

Information about how the guideline was developed is on the [guideline's page](#) on the NICE website. This includes the evidence reviews, the scope, and details of the committee and any declarations of interest.

The recommendations in this guideline were developed before the coronavirus pandemic. Please tell us if there are any particular issues relating to COVID-19 that we should take into account when finalising the guideline for publication.

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1 Recommendations

People have the right to be involved in discussions and make informed decisions about their care, as described in [making decisions about your care](#).

[Making decisions using NICE guidelines](#) explains how we use words to show the strength (or certainty) of our recommendations, and has information about prescribing medicines (including off-label use), professional guidelines, standards and laws (including on consent and mental capacity), and safeguarding.

2 **1.1 Assessing all types of chronic pain**

3 1.1.1 When assessing and managing any type of [chronic pain](#), follow the
4 recommendations in the [NICE guideline on patient experience in adult
5 NHS services](#), particularly on:

- 6 • knowing the patient as an individual
- 7 • enabling patients to actively participate in their care, including:
 - 8 – communication
 - 9 – information
 - 10 – shared decision making.

11 Recognise that chronic pain can cause distress. Foster a collaborative
12 supportive relationship.

13 1.1.2 Ask the person to describe how pain affects their life, and how their life
14 may affect their pain. This might include effects on:

- 15 • lifestyle and day-to-day activities, including work and sleep disturbance
- 16 • physical and psychological wellbeing
- 17 • social interaction and relationships.

18 1.1.3 Ask the person about their understanding and acceptance of their
19 condition, and that of their family, carers and significant others. This might
20 include:

- 1 • what causes the pain
2 • what might happen in the future, including expectations about the pain,
3 outcome of treatments and quality of life.
- 4 1.1.4 During discussions with the person and their family or carers (as
5 appropriate), acknowledge the fact that the pain may not improve or may
6 get worse.
- 7 1.1.5 Develop a care plan with the person with chronic pain. Explore their
8 priorities, strengths, preferences, interests and abilities to inform the plan.
- 9 1.1.6 Discuss the possible benefits, risks and uncertainties of all management
10 options for the person's condition when first developing the care plan and
11 at all stages of care.
- 12 1.1.7 Provide advice and information relevant to the person's individual
13 preferences, at all stages of care, to help them make decisions about
14 managing their condition.
- 15 1.1.8 When communicating negative or normal test results, be sensitive to the
16 risk of invalidating the person's experience of pain.

For a short explanation of why the committee made the recommendations on assessing chronic pain and how they might affect practice, see [rationale and impact](#).

Full details of the evidence and the committee's discussion are in evidence review A: Factors that may be barriers to successfully managing chronic pain, and evidence review B: Communication between healthcare professionals and people with chronic pain.

17 **1.2 Managing all types of chronic pain**

18 For guidance on specific conditions that cause pain, see the [NICE guidelines on](#)
19 [headaches](#), [low back pain and sciatica](#), [rheumatoid arthritis](#), [osteoarthritis](#),
20 [spondyloarthritis](#), [neuropathic pain](#), [endometriosis](#) and [irritable bowel syndrome](#).

1 **Pain management programmes**

- 2 1.2.1 Be aware that there was inconsistent evidence on the effectiveness of
3 pain management programmes, so the committee made a
4 [recommendation for research](#).

For a short explanation of why the committee made a recommendation for research on pain management programmes, see [rationale](#)

Full details of the evidence and the committee's discussion are in evidence review C: Pain management programmes.

5 **Social interventions**

- 6 1.2.2 Be aware that no evidence on social interventions for chronic pain was
7 identified, so the committee made a [recommendation for research](#).

For a short explanation of why the committee made a recommendation for research on social interventions, see [rationale](#)

Full details of the evidence and the committee's discussion are in evidence review D: Social interventions.

