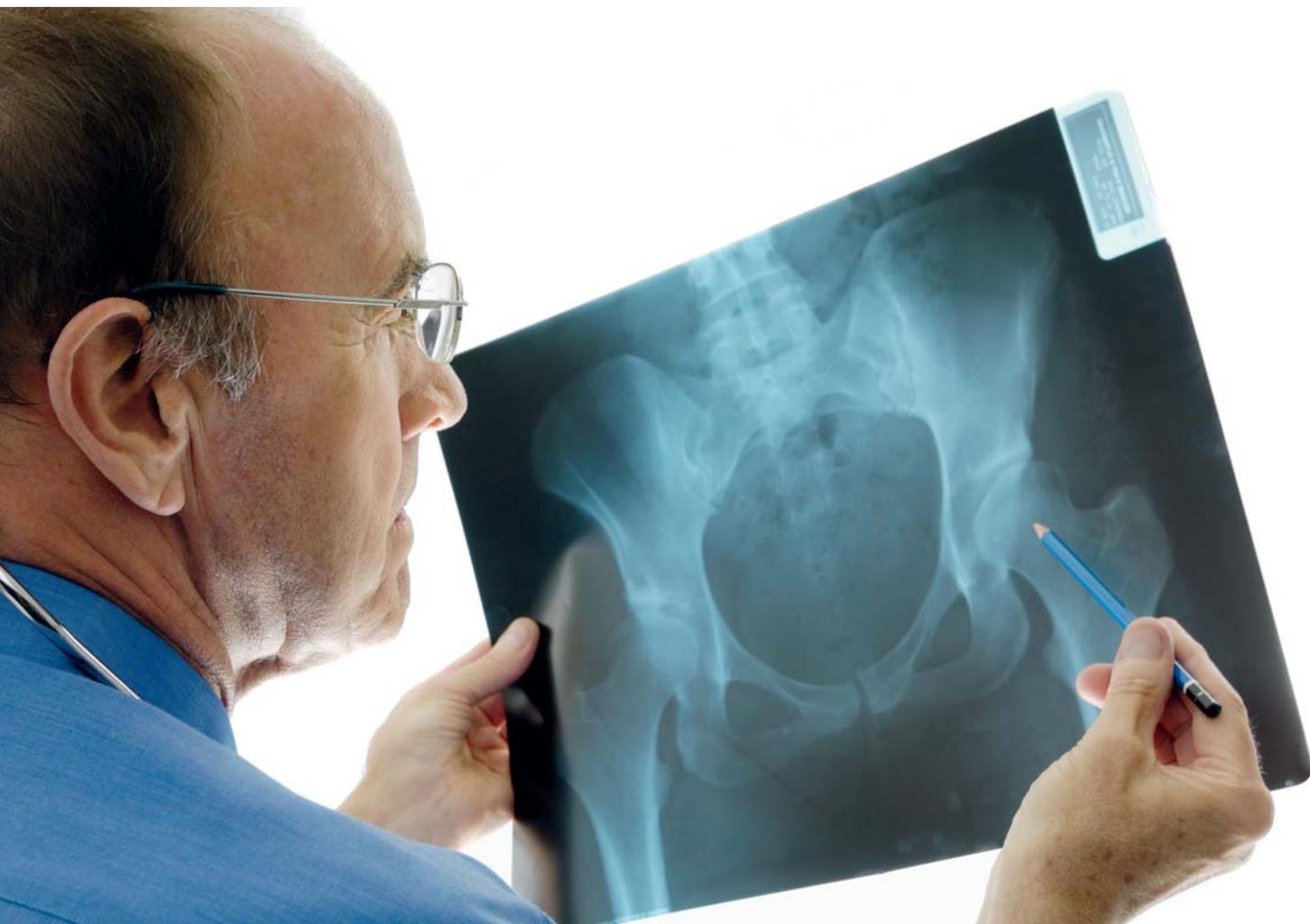
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Background

