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Special Article

# Chronic pain: a review of its epidemiology and associated factors in population-based studies

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## **Summary**

Chronic pain is a common, complex, and distressing problem that has a profound impact on individuals and society. It frequently presents as a result of a disease or an injury; however, it is not merely an accompanying symptom, but rather a separate condition in its own right, with its own medical definition and taxonomy. Studying the distribution and determinants of chronic pain allows us to understand and manage the problem at the individual and population levels. Targeted and appropriate prevention and management strategies need to take into account the biological, psychological, socio-demographic, and lifestyle determinants and outcomes of pain. We present a narrative review of the current understanding of these factors.

Keywords: chronic pain; epidemiology; genetics; incidence; prevalence; risk factors

Chronic pain is a common, complex, and distressing problem, which has a significant impact on society and individuals.  $^1$  It commonly presents as a result of an injury or a disease; however, it is a separate condition in its own right, not merely an accompanying symptom of other ailments. Chronic pain, therefore, has both its own taxonomy and medical definition.  $^{2-4}$ 

The Global Burden of Disease Study 2016 reaffirmed that the high prominence of pain and pain-related diseases is the leading cause of disability and disease burden globally. Worldwide, the burden caused by chronic pain is escalating: 1.9 billion people were found to be affected by recurrent tension-type headaches, which were the most common symptomatic chronic condition. Measuring years lived with disability, low back and neck pain have consistently been the leading causes of disability internationally, with other chronic pain conditions featuring prominently in the top 10 causes of disability.

In order to develop treatment plans and prevention strategies, chronic pain needs to be understood in the context of social, biological, psychological, and physical factors. This is a narrative synthesis of the epidemiology, particularly the risk factors and demographic associations, of chronic pain.

# Importance of epidemiology in chronic pain

Epidemiology, the 'study of the distribution and determinants of health-related states or events in specified populations and the applications of this study to control health problems', 6 is vital to understanding chronic pain. According to the International Association for the Study of Pain, chronic pain is 'pain which has persisted beyond normal tissue healing time', 7 which, in the absence of other factors, is generally taken to be 3 months. 7 There are many risk factors for chronic pain, including socio-demographic, psychological, clinical, and

biological factors. Insight into these risk factors will allow the identification of specific prevention and management approaches that take into account these predisposing factors and the impact of pain on those affected.

Chronic pain, like most diseases, often arises from a series or combination of multiple events. Even when there is a solitary precipitating event in the genesis of chronic pain (e.g. injury), there remains a series of factors that affect the duration, intensity, and effects (physical, psychological, social, and emotional) of chronic pain.9 Health-related behaviours and their outcomes are the most important modifiable risk factors in the genesis, duration, and impact of chronic pain. 10

## **Methods**

For this narrative review, we were informed by studies that investigated or demonstrated an association between any biological, psychological, socio-economic, cultural, or other risk factor for the development, persistence, or severity of chronic pain. In our search for these, we applied no limits for date of publication, country of origin, or study design. Articles published in languages other than English and abstracts for which no full text was available were excluded.

We searched six electronic databases (Medline, Embase, Cochrane Central Register of Controlled Trials, PubMed, Cochrane Database of Systematic Reviews, and Web of Science) using the following terms:

- (i) Chronic pain or persistent pain AND
- (ii) Epidemiology OR demography OR demographics OR incidence OR prevalence OR frequency OR pattern OR determinants OR causes OR risk factors OR public health

Databases were searched up to September 1, 2018. Through this search, we identified the main themes and important studies that have contributed to them. Under each of these themes, we further searched for systematic reviews. Additional records were identified through searching references of included articles and through literature searches of publications by leading academics working in the field of chronic pain research. Factors identified through this search were then search for specifically in order to identify further relevant publications.

## Prevalence and incidence

Prevalence is the proportion of the at-risk population affected by a condition. Population estimates for the prevalence of chronic pain vary widely according to case definition and ascertainment methods, and time place and population. Research suggests that chronic pain affects 13-50% of adults in the UK. 1,11 Of those who live with chronic pain, 10.4–14.3% were found to have moderate-to-severe disabling chronic

Incidence is the number of new cases of a disease developing during a particular time period in a population at risk of developing the disease. Although difficult to accurately determine because of a lack of longitudinal studies, the incidence of chronic pain in one region of the UK has been estimated at 8% per year. 12

# Factors associated with chronic pain

Chronic pain has multiple associated physical, psychological, and social factors (Table 1). Historically, risk factors have been

Table 1 Factors associated with the development of chronic pain

Factor	
Demographic	Age
	Gender
	Ethnicity and cultural background
	Socio-economic background
	Employment status and occupational factors
Lifestyle and	Smoking
behaviour	Alcohol
	Physical activity
	Nutrition
	Sunshine and vitamin D
Clinical	Pain
	Multi-morbidity and mortality
	Mental health
	Surgical and medical interventions
	Weight
	Sleep disorders
	Genetics
Other	Attitudes and beliefs about pain
	History of violent injury, abuse, or
	interpersonal violence

classified as 'modifiable' and 'non-modifiable'; however, this bio-medico-centric approach to epidemiology does not always take into account the complex interplay between both modifiable and non-modifiable elements of each risk factor. For example, the experience of violence or abuse in the past is often considered 'non-modifiable' because the event or events have already occurred, and the patient's history cannot be changed. However, the individual's interpretation of these events and their impact on their life and health is constantly evolving and will affect the impact that this has on their future health and life. Furthermore, interventions that improve the biopsychosocial determinants of health can prevent or reduce the future population exposure to risk factors implicated in the genesis of chronic pain.

Ascertaining and managing those risk factors and elements of risk factors that are modifiable may allow the prevention of chronic pain, or the reduction of its duration and severity. Factors that are not amenable to medical interventions are still important to discuss in the clinical context. Many are relevant to chronic pain predictions, assessments, management, and prognoses, and others will be important in the attempt to identify new targets for therapeutic intervention.8 An epidemiologically informed multidisciplinary and patient-centred approach is key to the successful management of chronic pain. 13,14

## Demographic

Age

Whilst there is a paucity of evidence examining chronic pain in children and adolescents, 15 the available literature suggests that older patients have a higher prevalence of chronic pain than younger groups of patients. 1 Advanced age and chronic pain (and its reporting) have a complex interrelationship, 16-18 whereby multi-morbidity is independently associated with chronic pain. 19 With increasing age comes increasing multi-

morbidity; the more advanced a patient's age, the more he or she is likely to have experienced noxious stimuli or injury that can trigger chronic pain. For example, in people with shingles, those aged 50-54 yr have an 8% chance of developing postherpetic neuralgia, whereas those aged 80-84 yr have a 21% chance of its development.<sup>20</sup> Assessing pain in older patients can be complex, particularly because older adults are often reticent to discuss or disclose the level of their pain.<sup>21</sup> Agerelated disease processes, such as cognitive decline and dementia, can make identifying and managing chronic pain more difficult.<sup>21</sup> Characteristics of pain, including duration and severity, area of body affected, and number of sites of pain, were found to be indicators for ongoing pain in older women, but these were less relevant in older men.<sup>22</sup> Chronic pain is not limited to older age groups, however. A study from across 42 countries identified that self-reported chronic pain amongst adolescent populations was common: 20.6% of young people experienced pain in at least two sites of headache, stomach, and backache.<sup>23</sup> Chronic pain affects up to 30% of those aged 18-39 yr. Younger age seems to be a risk factor for chronic post-surgical pain.<sup>24</sup>