8 **1.3 *Managing chronic primary pain***

9 **Non-pharmacological management of chronic primary pain**

10 ***Exercise for chronic primary pain***

- 11 1.3.1 Offer a supervised group exercise programme (for example,
12 cardiovascular, mind–body, strength, or a combination of approaches) to
13 people aged 16 years and over to manage [chronic primary pain](#). Take
14 people's specific needs, preferences and abilities into account.

- 15 1.3.2 Encourage people with chronic primary pain to carry on with their exercise
16 for longer-term general health benefits (also see [NICE guidelines on](#)
17 [physical activity](#) and [behaviour change: individual approaches](#)).

For a short explanation of why the committee made the recommendations on exercise and how they might affect practice, see [rationale and impact](#)

Full details of the evidence and the committee's discussion are in evidence review E: Exercise.

1 ***Psychological therapy for chronic primary pain***

2 1.3.3 Consider acceptance and commitment therapy (ACT) or cognitive–
3 behavioural therapy (CBT) for pain for people aged 16 years and over
4 with chronic primary pain.

5 1.3.4 Do not offer biofeedback to people aged 16 years and over to manage
6 chronic primary pain.

For a short explanation of why the committee made the recommendations on psychological therapy and how they might affect practice, see [rationale and impact](#)

Full details of the evidence and the committee's discussion are in evidence review F: Psychological therapy.

7 ***Acupuncture for chronic primary pain***

8 1.3.5 Consider a course of acupuncture or dry needling, within a traditional
9 Chinese or Western acupuncture system, for people aged 16 years and
10 over to manage chronic primary pain, **but only if** the course:

- 11
- 12 • is delivered in a community setting, and
 - 13 • is delivered by a band 7 (or lower) healthcare professional, and
 - 14 • is made up of no more than 5 hours of healthcare professional time (the
15 number and length of sessions can be adapted within these boundaries).

For a short explanation of why the committee made the recommendations on acupuncture and how they might affect practice, see [rationale and impact](#)

Full details of the evidence and the committee's discussion are in evidence review G: Acupuncture.

1 ***Electrical physical modalities for chronic primary pain***

2 1.3.6 Do not offer any of the following to people aged 16 years and over to
3 manage chronic primary pain:

- 4 • TENS
5 • ultrasound
6 • interferential therapy.

For a short explanation of why the committee made the recommendations on electrical physical modalities and how they might affect practice, see [rationale and impact](#)

Full details of the evidence and the committee's discussion are in evidence review H: Electrical physical modalities.

7

8 ***Manual therapy for chronic primary pain***

9 1.3.7 Be aware that there was not enough evidence on manual therapy for
10 chronic primary pain, so the committee made a [recommendation for](#)
11 [research](#).

For a short explanation of why the committee made a recommendation for research on manual therapy, see [rationale](#)

Full details of the evidence and the committee's discussion are in evidence review I: Manual therapy.

12

13 ***Pharmacological management of chronic primary pain***

14 1.3.8 Consider an antidepressant, either duloxetine, fluoxetine, paroxetine,
15 citalopram, sertraline or amitriptyline, for people aged 16 years and over

1 to manage chronic primary pain, after a full discussion of the benefits and
2 risks.

3
4 Note that this is an off-label use of these antidepressants. See [Prescribing](#)
5 [medicines](#) for more information.

6 1.3.9 For recommendations on reviewing treatments, see the [NICE guidelines](#)
7 [on medicines optimisation](#) and [medicines adherence](#).

8 1.3.10 For recommendations on stopping or reducing antidepressants , see the
9 [NICE guideline on depression in adults](#).

10 1.3.11 Do not offer any of the following, by any route, to people aged 16 years
11 and over to manage chronic primary pain:

- 12 • opioids
- 13 • non-steroidal anti-inflammatory drugs
- 14 • benzodiazepines
- 15 • anti-epileptic drugs including gabapentinoids, unless gabapentinoids
16 are offered as part of a clinical trial for complex regional pain
17 syndrome* (see [research recommendations](#))
- 18 • local anaesthetics, by any route, unless as part of a clinical trial for
19 complex regional pain syndrome (see [research recommendations](#))
- 20 • local anaesthetic/corticosteroid combinations
- 21 • paracetamol
- 22 • ketamine
- 23 • corticosteroids
- 24 • antipsychotics.