The impact of osteoporosis, and other bone diseases, on health and well-being is not well recognized. Currently, worldwide, osteoporosis causes in excess of 8.9 million fractures a year, this translates to 1 fracture every 3 seconds¹ and the prevalence in the UK is estimated at 3 million (Age UK; NOS, 2016). The rates of disabilities directly or indirectly associated with osteoporosis are higher than cancer¹ and once disease related fractures occur there is an 86% risk of further fracture leading to

a risk of continued, and potentially escalating, risk of long term osteoporotic disability and associated co-morbidities such as chronic pain.² In 2010, osteoporosis was the cause of death of 43,000 Europeans³ and the cost of treating osteoporosis in the EU in that year, including "medicines management", fracture treatment, and hospital stays, amounted to 37 billion euros. The burden for the population with osteoporosis was 1,180,000 lost quality-adjusted life years (QALYS), this number is estimated to increase by 20% between 2010 and 2025.³



Etiology

Osteoporosis occurs due to either a reduced peak bone mass or increased bone loss. Bone mass usually peaks around age 35, however a number of conditions, for example anorexia, malnutrition, Crohn's disease, celiac disease or alcoholism can reduce the peak bone mass achieved. Increased bone loss is associated with decreased sex hormone synthesis, particularly in women, malnutrition, hyperthyroidism, and certain medications; for example anti-seizure medications, chemotherapy, proton pump inhibitors, and steroids. Lifestyle can also be a factor with behaviors such as using sunscreen, smoking, and lack of exercise, all increasing risk of bone loss and, ultimately, bone diseases such as osteoporosis.⁴ Genetics, ethnicity, and age are also all risk factors for osteoporosis; for example, incidence of osteoporosis increases to 1 in 3 women and 1 in 5 men over 50 and in members of the same family.^{4,5} None of the factors mentioned here are mutually exclusive.

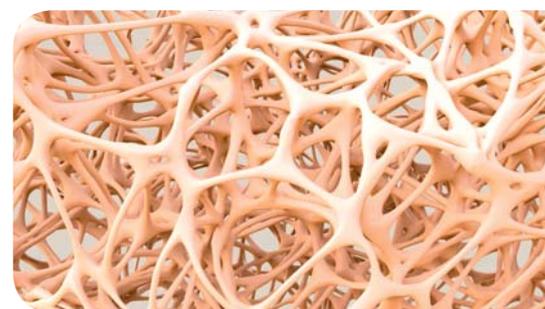
Osteoporosis and Celiac Disease

Increased levels of low bone mineral density, osteopenia, and osteoporosis have been frequently identified in people with celiac disease when compared to the general population.^{6,7,8,9} Estimates of fracture risk in the celiac population have varied, however a recent meta analysis found a 30% increased risk of fracture at baseline in comparison to controls (95% confidence interval [CI]: 1.14,

1.50) and 69% increase in the risk of hip fracture (95% CI: 1.10, 2.59) in comparison to controls.¹⁰ The meta analysis was based on observational and epidemiological studies and is therefore unable to demonstrate any causal inferences. It should also be noted that most of the subjects in the studies examined were reported to be following a gluten free diet which may have had an impact on the results.

Potential mechanisms for changes in bone metabolism and, therefore bone health, in celiac disease have been postulated over time. It is well established that the villous atrophy observed in celiac disease leads to the malabsorption of many micronutrients, for example, calcium and vitamin D. Both these micronutrients have essential roles in maintaining bone health and the increased risk of osteoporosis in people with celiac disease has been attributed to calcium malabsorption, stimulating release of parathyroid hormone which increases bone turnover and leads to increased bone loss.¹¹ Current research is also investigating the role of increased activation of inflammatory cytokines and autoimmune factors as potential mechanisms for altered bone metabolism. Further studies are needed to fully elucidate the exact nature and balance of the mechanisms involved.

Recent studies have analysed physiological changes associated with celiac disease in conjunction with bone health, with some interesting findings. In a largely female cohort, Garcia-Manzanares and colleagues (2012) found positive correlations between increased villous



Low bone mineral density is frequently observed in people with celiac disease.