## Gender

Men are less likely to report or experience chronic pain than women,<sup>25</sup> and girls are more likely to report pain in multiple sites than boys.<sup>23</sup> Several reviews have studied how gender (role) and sex (biological) differences are related to the way men and women experience pain. <sup>26–29</sup> One recent systematic review found that women who experience pain are more likely to use maladaptive coping strategies, which predispose them to chronic pain and poorer functional ability.<sup>26</sup> Women have been shown to have lower pain thresholds and tolerance, and are more likely to experience greater intensity and unpleasantness with pain.<sup>30</sup> The evidence also suggests that women have different sensitivities to analgesia.<sup>27</sup> When corrections are made for the prevalence of pain in the different genders, women are more likely to seek treatment for their pain. In a recent study from one specialist pain clinic, there were twice as many women as men. 31 Women reported a higher level of pain intensity and higher pain-related disability than men.<sup>32</sup> Although there is insufficient information on the mechanisms behind these sex-specific differences in pain perception and pain prevalence, <sup>26</sup> there is some evidence for the role of oestrogens<sup>33</sup> and genetics, including sex-specific differences in the contribution of pain-related genes.<sup>34</sup>

## Ethnicity and cultural background

There are substantial and complex ethnic variations in prevalence and outcomes of pain-related conditions, although the mechanisms behind these remain poorly understood. 11,35 Caucasian patients have been found to experience less pain and less pain-related disability than black patients.<sup>35,36</sup> A survey of 500 000 people in the UK showed that those who self-identified as white were less likely to report chronic pain than those reporting black, Asian, or mixed ethnicity.<sup>37</sup> However, once adjusted for income employment and adverse life events, the association between self-reported ethnicity and chronic pain was significantly attenuated. The prevalence of chronic pain and its associated disability has been found to be greater in developing countries than in developed countries.<sup>5</sup>

## Socio-economic background

Population studies reliably show that the prevalence of chronic pain is inversely related to socio-economic factors  $^{36,38-40}$  Those who are socio-economically deprived are not only more likely to experience chronic pain than people from more affluent areas, but they are also more likely to experience more severe pain and a greater level of painrelated disability. 36,41,42 People who have low levels of education, perceived income inequalities, and high levels of neighbourhood deprivation are more likely to experience chronic pain than those who have higher levels of education, less perceived income inequality, and who live in more affluent neighbourhoods.<sup>43</sup> The economic impact of chronic pain compounds the cycle of pain's complex interrelationship with socio-economic deprivation. 40 Although an individual's socioeconomic and educational background are non-modifiable, it is clear that political attention to these factors can have a great influence on the future prevalence and severity of chronic pain at a societal level.

## Employment status and occupational factors

People who are not in employment because of ill health or disability are more likely to have chronic pain than those who are employed. Occupational risk factors for chronic pain include poor job control, expectations of return to work (including fear around a recurring injury), 44 lack of work autonomy or the ability to modify work, job satisfaction, and higher perceived level of difficulty of job requirements.<sup>44,45</sup> Chronic neck and shoulder pain was found to be an independent predictor of chronic pain and related to work stress.<sup>46</sup> Non-manual workers were less likely to report chronic pain than people who hold manual occupations.<sup>47</sup> A recent study found that, amongst those who earned less than £18,000  $yr^{-1}$ , the prevalence of chronic pain was 52.5%, whilst amongst those with incomes greater that £100,000  $\mathrm{yr}^{-1}$  the prevalence was 33.5%.<sup>37</sup> The same study also demonstrated that chronic pain relates to working status: chronic pain was present in 78.9% of those who were unemployed, but only in 39.8% of those in paid employment and 42.4% of those in voluntary or unpaid work.<sup>37</sup> This relationship, however, may be bidirectional in that people with chronic pain may be less likely to be in work because of their pain. The extent of the burden on healthcare from chronic pain has also been related to education level and socio-economic level of patients.<sup>30</sup> A score to assess work disability from pain leading to absent days demonstrated that, in both genders, more locations of pain, smoking, less education, and obesity were related to higher risk of time off work as a result of pain.<sup>48</sup>

# Lifestyle and behaviour

# Smoking

People with chronic pain are more likely to smoke than those with no pain. 49-51 Patients who are heavy smokers report higher pain intensity scores than non-smokers, and report a higher number of painful sites. 52-54 Smoking is involved in the aetiology of several conditions that cause chronic pain,<sup>55</sup> and the relationship between smoking and chronic pain appears to be dose related.<sup>53</sup> Smokers affected by chronic pain are more likely to be dependent on tobacco, smoke more cigarettes a day, and have more difficulty in quitting smoking than those who do not have the condition.<sup>52</sup>

#### Alcohol

The analgesic properties of alcohol are short lived; however, it is commonly used by patients to 'self-medicate' for chronic pain. 57,58 When alcohol is used to excess, people with chronic pain can develop resistance to its limited analgesic effects. Additionally, alcohol withdrawal can increase pain sensitivity, which can promote a cycle of escalating alcohol abuse in order to seek analgesia effects at higher doses as tolerance builds, and to avoid alcohol withdrawal-related pain. 59,60 There is evidence that alcohol dependence may be caused by the same dysregulation of pain neurocircuitry and neurochemistry that causes chronic pain.60

## Physical activity

Systematic reviews conclude that exercise and physical activity have positive effects in chronic pain, with improved quality of life and physical function, reducing pain severity, and few adverse effects, although the quality of evidence is variable. 61,62 Compliance with exercise interventions is key to their success; interventions with a measured high compliance yielded significantly larger reduction in pain than those with uncertain or unmonitored compliance. 63 The specific effects of physical activity on chronic pain are hard to determine because of the heterogeneity of chronic pain and exercise regimes, and study limitations, including limited follow-up periods and small sample sizes. 61,64 However, there is evidence of benefit for some types of physical activity in particular chronic pain conditions: aquatic exercise can improve chronic back pain and improve physical function, 65 supervised aerobic and strength training has been shown to reduce pain in patients with fibromyalgia, 66 and t'ai chi had a beneficial effect in reducing arthritis pain. 67 Yoga has also been shown to have a beneficial impact on those with chronic pain.<sup>68</sup> Recommendations for including physical activity in treatment plans for the management of chronic pain have started to feature in national and international care guidelines. 14,69

# Nutrition

The role of nutrition in the development and prevention of chronic pain is unclear. Nutrition management plans may be of benefit to patients with chronic pain by improving pain management and reducing cardiovascular risk factors that are related to chronic pain. There have been calls for patients with chronic pain to be offered personalised nutrition assessment and counselling targeted at improving diet and supplement use, and emerging evidence that this may improve the quality of life and clinical outcomes in patients with chronic pain.<sup>70</sup> Omega-3 as a diet supplement in preclinical trials did show an improvement in inflammatory pain,  $^{71}$  whilst garlic has been suggested to reduce pain severity in overweight women with knee arthritis. 72 A recent systematic review and metaanalysis of 23 papers found that interventions based on nutrition, particularly those testing an altered overall diet or a single nutrient, had a significant effect on reducing participants' reported pain severity and intensity.<sup>73</sup> However, the studies in the field of nutrition and chronic pain, including those included in the meta-analysis, were of low quality, 73and there is insufficient evidence to make specific dietary recommendations. More rigorous studies examining nutrition with chronic pain as a primary outcome are needed in order to determine the role of nutrition in chronic pain.<sup>73</sup>