25
26 *Pregabalin and gabapentin are Class C controlled substances (under
27 the Misuse of Drugs Act 1971) and scheduled under the Misuse of
28 Drugs Regulations 2001 as Schedule 3. Evaluate patients carefully for
29 a history of drug misuse before prescribing and observe patients for
30 development of signs of misuse and dependence ([MHRA, Drug Safety](#)
31 [Update April 2019](#)).

1 1.3.12 If a person with chronic primary pain is already taking any of the
2 medicines in recommendation 1.3.11, explain the risks of continuing.

3 1.3.13 If a shared decision is made to stop antidepressants, opioids,
4 gabapentinoids or benzodiazepines, be aware of the problems associated
5 with withdrawal.

6 NICE is developing [a guideline on medicines associated with dependence or](#)
7 [withdrawal symptoms: safe prescribing and withdrawal management](#).

8 1.3.14 For recommendations on cannabis-based medicinal products, including
9 recommendations for research, see the [NICE guideline on cannabis-](#)
10 [based medicinal products](#).

For a short explanation of why the committee made the recommendations on pharmacological management and how they might affect practice, see [rationale and impact](#)

Full details of the evidence and the committee's discussion are in evidence review J: Pharmacological management.

11 ***Terms used in this guideline***

12 **Chronic pain**

13 Pain that persists or recurs for more than 3 months.

14 **Chronic primary pain**

15 Chronic primary pain is chronic pain in 1 or more anatomical regions that is
16 characterised by significant emotional distress (anxiety, anger/frustration or
17 depressed mood) or functional disability (interference in daily life activities and
18 reduced participation in social roles). Chronic primary pain is multifactorial:
19 biological, psychological and social factors contribute to the pain syndrome. The
20 diagnosis is appropriate unless another diagnosis would better account for the
21 presenting symptoms.

1 The ICD-11 definition of chronic primary pain includes chronic widespread pain,
2 complex regional pain syndrome, chronic primary headache or orofacial pain,
3 chronic primary visceral pain and chronic primary musculoskeletal pain.

4 **Recommendations for research**

5 The guideline committee has made the following recommendations for research.

6 ***Key recommendations for research***

7 **1 Pain management programmes for chronic pain**

8 What are the optimum characteristics of a clinically and cost-effective pain
9 management programme for people aged 16 years and over with chronic pain?

10 To find out why the committee made the research recommendation on pain
11 management programmes see [rationale](#).

12 **2 Psychological therapy – mindfulness for chronic primary pain**

13 What is the clinical and cost effectiveness of mindfulness therapy for managing
14 chronic primary pain in people aged 16 years and over?

15 To find out why the committee made the research recommendation on mindfulness
16 see [rationale](#).

17 **3 Psychological therapy – CBT for insomnia in chronic primary pain**

18 What is the clinical and cost effectiveness of cognitive–behavioural therapy (CBT) for
19 insomnia or CBT for insomnia and pain for managing chronic primary pain in people
20 aged 16 years and over?

21 To find out why the committee made the research recommendation on CBT for
22 insomnia see [rationale](#).

23 **4 Pharmacological interventions – gabapentinoids and local anaesthetics for 24 complex regional pain syndrome**

25 What is the clinical and cost effectiveness of gabapentinoids or local anaesthetics for
26 managing complex regional pain syndrome in people aged 16 years and over?

1 To find out why the committee made the research recommendation on
2 gabapentinoids and local anaesthetics see [rationale](#).

3 **5 Manual therapies for chronic primary pain**

4 What is the clinical and cost effectiveness of manual therapy for managing chronic
5 primary pain in people aged 16 years and over?