Calcium

atrophy and loss of bone mass on the lumbar spine.⁸ In a female only population, Stein et al (2015) found significant differences in bone microarchitecture in participants with celiac disease compared to healthy age matched controls.⁹ These differences were principally found in trabecular rather than cortical bone and were associated with reduced performance in measures of skeletal strength. Stein et al (2015) also noted that despite a substantially increased calcium intake compared to the age matched controls, serum calcium levels were lower in the celiac group.⁹ This is suggestive of calcium malabsorption and consistent with long-held beliefs about the mechanisms for increased osteoporosis in the celiac population.

Further studies looking at larger groups of both male and females with celiac disease are needed to establish the relevance and applicability of these studies to the larger celiac population.

Diagnosis

Osteoporosis is most commonly diagnosed by using dual-energy X-ray absorptiometry (DEXA) to measure bone density. Other conventional radiotherapy techniques such as quantitative computed tomography (QCT) can also be used.¹² The measured bone mineral density is compared to standardized results from the general population using a T score; a T score of 2.5 standard deviations, or more, lower than the standard is defined as osteoporosis. The techniques for diagnosing for the celiac population are the same as those used for the general population. Standard recommendations for the diagnosis of celiac disease continue to be serology and duodenal biopsy;

this recommendation takes into account the variability that can be found with serology and the fact that ultimately the biopsy reveals if malabsorption is present.¹³ This recommendation does not apply to diagnosis in pediatric patients.¹³ The biopsy, if performed, informs not only on the presence of villous atrophy of the small intestine but also on degrees of change seen, usually measured using the Marsh scale¹³; a high Marsh score correlates with a high degree of villous atrophy and is associated with a high degree of malabsorption.^{8,13} Repeat biopsies, when following a gluten free diet, are used to confirm continued atrophy of the villi and likelihood of continued malabsorption.

Treatment

Current management of osteoporosis is focused on medical therapy such as bisphosphonates, hormone replacement therapy, recombinant parathyroid therapy and, supplements such as calcium and calcium combined with vitamin D.

However diet and lifestyle also have a role to play in both prevention and treatment of osteoporosis, including osteoporosis in those with celiac disease. The British Society of Gastroenterology 2014 guidelines for the management of celiac disease emphasise the importance not only of a gluten free diet, to reduce villous atrophy, promote mucosal healing and lead to enhanced nutrient absorption, but also to ensure an adequate intake of a range of nutrients, including calcium.¹³ Adequate ingestion of dietary calcium can reduce the need for calcium supplements, and some people with celiac disease can achieve the recommended

intake of 1000 mg/day from diet alone¹⁴, although very few achieve the previously recommended intake of 1500 mg/day¹⁴, which is still the recommended intake for post-menopausal women and elderly men¹³. The prevalence of osteoporosis within the celiac population emphasises the importance of a consistent and adequate calcium intake.

To conclude, current evidence suggests that preventative measures are essential for all those diagnosed or at risk of osteoporosis, whether they have celiac disease or not. These

measures include ensuring an adequate intake of a wide range of nutrients to optimise health, maintain an active lifestyle and avoid malnutrition. For those at increased risk, for example people with celiac disease, calcium intake should meet the 1000 mg/day recommended guidelines¹³. Supplements should be used if intake is insufficient from dietary sources.