#### Sunshine and vitamin D

Colder climates and lack of sunshine correlate with chronic pain; a study showed less pain was experienced on longer, sunnier days. A relationship between high levels of reported pain and low levels of vitamin D has been demonstrated, with the suggestion that low vitamin D levels cause anatomic, endocrine, neurological, and immunological changes, which predispose to onset and perpetuation of chronic pain. //,/8 However, the effect is not replicated across all studies with only 25% of studies concluding that there is a correlation between low levels of vitamin D and chronic pain. 79,80

## Clinical

#### Pain

The most important clinical risk factor for the development of chronic pain is the presence of another site of acute or chronic pain within the body. 12 The greater the severity and the greater number of sites, the more likely chronic pain is to develop. 12,81 The presence of painful stimuli alters brain chemistry in such a way as to predispose individuals to develop chronic pain. 82 This increased susceptibility to pain can develop within days of exposure to continuous painful stimuli and can persist for up to a year after the pain has resolved.83 Having more than one cause of chronic pain and pain of longer duration are both associated with poorer quality of life.84 Effective analgesia for acute pain is protective against the development of chronic pain. 85 One of the most important ways to reduce the incidence of chronic pain is to prevent acute pain from occurring and managing it well when it does occur.

## Multi-morbidity and mortality

Patients with co-morbid physical and mental chronic diseases are more liable to suffer chronic pain than those without. 19,86 Up to 88% of those with chronic pain have additional chronic diagnoses. 11,19,87 Even after adjusting for known socioeconomic and environmental confounders (e.g. age, gender, smoking, deprivation, and education), there is an increased co-occurrence of chronic pain with depression and cardiovascular disease.<sup>88</sup> Approximately a third of patients with chronic pulmonary disease and coronary heart disease report experiencing chronic pain. <sup>19</sup> The presence of co-morbidities also complicates the clinical management of people with chronic pain by limiting the applicability of disease-specific clinical guidelines and reducing the treatment options available for optimal pain control.<sup>89</sup> In people with other medical co-morbidities, chronic pain is an independent risk factor for all-cause mortality. 90,91 Patients diagnosed with severe chronic pain are twice as likely to have died 10 yr later from ischaemic heart disease or respiratory disease than those who report mild chronic pain or were pain free. 92-95 Those who were 'resilient to pain'-people who experienced a high pain intensity, but documented a low pain disability-had an improved 10 yr survival rate compared with people who were not resilient to pain. 96

Improvements in cancer treatments and care have led to improved survival rates, meaning that chronic pain is becoming more prevalent in those who have had cancer. A French study discovered that a third of patients with cancer had chronic pain; within this group, approximately a fifth had pain with neuropathic characteristics.<sup>97</sup> In patients with chronic obstructive pulmonary disease (COPD), chronic pain is common, and those with COPD and chronic pain were found to have more depression, do less physical exercise, and have higher breathlessness scores than those without COPD. 98 For those living with neurological conditions, the prevalence of chronic pain is double that of the general population, and those with spinal cord injury have the highest levels of pain. 95 Chronic pain is also more prevalent in adolescents with physical disabilities; however, other mental health concerns, such as insomnia, low mood, and anxiety associated with chronic pain, were as prevalent in those without any physical disability who had chronic pain. 100

## Mental health

Chronic pain is linked with depression, and the combination of these conditions can cluster in families. 101 Depression, anxiety, and negative beliefs about pain are all related not only to developing chronic pain, but also to having worse outcomes from chronic pain. 102-105 Depression is strongly linked with chronic pain: 20-50% of patients with chronic pain have comorbid depression, 11,19 and patients with severe pain are more likely to be depressed. 106 Depression often goes unrecognised, and therefore, untreated in patients with chronic pain. 107 Even once depression has improved, people with a history of depression remain at higher risk of chronic pain. 108 It is likely that the aetiology is bidirectional with chronic pain both causing and resulting from poor mental health. 109 Neuroimaging studies on patients with depression showed abnormal prefrontal brain activity and dysfunctional emotional regulation when exposed to experimentally induced painful stimuli. 110,111 Anxiety and fear about pain are linked to a higher likelihood of developing chronic pain and a poorer prognosis of recovery from chronic pain. 102 Fear avoidance behaviours and associated lack of movement are independent risk factors for developing chronic pain. 112 A large population study demonstrated that patients who had consulted their general practitioner (GP) for 'nerves, anxiety, tension, or depression' had a higher risk of consulting about chronic pain than those who had not consulted their GP in this way (52.2% vs 38.0%).37 In post-traumatic stress disorder (PTSD), chronic pain is common, 113,114 although the strength of the relationship varies depending on the cause of the PTSD and the type of chronic pain. The resolution of anxiety and depression may need to be achieved, and effective analgesia, in order to enable effective pain control in chronic pain. 116 Because of the bidirectional relationship between chronic pain and mental health conditions, screening for mental health issues in people with chronic pain, and for chronic pain in people with mental health issues, should be considered. 117

## Surgical and medical interventions

Postoperative chronic pain is a significant complication of many surgical procedures. 118 Whilst rates of post-surgical pain vary, it has been suggested that up to 80% of patients experience some degree of significant postoperative pain. 119 Chronic post-surgical pain affects up to 10% of patients, 120 and is particularly common after amputations (50-85%), thoracotomies (5-65%), cardiac surgery (30-55%), and breast surgery (20-50%). 121 In spite of it becoming a more widely recognised pathophysiology, rates of diagnosis of post-surgical pain have remained unchanged. 122 In foot surgery patients, the risk of developing chronic postoperative pain was not correlated with the severity of surgery. 123 Those with pre-existing severe

chronic pain were more likely to develop chronic postoperative pain. 118,123 The presence of moderate-to-severe pain before the operation and acute post-surgical pain are both independent predictors of whether patients develop chronic post-surgical pain.<sup>24</sup> Patients who were anxious about their operation, younger patients, and those who developed postoperative infections were also more likely to suffer from chronic post-surgical pain. <sup>24,123</sup> Chronic post-surgical pain can have significant negative effects on patients' quality of life, particularly in conditions, such as breast cancer, where survivors are living longer, but living with chronic pain as a result of their treatment. 12

## Weight

Obesity, defined as a BMI of greater than 30, is related to multimorbidity and is an independent predictor of chronic pain. Obesity increases chronic pain in several ways, including placing strain on weight-bearing joints, reducing physical activity, and contributing to overall body deconditioning. 125 One study suggests that nearly 40% of people who are obese experience chronic pain, and that the pain they report is more likely to be moderate to severe than chronic pain in those who are not obese. 126 A large-scale population study found that the likelihood of reporting chronic pain increased proportionately with BMI: compared with groups of people with a healthy BMI, in people with a BMI of 30-34, the rates of pain were 68%; the relative rates of chronic pain were 136% in those with a BMI of 35-39, and 254% in those with BMI >40.127 This increased prevalence of chronic pain is seen even after adjusting for the impact of obesity on other medical conditions, which contribute to multi-morbidity and which are independently associated with pain. 128 There are both environmental and genetic elements to the relationship between pain and obesity. 129 There is, however, limited evidence that weight loss improves chronic pain. 130 However, there is evidence that being underweight is a consideration when managing patients with chronic pain: one study showed a higher chronic pain prevalence in men over 50 who had a BMI of less than 18.5; they also had higher rates of severe depression. 131