6 To find out why the committee made the research recommendation on manual
7 therapy see [rationale](#).

8 ***Other recommendations for research***

9 **Factors that may be barriers to successfully managing chronic pain**

10 What risk factors enable stratification of treatment for people aged 16 years and over
11 with chronic pain?

12 **Repeat courses of acupuncture for chronic primary pain**

13 What is the clinical and cost effectiveness of repeat courses of acupuncture or dry
14 needling for managing chronic primary pain in people aged 16 years and over?

15 **Psychotherapy for chronic primary pain**

16 What is the clinical and cost effectiveness of psychotherapy for managing chronic
17 primary pain in people aged 16 years and over?

18 **Relaxation therapy for chronic primary pain**

19 What is the clinical and cost effectiveness of relaxation therapies for managing
20 chronic primary pain in people aged 16 years and over?

21 **Social interventions for chronic pain**

22 What is the clinical and cost effectiveness of social interventions aimed at improving
23 the quality of life of people aged 16 years and over with chronic pain?

24 **Laser therapy for chronic primary pain**

25 What is the clinical and cost effectiveness of laser therapy for managing chronic
26 primary pain in people aged 16 years and over?

1 **Transcranial magnetic stimulation for chronic primary pain**

2 What is the clinical and cost effectiveness of transcranial magnetic stimulation for
3 managing chronic primary pain in people aged 16 years and over?

4 **Rationale and impact**

5 These sections briefly explain why the committee made the recommendations and
6 how they might affect practice. They link to details of the evidence and a full
7 description of the committee's discussion.

8 ***Assessing chronic pain***

9 Recommendations [1.1.1 to 1.1.8](#)

10 **Why the committee made the recommendations**

11 ***Possible barriers to successfully managing chronic pain***

12 There was not enough evidence to indicate whether any psychological, biological or
13 social factors predict successful pain management. The committee acknowledged
14 the importance of a comprehensive biopsychosocial approach to assessment and
15 management. They agreed that it is important for the healthcare professional to
16 understand how pain is affecting a person's life. A care plan should be based on the
17 effects of pain on day-to-day activities, while acknowledging that it is not possible to
18 predict what might happen in the future.

19 ***Communication between healthcare professionals and people with chronic
20 pain***

21 The committee agreed that the evidence on communication was in line with what
22 was generally considered best practice. However, evidence demonstrated
23 shortcomings in people's experience of consultations with healthcare professionals.
24 The committee agreed that this area needs addressing. They emphasised the
25 fundamental importance of good communication to the experience of care for people
26 with chronic pain, especially when many or all treatments are ineffective or not well
27 tolerated. The committee reviewed the recommendations from the NICE guideline on
28 patient experience in adult NHS services (CG138) alongside the qualitative evidence
29 to identify any areas needing specific recommendations for people with chronic pain.

1 They agreed that the heterogeneous, complex and potentially distressing nature of
2 the condition should be reflected in the recommendations. More specifically, a
3 comprehensive assessment should elicit an understanding of the effects of the pain,
4 and how this is viewed by the person and those around them. Understanding what is
5 important to the person is the first step in developing a care plan. The committee
6 agreed that it is important to explore a person's priorities, strengths, preferences,
7 interests and abilities, because these can help inform the plan.

8 The committee highlighted the importance of honesty about the uncertainty of the
9 prognosis, because the evidence suggested that this is valued by people with
10 chronic pain. Evidence showed that discussions about self-management often
11 happen late in the care pathway, or not at all. The committee considered that all
12 relevant management options should be considered at all stages of care, including
13 the first contact, and therefore made a recommendation to provide advice and
14 information, relevant to the person's individual preferences, at all stages of care, to
15 help them make decisions about managing their condition. Evidence showed that
16 normal or negative test results can be communicated in a way that is perceived as
17 being dismissive of pain. Therefore, the committee made a recommendation to
18 promote sensitivity around communicating test results.

19 **How the recommendations might affect practice**

20 The recommendations reflect best practice, but are currently implemented to varying
21 degrees across NHS settings and will involve a change of practice for some
22 providers. To fully implement these recommendations for people with chronic pain,
23 longer consultations or additional follow-up may be needed to discuss self-
24 management and treatment options.