REFERENCES

- 1 Johnell O and Kanis JA An estimate of the worldwide prevalence and disability associated with osteoporotic fractures. *Osteoporos Int* (2006) 17:1726.
- 2 Kanis JA, Johnell O, De Laet C, et al. (2004) A meta-analysis of previous fracture and subsequent fracture risk. *Bone* 35:375.
- 3 Hernlund E, Svedbom A, Ivergard M, Compston J. Osteoporosis in the European Union: Medical Management, Epidemiology and Economic Burden. A report prepared in collaboration with the International Osteoporosis Foundation (IOF) and the European Federation of Pharmaceutical Industry Associations (EFPIA). *Arch Osteoporos* (2013) 8:136 DOI 10.1007/s11657-013-0136-1
- 4 Hendrickx, G., Boudin, E. and Van Hul, W. A look behind the scenes: the risk and pathogenesis of primary osteoporosis. *Nature Reviews Rheumatology* (2015), 11, 462–474 doi:10.1038/nrrheum.2015.48
- 5 Van Staa TP, Dennison EM, Leufkens HG, Cooper C. Epidemiology of fractures in England and Wales. *Bone* (2001) 29:517
- 6 Van Heel, D.A. & West, J. Recent advances in coeliac disease. *GUT* (2006), 55, 1037-1046.
- 7 Corazza, G.R., Di Sario, A., Cecchetti, L., et al. Influence of pattern of clinical presentation and of gluten-free diet on bone mass and metabolism in adult coeliac disease. *Bone*, (1996) 18(6), 525-530.
- 8 García-Manzanares A, Tenias JM, Lucendo AJ. Bone mineral density directly correlates with duodenal Marsh stage in newly diagnosed adult celiac patients. *Scand J Gastroenterol.* (2012) 47(8-9):927-36. doi: 10.3109/00365521.2012.688217
- 9 Stein, E.M., Rogers, H., Leib, A., McMahon, D.J., Young, P., Nishiyama, K., Guo, X.E., Lewis, S, Green, P.H., Shane, E. Abnormal Skeletal Strength and Microarchitecture in women with Celiac Disease. *J. Clin Endocrinol Metab*, (2015) 100(6), 2347-2353
- 10 Heikkila K, Pearce J, Maki M, Kaukinen K. Coeliac disease and bone fractures: a systematic review and meta-analysis. *J Clin Endocrinol Metab.* 2015;100(1),25–34
- 11 Walters JRF. Bone mineral density in coeliac disease. *Gut* (1994) 35,150–1
- 12 <https://www.iofbonehealth.org/diagnosing-osteoporosis> accessed 10/8/16 at 2.13
- 13 Ludvigsson, J.F., Bai, J.C., Biagi, F., et al. Diagnosis and management of adult coeliac disease: guidelines from the British Society of Gastroenterology, *GUT* (2014) 63(8). 1210-1228.
- 14 Martin, K. and Woodall, A. Optimising the Management of Bone Disease for Coeliac Patients in a Dietetic-led Clinic. *International Journal of Celiac Disease.* 2016, 4(2), 48-54. DOI: 10.12691/ijcd-4-2-6

News

Virtual Tour of Dr. Schar Plant

In the February issue of Dr. Schar Institute (DSI) Newsletter, we talked about the tour of our gluten free plant for Retail Dietitians. Since we've received many requests from other Dietitians to tour our plant in South Jersey. For out-of-state Dietitians, we're currently working on a virtual tour of Dr. Schar plant. This tour would allow Dietitians in other states to experience firsthand how Schar gluten free products are made and all the precautions we take to prevent gluten cross-contamination.

Either in-person or virtual, the tour will allow us to introduce Dr. Schar, the company, to those who know our products but haven't heard yet the story behind the brand. Dr. Schar was an actual Medical Doctor who started the company in Europe in the 20's. He worked closely with other medical doctors and hospitals to provide patients with allergen-free products. In the 80's, we decided to focus our efforts on gluten free foods. At all Dr. Schar facilities across the world, we solely make gluten free products. This allows us to fully control the production process and prevent cross-contamination with gluten and other allergens. We have our own farmers growing gluten free grains for us. We continuously test all raw in-

gredients entering our facilities, all our products throughout the entire production process, and even our packaging for gluten.

We also try to meet the demands of our consumers and healthcare professionals for products that are not only gluten but also dairy, egg, and nut free. Some of Schar products are also soy or corn free. We listen to our consumers, either directly or via Dietitians who work with gluten free patients, and try to accommodate their needs whenever possible. We welcome new ideas, and any feedback we may receive is forwarded to our R&D team in Italy.

Dietitians are the first line of defense when it comes to educating patients about celiac and non-celiac gluten sensitivity. Recently, many healthcare professionals also started recommending the gluten-free diet for patients with IBS and autism.

For more on different conditions related to gluten and other free resources, please register on our website solely dedicated to for healthcare professionals working with gluten free patients.



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