# Sleep disorders

Sleep disorders have been shown to affect nearly half of people reporting chronic pain, with a quarter of chronic pain patients suffering from clinical insomnia. 132 The association is bidirectional, with chronic pain causing poor sleep, and poor sleep increasing the intensity and duration of chronic pain. 132 Sleep deprivation was found to be a risk factor for chronic pain in a prospective survey of women over a 17 yr period. 133 Another study showed that having chronic pain made people more likely to suffer from sleep problems and depression, and suggested that treating sleep disorders should be considered as part of chronic pain management. 134 Severe chronic pain after concussion was significantly related to insomnia. 135 There is a high prevalence of sleep apnoea in patients who take opioid medications long term, but patients with chronic pain are at higher risk of developing sleep apnoea irrespective of opioid medication. 136

## Genetics

The relationship between chronic pain and genes is complex. Genes act at many levels to shape the experience of chronic pain, influencing emotional, behavioural, and biological processes. 137 Sensitivity to painful stimuli and pain tolerance are partly genetically determined. 138,139 Chronic pain is a heritable phenotype, and the presence of chronic pain clusters in family groups 140,141 through genetic and 'maternal' effects. It also may be as a result of important genetic contributions to underlying diseases, which will include chronic pain. 63,140,141

One of the current challenges in chronic pain epidemiology is to determine which genes contribute to chronic pain and what their individual roles are. Currently, there are known to be at least 150 genes associated with chronic pain in humans, and this number is ever expanding. 137,142 Amongst others, they include genes from immune, inflammatory, and stressrelated pathways, including COMT and OPRM. 143 Specific genetic variants have been identified with rare chronic pain conditions, such as SNC9A with erythromelalgia. 144 A recent systematic review of genetic factors associated with chronic neuropathic pain found that variants in HLA genes, COMT, OPRM1, TNFA, IL6, and GCH1, were identified in more than one study. 145

At a human population level, research has failed to identify any single genetic variant that contributes substantially to the population risk of developing chronic pain; there is no 'chronic pain gene'. It is more likely that a combination of genetic variants increases the risk of developing chronic pain. However, identifying relevant genes may help to understand underlying biological mechanisms and the search for therapeutic targets. Gene identification from genome-wide association studies (GWAS) may offer hope, particularly as genetic data from large numbers of samples, such as the UK Biobank, 146 are accessible. In one GWAS, a genetic variant on Chromosome 5 was found to be associated with chronic widespread pain in both human genome- and animal-wide association study meta-analyses. 147 A more recent GWAS and meta-analysis of 158 000 individuals identified three novel genetic variants associated with chronic back pain. 148

## Other

## Attitudes and beliefs about pain

Personal beliefs and attitudes can affect a person's likelihood of developing long-term pain or pain-related disability. 149,150 Patients who adopted passive coping strategies, such as 'resting and taking medications', were found to use three times the amount of healthcare appointments and have double the level of disability from pain in comparison with those who adopted active strategies (e.g. exercise). 151 Changing patients' attitudes to their pain can reduce their level of pain and the impact of their quality of life. 152

## History of violent injury, abuse, or interpersonal violence

The severity and development of chronic pain experience are affected by early life factors: people who experience adversity or emotional trauma (e.g. death of parent and being raised in the care system) or physical trauma (e.g. substantial hospitalisation and preterm birth) in childhood have a higher risk of chronic pain in their adult lives. 153 Early stress in life can alter the function of the hypothalamic pituitary adrenal axis, affecting the stress response.<sup>153</sup> Young people who have experienced traumatic adverse childhood experiences (ACEs) have a greater chance of developing chronic pain than those who have not. A study of children and 9-19 years with chronic pain found that the most common ACE in children with chronic pain was having family members with mental health illnesses; 55% of children with multiple ACEs experience chronic pain. 154 The more ACEs, the greater the level of chronic widespread pain and psychological distress, such as anxiety and depression (which have been noted previously to be related to the development and severity of chronic pain). 154

People who have experienced personal violence or abusive relationships are more likely to experience subsequent chronic pain. 43,155 This has been found to be true regardless of the age at which the violence or abuse was experienced, or whether it was domestic or public violence or abuse. 155,156

## Limitations

It is challenging to draw conclusions on chronic pain epidemiology given the heterogeneity of the research in this field. whose sources include variations in how chronic pain is defined in each study and of the populations studied. The latter may impact on the generalisability of study findings in other populations, regions, or countries. Differences in study design also limit the applicability of individual research findings. These differences make robust data synthesis, including meta-analysis, particularly challenging in this field. In particular, it is generally impossible to determine causality from cross-sectional studies, and therefore, to distinguish risk factors from outcomes of chronic pain.

#### Conclusion

The significant global disease burden of chronic pain needs to be addressed by managing the causes and effects of chronic pain, targeted at both individual and population levels. To prevent and reduce the impact of chronic pain, modifiable risk factors (e.g. acute pain, lifestyle, and behaviour) need to be addressed, with the patient at the centre of the management. In the longer term, attention to the local and global distributions of wealth and access to education are likely to be more important. Employing modern techniques in further epidemiological research, such as neuroimaging and genetics, is important in identifying underlying biological mechanisms and potential therapeutic targets.

## Authors' contributions

Literature search: SEEM, KPN.

Identifying factors used in the thematic synthesis: all authors.

Drafting paper: SEEM, KPN.

Critically revised the article for important intellectual content and had final approval of the version to be published: BHS.

## **Declaration of interest**

The authors declare that they have no conflicts of interest.