25 Full details of the evidence and the committee's discussion are in evidence review A:
26 Factors that may be barriers to successfully managing chronic pain and evidence
27 review B: Communication between healthcare professionals and people with chronic
28 pain.

29 [Return to recommendations](#)

1 ***Pain management programmes for chronic pain***

2 [Research recommendation](#)

3 **Why the committee made the recommendations**

4 Evidence from 8 studies showed a very small improvement in quality of life with pain
5 management programmes led by professionals compared with usual care or waiting
6 list controls. However, benefits to quality of life were not consistent across studies
7 and there were no benefits observed in terms of physical function and psychological
8 distress. Where benefits were observed, they were only small. Differences in
9 programme delivery methods, including intensity, duration, components, structure,
10 and staffing, and aims meant that the committee were not able to determine whether
11 there was a particular content and characteristics of a programme that could be
12 effective. This, together with the uncertainty about cost effectiveness meant that the
13 committee were unable to make a recommendation for or against their use. The
14 committee decided to make a research recommendation to help determine the
15 elements that could make up an effective pain management programme for people
16 with chronic pain. They hoped that this research would inform future guidance.

17 Full details of the evidence and the committee's discussion are in evidence review C:
18 Pain management programmes.

19 [Return to recommendations](#)

20 ***Social interventions for chronic pain***

21 [Research recommendation](#)

22 **Why the committee made the recommendations**

23 No evidence was identified. The committee noted that provision of social prescribing
24 link workers is part of the NHS long term plan, and so there is already a move
25 towards social interventions within the NHS. The committee were aware of evidence
26 for social interventions in conditions other than chronic pain, but they agreed that this
27 evidence could not be extrapolated as the issues faced by people with chronic pain
28 are likely to be different from those populations. They could not make a
29 recommendation for chronic pain without evidence on clinical and cost effectiveness.
30 The committee decided to make a research recommendation to gather high-quality

1 evidence on social interventions in the NHS, specifically for adults with chronic pain.
2 This will hopefully inform future guidance.

3 Full details of the evidence and the committee's discussion are in evidence review D:
4 Social interventions.

5 [Return to recommendations](#)

6 ***Exercise for chronic primary pain***

7 [Recommendations 1.3.1 and 1.3.2](#)

8 **Why the committee made the recommendations**

9 Evidence from many studies showed that exercise reduced pain (23 studies) and
10 improved quality of life (22 studies) compared with usual care in people with chronic
11 primary pain. Benefit was seen for both short- and long-term follow-up and was
12 consistent across different types of exercise. Most of the evidence was for
13 professionally led supervised group exercise and for women with fibromyalgia or
14 people with chronic neck pain. There was limited evidence comparing different types
15 of exercise with each other although, from what was available, there was minimal
16 difference between the types. For this reason, the committee did not specify what
17 type of exercise should be used.

18 An economic model comparing exercise (all types) with no exercise was developed
19 for this guideline and showed that exercise was likely to be cost effective (both if
20 using only the time horizon of the trials and also when extrapolating the quality of life
21 gain beyond the trials). The analysis used studies in which exercise was
22 predominantly group based. The committee considered the results to be robust, and
23 agreed that the studies used in the model were generalisable to the whole evidence
24 review and chronic primary pain population. Exercise remained cost effective when
25 the assumed benefits and costs were varied (sensitivity analysis).

26 There were no negative effects demonstrated except for more people discontinuing
27 from exercise programmes. The committee agreed that people are more likely to
28 continue with exercise if the programme offered suits their lifestyle and physical
29 ability and addresses their individual health needs. They agreed that the choice of
30 programme as well as the content should take into account people's abilities and

1 preferences. This might include providing individual exercise advice for different
2 members of a group.