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## References

1. Fayaz A, Croft P, Langford R, Donaldson J, Jones G. Prevalence of chronic pain in the UK: a systematic

- review and meta-analysis of population studies. BMJ Open 2016; 6
- 2. International Association for the Study of Pain, Subcommittee on Taxonomy. Classification of chronic pain. Descriptions of chronic pain syndromes and definitions of pain terms. Pain Suppl 1986; 3: S1-226
- 3. Tracey I, Bushnell M. How neuroimaging studies have challenged us to rethink: is chronic pain a disease? J Pain 2009; 10: 1113-20
- 4. World Health Organization. ICD-11 international classification of diseases for mortality and morbidity statistics, 2018, https://icd.who.int/browse11/l-m/en, accessed 7 September 2018. 14: SIGN guidance is at https://www.sign. ac.uk/assets/sign136.pdf, accessed 1 September 2018.
- 5. Vos T, Allen C, Arora M, et al. Global, regional, and national incidence, prevalence, and years lived with disability for 328 diseases and injuries for 195 countries, 1990-2016: a systematic analysis for the Global Burden of Disease Study 2016. Lancet 2017; 390: 1211-59
- 6. Last R. A Dictionary of Epidemiology. 4th ed. Oxford: International Epidemiological Association; 2001
- 7. Merskey H, Bogduk N, editors. IASP task force on taxonomy, Part III: Pain Terms, A Current List with Definitions and Notes on Usage. Seattle, WA: IASP Press; 1994. p. 209-14
- 8. van Hecke O, Torrance N, Smith B. Chronic pain epidemiology and its clinical relevance. Br J Anaesth 2013; 111:
- 9. Diatchenko L, Fillingim RB, Smith SB, Maixner W. The phenotypic and genetic signatures of common musculoskeletal pain conditions. *Nat Rev Rheumatol* 2013; **9**: 340–50
- 10. van Hecke O, Torrance N, Smith B. Chronic pain epidemiology—where do lifestyle factors fit in? Br J Pain 2013; 7: 209-17
- 11. Breivik H, Collett B, Ventafridda V, Cohen R, Gallacher D. Survey of chronic pain in Europe: prevalence, impact on daily life, and treatment. Eur J Pain 2006; 10: 287-333
- 12. Elliott A, Smith B, Hannaford P. The course of chronic pain in the community: results of a 4-year follow-up study. Pain 2002; 99: 299-307
- 13. Dorner TE. Pain and chronic pain epidemiology: implications for clinical and public health fields. Wien Klin Wochenschr 2018; 130: 1-3
- 14. Scottish Intercollegiate Guideline Network. SIGN 136: management of chronic pain 2012
- 15. King S, Chambers CT, Huguet A, et al. The epidemiology of chronic pain in children and adolescents revisited: a systematic review. Pain 2011; 152: 2729-38
- 16. Thomas E. In: Croft P, Blyth FM, van der Windt D, editors. Chapter 4: Chronic pain epidemiology. Oxford: Oxford University Press; 2010. p. 185
- 17. Docking R, Fleming J, Brayne C, et al. Epidemiology of back pain in older adults: prevalence and risk factors for back pain onset. Rheumatology 2011; 50: 1645-53
- 18. Thomas E, Peat G, Harris L, Wilkie R, Croft PR. The prevalence of pain and pain interference in a general population of older adults: cross-sectional findings from the North Staffordshire Osteoarthritis Project (NorStOP). Pain 2004; 110: 361-8
- 19. Barnett K, Mercer SW, Norbury M, Watt G, Wyke S, Guthrie B. Epidemiology of multimorbidity and implications for health care, research, and medical education: a cross-sectional study. Lancet 2012; 380: 37-43
- 20. Johnson R, McElhaney J. Postherpetic neuralgia in the elderly. Int J Clin Pract 2009; 63: 1386-91

- 21. Schofield P. The assessment of pain in older people: UK National Guidelines. Age Ageing 2018; 47: i1-22
- 22. Larsson C, Hansson EE, Sundquist K, Jakobsson U. Chronic pain in older adults: prevalence, incidence, and risk factors. Scand J Rheumatol 2017; 46: 317-25
- 23. Gobina I, Villberg J, Välimaa R, et al. Prevalence of selfreported chronic pain among adolescents: evidence from 42 countries and regions. Eur J Pain 2019 Feb; 23: 316-26
- 24. Bruce J, Quinlan J. Chronic post surgical pain. Rev Pain 2011; **5**: 23–9
- 25. Greenspan J, Craft R, LeResche L. Studying sex and gender differences in pain and analgesia: a consensus report. Pain 2007; 132: S26-45
- 26. El-Shormilisy N, Strong J, Meredith PJ. Associations among gender, coping patterns and functioning for individuals with chronic pain: a systematic review. Pain Res Manag 2015; 20
- 27. Wiesenfeld H. Sex differences in pain perception. Gend Med 2005; 2
- 28. Unruh A. Gender variations in clinical pain experience. Pain 1996; 65: 123-67
- 29. Craft R, Mogil J, Aloisi A. Sex differences in pain and analgesia: the role of gonadal hormones. Eur J Pain 2004;
- 30. Malon J, Shah P, Koh WY, Cattabriga G, Li E, Cao L. Characterizing the demographics of chronic pain patients in the state of Maine using the Maine all payer claims database. BMC Public Health 2018; 18: 810
- 31. Ferreira Kdos S, Speciali J. Epidemiology of chronic pain in the office of a pain specialist neurologist. Arg Neuropsiquiatr 2015; 73: 582-5
- 32. Bartley EJ, Fillingim RB. Sex differences in pain: a brief review of clinical and experimental findings. Br J Anaesth 2013: 111: 52-8
- 33. Craft R. Modulation of pain by estrogens. Pain 2007; 132:
- 34. Meng W, Deshmukh HA, Donnelly LA, et al. A genomewide association study provides evidence of sex-specific involvement of Chr1p35.1 (ZSCAN20-TLR12P) and Chr8p23.1 (HMGB1P46) with diabetic neuropathic pain. EBioMedicine 2015; 2: 1386-93
- 35. Campbell C, Edwards RR. Ethnic differences in pain and pain management. Pain Manag 2012; 2: 219-30
- 36. Janevic MR, McLaughlin SJ, Heapy AA, Thacker C, Piette JD. Racial and socioeconomic disparities in disabling chronic pain: findings from the Health and Retirement Study. J Pain 2017; 18: 1459-67
- 37. Macfarlane G, Beasley M, Smith B, Jones G, Macfarlane T. Can large surveys conducted on highly selected populations provide valid information on the epidemiology of common health conditions? An analysis of UK Biobank data on musculoskeletal pain. Br J Pain 2015; 9: 203 - 12
- 38. Blyth F. Chronic pain—is it a public health problem? Pain 2008; **137**: 465–6
- 39. Poleshuck E, Green C. Socioeconomic disadvantage and pain. Pain 2008; 136: 235-8
- 40. Maly A, Vallerand AH. Neighborhood, socioeconomic, and racial influence on chronic pain. Pain Manag Nurs 2018; 19: 14-22
- 41. Brekke M, Hjortdahl P, Kvien T. Severity of musculoskeletal pain: relationship to socioeconomic inequality. Soc Sci Med 2002; 54: 221-8

- **42.** Eachus J, Chan P, Pearson N, Propper C, Davey-Smith G. An additional dimension to health inequality: disease severity and socioeconomic position. *J Epidemiol Community Health* 1999; **53**: 603–11
- 43. Jordan K, Thomas E, Peat G, Wilkie R, Croft P. Social risks for disabling pain in older people: a prospective study of individual and area characteristics. Pain 2008; 137: 652–66
- 44. Shaw W, Linton S, Pransky G. Reducing sickness absence from work due to low back pain: how well do intervention strategies match modifiable risk factors? *J Occup Rehabil* 2006; **16**: 591–605
- **45.** Teasell R, Bombardier C. Employment-related factors in chronic pain and chronic pain disability. *Clin J Pain* 2001; **17**: S39–45
- 46. Kerkhof EVD, Mann EG, Torrance N, Smith BH, Johnson A, Gilron I. An epidemiological study of neuropathic pain symptoms in Canadian adults. Pain Res Manag 2016
- Saastamoinen P, Leino-Arjas P, Laaksonen M, Lahelma E. Socio-economic differences in the prevalence of acute, chronic and disabling chronic pain among ageing employees. Pain 2005; 114: 364–71
- **48.** Ruokolainen O, Auvinen J, Linton SJ, et al. OMPSQ-Short Score and determinants of chronic pain: cross-sectional results from a middle-aged birth cohort. *Eur J Phys Rehabil Med* 2018; **54**: 34–40
- 49. Vogt M, Hanscom B, Lauerman WC, Kang JD. Influence of smoking on the health status of spinal patients: the National Spine Network database. Spine (Phila Pa 1976) 2002; 27: 313–9
- 50. Ekholm O, Gronbaek M, Peuckmann V, Sjogren P. Alcohol and smoking behavior in chronic pain patients: the role of opioids. Eur J Pain 2009; 13: 606–12
- Orhurhu VJ, Pittelkow TP, Hooten WM. Prevalence of smoking in adults with chronic pain. Tob Induc Dis 2015; 13: 17
- 52. Ditre JW, Zale EL, Heckman BW, Hendricks PS. A measure of perceived pain and tobacco smoking interrelations: pilot validation of the pain and smoking inventory. *Cogn Behav Ther* 2017; 46: 339–51
- 53. John U, Hanke M, Meyer C, Völzke H, Baumeister SE, Alte D. Tobacco smoking in relation to pain in a national general population survey. *Prev Med* 2006; 43: 477–81
- 54. Weingarten TN, Moeschler SM, Ptaszynski AE, Hooten WM, Beebe TJ, Warner DO. An assessment of the association between smoking status, pain intensity, and functional interference in patients with chronic pain. Pain Physician 2008; 11: 643–53
- 55. Ditre JW, Brandon TH, Zale EL, Meagher MM. Pain, nicotine, and smoking: research findings and mechanistic considerations. Psychol Bull 2011; 137: 1065–93
- 56. van Hecke O, Torrance N, Cochrane L, et al. Does a history of depression actually mediate smoking-related pain? Findings from a cross-sectional general population-based study. Eur J Pain 2014; 18: 1223–30
- 57. Alford DP, German JS, Samet JH, Cheng DM, Lloyd-Travaglini CA, Saitz R. Primary care patients with drug use report chronic pain and self-medicate with alcohol and other drugs. J Gen Intern Med 2016; 31: 486–91
- 58. Riley J, King C. Self-report of alcohol use for pain in a multi-ethnic community sample. *J Pain* 2009; 10: 944–52
- 59. Brennan PL, Schutte KK, Moos RH. Pain and use of alcohol to manage pain: prevalence and 3-year outcomes

- among older problem and non-problem drinkers. Addiction 2005; **100**: 777–86
- Egli M, Koob GF, Edwards S. Alcohol dependence as a chronic pain disorder. Neurosci Biobehav Rev 2012; 36: 2179—92
- 61. Geneen LJ, Moore RA, Clarke C, Martin D, Colvin LA, Smith BH. Physical activity and exercise for chronic pain in adults: an overview of Cochrane Reviews. Cochrane Database Syst Rev 2017; 1, CD011279
- 62. Marley J, Tully MA, Porter-Armstrong A, et al. The effectiveness of interventions aimed at increasing physical activity in adults with persistent musculoskeletal pain: a systematic review and meta-analysis. BMC Musculoskelet Disord 2017; 18: 482
- **63.** Moseng T, Dagfinrud H, Smedslund G, Osteras N. The importance of dose in land-based supervised exercise for people with hip osteoarthritis. A systematic review and meta-analysis. Osteoarthritis Cartilage 2017; **25**: 1563–76
- **64.** Parreira P, Heymans MW, van Tulder MW, et al. Back schools for chronic non-specific low back pain. Cochrane Database Syst Rev 2017; **8**
- 65. Shi Z, Zhou H, Lu L, et al. Aquatic exercises in the treatment of low back pain: a systematic review of the literature and meta-Analysis of eight studies. Am J Phys Med Rehabil 2017
- 66. Busch AJ, Schachter CL, Overend TJ, Peloso PM, Barber KA. Exercise for fibromyalgia: a systematic review. J Rheumatol 2008; 35: 1130–44
- 67. Hall A, Maher C, Latimer J, Ferreira M. The effectiveness of Tai Chi for chronic musculoskeletal pain conditions: a systematic review and meta-analysis. *Arthritis Rheum* 2009; 61: 717–24
- 68. Schmid AA, Atler KE, Malcolm MP, et al. Yoga improves quality of life and fall risk-factors in a sample of people with chronic pain and type 2 diabetes. Complement Ther Clin Pract 2018; 31: 369–73
- 69. Rausch Osthoff AK, Niedermann K, Braun J, et al. 2018 EULAR recommendations for physical activity in people with inflammatory arthritis and osteoarthritis. Ann Rheum Dis 2018; 77: 1251–60
- 70. De Gregori M, Muscoli C, Schatman ME, et al. Combining pain therapy with lifestyle: the role of personalized nutrition and nutritional supplements according to the SIMPAR Feed Your Destiny approach. *J Pain Res* 2016; 9: 1179–89
- Goldberg R, Katz J. A meta-analysis of the analgesic effects of omega-3 polyunsaturated fatty acid supplementation of inflammatory joint pain. Pain 2007; 152: 210–23
- 72. Dehghani S, Alipoor E, Salimzadeh A, et al. The effect of a garlic supplement on the pro-inflammatory adipocytokines, resistin and tumor necrosis factor-alpha, and on pain severity, in overweight or obese women with knee osteoarthritis. *Phytomedicine* 2018; 48: 70–5
- 73. Brain K, Burrows TL, Rollo ME, et al. A systematic review and meta-analysis of nutrition interventions for chronic noncancer pain. *J Hum Nutr Diet* 2018
- 74. Scottish Intercollegiate Guideline Network. SIGN 136: management of chronic pain 2013
- 75. Sesti F, Capozzolo T, Pietropolli A, Collalti M, Bollea MR, Piccione E. Dietary therapy: a new strategy for management of chronic pelvic pain. Nutr Res Rev 2011; 24: 31–8
- 76. Hagen K, Byfuglien M, Falzon L, Olsen SU, Smedslund G. Dietary interventions for rheumatoid arthritis. Cochrane