3 The committee's experience was that many people with chronic primary pain find it
4 difficult to be physically active. The committee agreed that it is important for these
5 people to continue physical activity after a formal exercise programme ends, but the
6 exercise should be sustainable for the person.

7 **How the recommendations might affect practice**

8 The types of exercise programmes currently offered vary from place to place, often
9 determined by the needs of the local population. In areas where supervised group
10 exercise is currently not provided, implementing the recommendation will lead to
11 increased resource use.

12 The committee discussed that the cost of engaging in physical activity after a formal
13 exercise programme ends would be a personal cost for people with chronic primary
14 pain, and would not fall to the NHS.

15 Full details of the evidence and the committee's discussion are in evidence review E:
16 Exercise.

17 [Return to recommendations](#)

18 ***Psychological therapy for chronic primary pain***

19 Recommendations [1.3.3 and 1.3.4](#)

20 **Why the committee made the recommendations**

21 ***ACT for chronic primary pain***

22 Most of the evidence showed that acceptance and commitment therapy (ACT)
23 improved quality of life and sleep, and reduced pain and psychological distress.

24 Although clinical evidence was from a fairly small number of studies, one economic
25 evaluation also showed ACT to be cost effective. The committee agreed that ACT
26 was likely to offer a good balance of benefits and costs and so recommended that it
27 should be considered as a psychological therapy for chronic primary pain. There was

1 not enough evidence to support a preference for ACT over cognitive–behavioural
2 therapy (CBT) or CBT over ACT.

3 ***CBT for chronic primary pain***

4 Most of the evidence showed that CBT for pain improved quality of life for people
5 with chronic primary pain. A consistent benefit was not demonstrated in other
6 outcomes, but the committee considered that the evidence may have
7 underestimated the benefits because the studies varied in terms of the level of
8 training of the therapists and the way the therapy was delivered. There was no
9 strong evidence of harm. Two economic evaluations also showed CBT to be cost
10 effective. The committee agreed that the evidence was not of high quality so they
11 decided to recommend that CBT (for pain) is considered.

12 Although there was some benefit of CBT for insomnia (CBT-I), particularly for quality
13 of life and sleep, the amount of evidence was smaller and did not include economic
14 evidence, so was insufficient to justify a recommendation. The committee agreed to
15 make a research recommendation for CBT-I to inform future guidance.

16 ***Biofeedback for chronic primary pain***

17 Evidence for biofeedback was conflicting, with little evidence of benefit and some
18 evidence of harm. For this reason, the committee decided that this should not be
19 offered as a management option for people with chronic primary pain.

20 ***Relaxation, mindfulness and psychotherapy for chronic primary pain***

21 There was not enough evidence for relaxation therapy, mindfulness or
22 psychotherapy for the committee to make recommendations, but what evidence
23 there was suggested there may be some benefit. The committee decided to make
24 research recommendations to inform future guidance.

25 ***Hypnosis, pain education and sleep hygiene for chronic primary pain***

26 Limited evidence showed little benefit of hypnosis and no clinically important effect of
27 pain education, but no evidence of harm. The committee noted that education should
28 be part of good clinical practice and is not specific to chronic primary pain. This is
29 already addressed by the NICE guideline on patient experience in adult NHS
30 services (CG138). It was also agreed that hypnosis is not widely used to manage

1 chronic primary pain in current clinical practice. The committee therefore decided not
2 to make recommendations for these therapies.

3 Limited evidence showed a benefit of sleep hygiene for improving quality of life,
4 sleep and pain. The committee considered that sleep hygiene is a component of
5 CBT-I and evidence showed that sleep hygiene was no more effective than CBT-I.
6 Therefore the committee decided not to make a recommendation for sleep hygiene.

7 **How the recommendations might affect practice**

8 The resource impact will depend on the uptake of the recommendations. CBT is
9 used in the NHS for chronic primary pain, although it is not standard practice
10 everywhere. ACT is a relatively new intervention but is also used to varying degrees
11 in practice. The costs of both interventions depend on the number and length of
12 sessions, whether they are group or individual (or face to face or virtual/online), and
13 who they are run by. Therefore costs can vary.