- Database Syst Rev 2009; 1. https://www.cochranelibrary. com/cdsr/doi/10.1002/14651858.CD006400.pub2/media/ CDSR/CD006400/CD006400.pdf. accessed 1 September 2018
- 77. Shipton EE, Shipton EA. Vitamin D deficiency and pain: clinical evidence of low levels of vitamin D and supplementation in chronic pain states. Pain Ther 2015; 4: 67–87
- 78. Shipton EA, Shipton EE. Vitamin D and pain: vitamin D and its role in the aetiology and maintenance of chronic pain states and associated comorbidities. Pain Res Treat 2015; **2015**: 1-12
- 79. Straube S, Andrew Moore R, Derry S, McQuay HJ. Vitamin D and chronic pain. Pain 2009; 141: 10-3
- 80. Straube S, Derry S, Straube C, Moore RA. Vitamin D for the treatment of chronic painful conditions in adults. Cochrane Database Syst Rev 2010; 1
- 81. Bergman S, Herrstrom P, Jacobsson L, Petersson IF. Chronic widespread pain: a three year followup of pain distribution and risk factors. J Rheumatol 2002; 29: 818-25
- 82. Rodriguez-Raecke R, Niemeier A, Ihle K, Ruether W, May A. Brain gray matter decrease in chronic pain is the conse- quence and not the cause of pain. J Neurosci 2009; **29**: 13746-50
- 83. Teutsch S, Herken W, Bingel U. Changes in brain gray matter due to repetitive painful stimulation. Neuroimage 2008; **42**: 845-9
- 84. Pagé G, Fortier M, Ware M, Choinière M. As if one pain problem was not enough: prevalence and patterns of coexisting chronic pain conditions and their impact on treatment outcomes. J Pain Res 2018: 237-54
- 85. Gwilym S, Filippini N, Douaud G, Carr AJ, Tracey I. Thalamic atrophy associated with painful osteoarthritis of the hip is reversible after arthroplasty; a longitudinal voxel-based morphometric study. Arthritis Rheum 2010;
- 86. Dominick C, Blyth F, Nicholas M. Unpacking the burden: understanding the relationships between chronic pain and co-morbidity in the general population. Pain 2012; **153**: 292-304
- 87. Donaldson L. 150 years of the Annual Report of the Chief Medical Officer: On the state of public health 2008. Department of Health, Richmond House, 79 Whitehall, London SW1A 2NJ, UK 2009. dhmail@dh.gsi.gov.uk
- 88. van Hecke O, Hocking L, Torrance N, et al. Chronic pain, depression and cardiovascular disease linked through a shared genetic predisposition: analysis of a family-based cohort and twin study. PloS One 2017; 12: 1-19
- 89. Guthrie B, Payne K, Alderson P, McMurdo ME, Mercer SW. Adapting clinical guidelines to take account of multimorbidity. BMJ 2012; 345: 5
- 90. Smith D, Wilkie R, Uthman O, Jordan JL, McBeth J. Chronic pain and mortality: a systematic review. PloS One 2014; 9, e99048
- 91. Macfarlane G, Barnish M, Jones G. Persons with chronic widespread pain experience excess mortality: longitudinal results from UK Biobank and meta-analysis. Ann Rheum Dis 2017; 76
- 92. Torrance N, Elliott A, Lee A, Smith B. Severe chronic pain is associated with increased 10 year mortality. A cohort record linkage study. Eur J Pain 2010; 14: 380-6
- 93. Kareholt I, Brattberg G. Pain and mortality risk among elderly persons in Sweden. Pain 1998; 77: 271-8
- 94. Smith B, Elliott A, Hannaford P. Pain and subsequent mortality and cancer among women in the royal college

- of general practitioners oral contraception study. Br J Gen Pract 2003; **53**: 45-6
- 95. Andersson H. The course of non-malignant chronic pain: a 12-year follow-up of a cohort from the general population. Eur J Pain 2004; 8: 47-53
- 96. Elliott A, Burton C, Hannaford P. Resilience does matter: evidence from a 10-year cohort record linkage study. BMJ Open 2013
- 97. Bouhassira D, Luporsi E, Krakowski I. Prevalence and incidence of chronic pain with or without neuropathic characteristics in patients with cancer. Pain 2017; 158: 1118-25
- 98. Lee AL, Goldstein RS, Brooks D. Chronic pain in people with chronic obstructive pulmonary disease: prevalence, clinical and psychological implications. Chronic Obstr Pulm Dis 2017; 4: 194-203
- 99. Cragg JJ, Warner FM, Shupler MS, et al. Prevalence of chronic pain among individuals with neurological conditions. Health Rep 2018; 29: 11-6
- 100. de la Vega R, Groenewald C, Bromberg MH, Beals-Erickson SE, Palermo TM. Chronic pain prevalence and associated factors in adolescents with and without physical disabilities. Dev Med Child Neurol 2018; 60: 596-601
- 101. McIntosh AM, Hall LS, Zeng Y, et al. Genetic and environmental risk for chronic pain and the contribution of risk variants for major depressive disorder: a familybased mixed-model analysis. PLoS Med 2016; 13, e1002090
- 102. Boersma K, Linton S. Expectancy, fear and pain in the prediction of chronic pain and disability: a prospective analysis. Eur J Pain 2006; 10: 551-7
- 103. van der Windt D, Croft P, Penninx B. Neck and upper limb pain: more pain is associated with psychological distress and consultation rate in primary care. J Rheumatol 2002;
- 104. van der Windt D, Kuijpers T, Jellema P, van der Heijden G, Bouter L. Do psychological factors predict outcome in both low-back pain and shoulder pain? Ann Rheum Dis 2007; 66: 313-9
- 105. Nijrolder I, van der Windt D, van der Horst H. Prediction of outcome in patients presenting with fatigue in primary care. Br J Gen Pract 2009; **59**: e101-9
- 106. de Heer E, Ten Have M, van Marwijk HWJ, et al. Pain as a risk factor for common mental disorders. Results from The Netherlands Mental Health Survey and Incidence Study-2: a longitudinal population-based study. Pain 2018; **159**: 712-8
- 107. Lee HJ, Choi EJ, Nahm FS, Yoon IY, Lee PB. Prevalence of unrecognized depression in patients with chronic pain without a history of psychiatric diseases. Korean J Pain 2018; **31**: 116–24
- 108. Gerrits MM, van Marwijk HW, van Oppen P, van der Horst H, Penninx BW. Longitudinal association between pain, and depression and anxiety over four years. J Psychosom Res 2015; 78: 64-70
- 109. Van Korff M, Le Resche L, Dworkin SF. First onset of common pain symptoms: a prospective study of depression as a risk factor. Pain 1993; 55: 251-8
- 110. Bär KJ, Wagner G, Koschke M, et al. Increased prefrontal activation during pain perception in major depression. Biol Psychiatry 2007; **62**: 1281-7
- 111. Strigo I, Simmons A, Matthews S, Craig A, Paulus M. Association of major depressive disorder with altered functional brain response during anticipation and

- processing of heat pain. Arch Gen Psychiatry 2008; 65: 1275-84
- 112. Wakaizumi K, Yamada K, Oka H, et al. Fear-avoidance beliefs are independently associated with the prevalence of chronic pain in Japanese workers. J Anesth 2017; 31: 255-62
- 113. Akhtar E, Ballew AT, Orr WN, Mayorga A, Khan TW. The prevalence of post-traumatic stress disorder symptoms in chronic pain patients in a tertiary care setting: a crosssectional study. Psychosomatics 2018
- 114. Siqveland J, Hussain A, Lindstrom JC, Ruud T, Hauff E. Prevalence of posttraumatic stress disorder in persons with chronic pain: a meta-analysis. Front Psychiatry 2017; 8: 164
- 115. Fishbain DA, Pulikal A, Lewis JE, Gao J. Chronic pain types differ in their reported prevalence of post-traumatic stress disorder (PTSD) and there is consistent evidence that chronic pain is associated with PTSD: an evidencebased structured systematic review. Pain Med 2017; 18: 711-35
- 116. Kroenke K, Bair MJ, Damush T, et al. Optimized antidepressant therapy and pain self-management in primary care patients with depression and musculo- skeletal pain: a randomized controlled trial. JAMA 2009; 301: 2099-110
- 117. Pereira FG, Franca MH, Paiva MCA, Andrade LH, Viana MC. Prevalence and clinical profile of chronic pain and its association with mental disorders. Rev Saude Publica 2017; 51: 96
- 118. Gan TJ. Poorly controlled postoperative pain: prevalence, consequences, and prevention. J Pain Res 2017; 10: 2287-98
- 119. Institute of Medicine (US). Committee on advancing pain research, care, and education. Relieving pain in America: a blueprint for transforming prevention, care, education, and research. Washington, DC: The National Academies Collection: reports funded by National Institutes of Health; 2011
- 120. Fletcher D, Stamer UM, Pogatzki-Zahn E, et al. Chronic postsurgical pain in Europe: an observational study. Eur J Anaesthesiol 2015; 32: 725-34
- 121. Macrae W. Chronic post-surgical pain: 10 years on. Br J Anaesth 2008; 101: 77-86
- 122. Correll D. Chronic postoperative pain: recent findings in understanding and management. F1000Res 2017; 6: 1054
- 123. Remérand F, Godfroid HB, Brilhault J, et al. Chronic pain 1 year after foot surgery: epidemiology and associated factors. Orthop Traumatol Surg Res 2014; 100: 767-73
- 124. Hamood R, Hamood H, Merhasin I, Keinan-Boker L. Chronic pain and other symptoms among breast cancer survivors: prevalence, predictors, and effects on quality of life. Breast Cancer Res Treat 2018; 167: 157-69
- 125. Hitt HC, McMillen RC, Thornton-Neaves T, Koch K, Cosby AG. Comorbidity of obesity and pain in a general population: results from the Southern Pain Prevalence Study. Pain 2007; 430-6
- 126. Coaccioli S, Masia F, Celi G, Grandone I, Crapa ME, Fatati G. Chronic pain in the obese: a quali-quantitative observational study. Recenti Prog Med 2014; 105: 151-4
- 127. Stone AA, Broderick JE. Obesity and pain are associated in the United States. Obesity 2012; **20**: 11491-5
- 128. Ray L, Lipton R, Zimmerman M, Katz MJ, Derby CA. Mechanisms of association between obesity and chronic pain in the elderly. Pain 2011; **152**: 53-9

- 129. Wright L, Schur E, Noonan C, Ahumada S, Buchwald D, Afari N. Chronic pain, overweight, and obesity: findings from a community-based twin registry. Pain 2010; 11: 628-35
- 130. Okifuji A, Hare B. The association between chronic pain and obesity. J Pain Res 2015; 8: 399-408
- 131. Yamada K, Kubota Y, Iso H, Oka H, Katsuhira J, Matsudaira K. Association of body mass index with chronic pain prevalence: a large population-based crosssectional study in Japan. J Anesth 2018; 32: 360-7
- 132. Jank R, Gallee A, Boeckle M, Fiegl S, Pieh C. Chronic pain and sleep disorders in primary care. Pain Res Treat 2017; **2017**: 9081802
- 133. Nittera AK, Pripp AH, Forsetha KØ. Are sleep problems and non-specific health complaints risk factors for chronic pain? A prospective population-based study with 17 year follow-up. Scand J Pain 2012; 3: 210-7
- 134. Campbell P, Tang N, McBeth J, et al. The role of sleep problems in the development of depression among those with chronic pain: a prospective cohort study. Sleep 2013; **36**: 1693-8
- 135. Theunissen M, Peters ML, Schepers J, et al. Recovery 3 and 12 months after hysterectomy epidemiology and predictors of chronic pain, physical functioning, and global surgical recovery. Medicine (Baltimore) 2016: 95
- 136. Tentindo GS, Fishman SM, Li CS, Wang Q, Brass SD. The prevalence and awareness of sleep apnea in patients suffering chronic pain: an assessment using the STOP-Bang sleep apnea questionnaire. Nat Sci Sleep 2018; 10: 217 - 24
- 137. Zorina-Lichtenwalter K, Meloto C, Khoury S, Diatchenko L. Genetic predictors of human chronic pain conditions. Neuroscience 2016; 338: 36-62
- 138. Norbury TA, MacGregor AJ, Urwin J, Spector TD, McMahon SB. Heritability of responses to painful stimuli in women: a classical twin study. Brain 2007; 130: 3041-9
- 139. Nielsen CS, Stubhaug A, Price DD, Vassend O, Czajkowski N, Harris JR. Individual differences in pain sensitivity: genetic and environmental contributions. Pain 2008; 136: 21-9
- 140. Hocking LJ, Generation Scotland, Morris AD, Dominiczak AF, Porteous DJ, Smith BH. Heritability of chronic pain in 2195 extended families. Eur J Pain 2012;
- 141. Grøholt EK, Stigum H, Nordhagen R, Köhler L. Recurrent pain in children, socio-economic factors and accumulation in families. Eur J Epidemiol 2003; 18: 965-75
- 142. Diatchenko L, Nackley A, Tchivileva I, Shabalina S, Maixner W. Genetic architecture of human pain perception. Trends Genet 2006; 23: 605-13
- 143. James S. Human pain and genetics: some basics. Br J Pain 2013; **7**: 171-8
- 144. Hisama F, Dib-Hajj S, Waxman S. SCN9A-Related inherited erythromelalgia 2006
- 145. Veluchamy A, Hébert HL, Meng W, Palmer CNA, Smith BH. Systematic review and meta-analysis of genetic risk factors for neuropathic pain. Pain 2018; 159: 825 - 48
- 146. UK Biobank. Available from: http://www.ukbiobank.ac.uk; 2017
- 147. Peters MJ, Broer L, Willemen HL, et al. Genome-wide association study meta-analysis of chronic widespread pain: evidence for involvement of the 5p15.2 region. Ann Rheum Dis 2013; 72: 427-36

- 148. Suri P, Palmer MR, Tsepilov YA, et al. Genome-wide meta-analysis of 158,000 individuals of European ancestry identifies three loci associated with chronic back pain. PLoS Genet 2018; 14, e1007601
- 149. Rainville J, Smeets RJ, Bendix T, Tveito TH, Poiraudeau S, Indahl AJ. Fear-avoidance beliefs and pain avoidance in low back pain-translating research into clinical practice. Spine J 2011; 9: 895-903
- 150. Darlow B, Fullen BM, Dean S, Hurley DA, Baxter GD, Dowell A. The association between health care professional attitudes and beliefs and the attitudes and beliefs, clinical management, and outcomes of patients with low back pain: a systematic review. Eur J Pain 2012; 16: 3-17
- 151. Blyth F, March L, Nicholas M, Cousins M. Self-management of chronic pain: a population-based study. Pain 2005; 113: 285-92
- 152. Butchart A, Kerr E, Heisler M, Piette J, Krein S. Experience and management of chronic pain among patients with

- other complex chronic conditions. Clin J Pain 2009; 25: 293-8
- 153. Macfarlane G. The epidemiology of chronic pain. Pain 2016; **157**: 2158-9
- 154. Nelson S, Simons LE, Logan D. The incidence of adverse childhood experiences (ACEs) and their association with pain-related and psychosocial impairment in youth with chronic pain. Clin J Pain 2018; 34: 402-8
- 155. Sachs-Ericsson N, Kendall-Tackett K, Hernandez A. Childhood abuse, chronic pain, and depression in the National Comorbidity Survey. Child Abuse Negl 2007; 31: 531-47
- 156. Ellsberg M, Jansen HA, Heise L, et al. Intimate partner violence and women's physical and mental health in the WHO multi-country study on women's health and domestic violence: an observational study. Lancet 2008; 371: 1165-72

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