14 Biofeedback is usually used in physiotherapy as a method of monitoring progress,
15 rather than as a treatment in itself. The recommendation is therefore unlikely to have
16 a significant impact on current practice.

17 Full details of the evidence and the committee's discussion are in evidence review F:
18 Psychological therapy.

19 [Return to recommendations](#)

20 ***Acupuncture for chronic primary pain***

21 [Recommendation 1.3.5](#)

22 **Why the committee made the recommendations**

23 Many studies (27 in total) showed that acupuncture reduced pain and improved
24 quality of life in the short term (3 months) compared with usual care or sham
25 acupuncture. There was not enough evidence to determine longer term benefits. The
26 committee acknowledged the difficulty in blinding for sham procedures, but agreed
27 that the benefit compared with a sham procedure indicated a specific treatment
28 effect of acupuncture. There was a wide variation among the studies in the type and
29 intensity of the intervention used, and the studies were from many different

1 countries. The committee agreed that the type of acupuncture or dry needling should
2 depend on the individual needs of the person with pain.

3 Two economic evaluations (1 in the UK) showed that acupuncture offered a good
4 balance of benefits and costs for people with chronic neck pain. However, both
5 studies had limitations; a notable limitation being that the costs of acupuncture
6 seemed low. Threshold analysis based on these studies indicated the maximum
7 number of hours of a band 6 and 7 healthcare professional's time that would make
8 the intervention cost effective.

9 An original economic model was developed, which compared acupuncture with no
10 acupuncture. The model used data from studies with usual care comparisons, not
11 comparisons with sham acupuncture, because the committee agreed that a usual
12 care comparison in an economic model better reflects the real world benefit of the
13 intervention. The model showed that acupuncture was likely to be cost effective. The
14 committee considered the results to be robust, and agreed that the studies used in
15 the model were generalisable to the whole evidence review and chronic primary pain
16 population. Acupuncture remained cost effective when the assumed benefits and
17 costs were varied (sensitivity analysis).

18 Overall, the committee agreed that there was a large evidence base showing
19 acupuncture to be clinically effective in the short term (3 months); the original
20 economic modelling also showed it is likely to be cost effective. However, they were
21 uncertain whether the beneficial effects would be sustained long term and were
22 aware of the high resource impact of implementation. Taking these factors into
23 account, the committee made a recommendation to consider acupuncture or dry
24 needling for chronic primary pain, caveated by the factors likely to make the
25 intervention cost effective. These were: only if delivered in the community, and with a
26 maximum of 5 treatment hours (based on the average resource use in the trials in
27 the model and on the threshold analysis), and from a band 7 (or lower) healthcare
28 professional (based on the threshold analysis). The committee agreed that
29 discontinuing before this total amount of course time would be an option if the person
30 finds that the first few sessions are not effective.

1 No evidence was found to inform a recommendation for repeat courses of
2 acupuncture. The committee agreed that further research would help to inform future
3 practice and made a research recommendation.

4 **How the recommendations might affect practice**

5 There is variation in the availability and use of acupuncture for chronic primary pain,
6 with a recent reduction in these services. The recommendation is expected to lead to
7 increased use and need for acupuncture services and therefore to have a resource
8 impact. This is due to the number of people with chronic primary pain, and
9 acupuncture being an individual patient intervention and so staff intensive.

10 Full details of the evidence and the committee's discussion are in evidence review G:
11 Acupuncture.

12 [Return to recommendations](#)

13 ***Electrical physical modalities for chronic primary pain***

14 [Recommendation 1.3.6](#)

15 **Why the committee made the recommendations**

16 Limited evidence showed some benefit of electrical therapies for chronic primary
17 pain, but sample sizes were small and benefit beyond 3 months was unclear.

18 ***Laser therapy for chronic primary pain***

19 The exception was laser therapy, which showed a benefit for patient-reported pain
20 and quality of life in larger studies than for other electrical physical modalities.

21 However, the therapy used in the studies varied widely, particularly in terms of
22 wavelength, power, and the time the laser was applied to each painful area.

23 Evidence at more than 3 months' follow-up was limited, and there was no evidence
24 on cost effectiveness.

25 Taking into account the quality of the evidence, the limited long-term data and the
26 lack of evidence on cost effectiveness, the committee decided not to make a practice
27 recommendation for laser therapy. However, because the limited evidence was
28 promising, they agreed to make a research recommendation to inform future
29 guidance.

1 ***TENS, ultrasound and interferential therapy for chronic primary pain***

2 Limited evidence for TENS showed no clinically important difference compared with
3 sham TENS and usual care across several outcomes at less than 3 months, and no
4 longer term evidence was identified. There was no evidence for ultrasound or
5 interferential therapy. The committee noted these technologies have been around for
6 some time so it is unlikely that new research would be undertaken. These treatments
7 are being used by some in the NHS without evidence of benefit, so the committee
8 agreed that TENS, ultrasound and interferential therapy should not be offered for
9 chronic primary pain. Resources should be re-allocated to areas with more evidence
10 of clinical and cost effectiveness.

11 ***PENS and transcranial direct current stimulation for chronic primary pain***

12 There was a very limited amount of evidence for PENS and transcranial direct
13 current stimulation (TDCS), which suggested inconsistent benefits in some outcomes
14 only. The committee agreed this was insufficient for a recommendation. As neither
15 intervention is widely used in current practice for chronic primary pain, they did not
16 think further research was warranted.

17 Full details of the evidence and the committee's discussion are in evidence review H:
18 Electrical physical modalities.

19 [Return to recommendations](#)

20 ***Manual therapy for chronic primary pain***

21 [Research recommendation](#)

22 **Why the committee made the recommendations**

23 There was only a small amount of evidence available for each of the types of manual
24 therapy from studies of small sample sizes. The committee considered the lack of
25 evidence for the different types of manual therapy as well as the limitations of the
26 evidence. The committee were concerned about the quality of the evidence and the
27 variation in the type and intensity of the therapy. For example, vigorous soft tissue
28 techniques might be very similar in practice to mobilisation. For some types of
29 manual therapy, there was no evidence for outcomes beyond 3 months. The
30 committee were not able to draw any definite conclusions from the evidence about

1 the best type of manual therapy and so could not make recommendations for
2 practice. However, the committee agreed that the benefits compared with usual care
3 were promising and there was no evidence of harm. Therefore, they decided to
4 make a research recommendation.

5 Full details of the evidence and the committee's discussion are in evidence review I:
6 Manual therapy.

7 [Return to recommendations](#)

8 ***Pharmacological management for chronic primary pain***

9 [Recommendations 1.3.7 to 1.3.14](#)

10 **Why the committee made the recommendations**

11 ***Antidepressants for chronic primary pain***

12 Evidence indicated that antidepressants (duloxetine, amitriptyline and the SSRIs
13 fluoxetine, paroxetine, citalopram and sertraline) improved quality of life, pain and
14 psychological distress compared with placebo. But there were some limitations in the
15 quality and amount of the evidence. Most of the evidence was for women with
16 fibromyalgia. However, the committee agreed that for most medicines, response to
17 treatment would be sufficiently similar to allow recommendations to be made across
18 all chronic primary pain conditions, even when evidence was available for only one
19 condition. When the committee thought there was reason to distinguish between
20 chronic primary pain conditions, this is reflected in the recommendations.

21 The antidepressants were considered by class, but evidence was only available for
22 certain drugs within each class. The committee agreed these should be stated in the
23 recommendation. No evidence was identified that compared antidepressant classes
24 with each other, and the committee agreed that although there were some
25 inconsistencies in benefits observed between classes, they could not assume one
26 class to be more or less effective than another. Duloxetine (the only SNRI with
27 evidence for chronic primary pain) had a larger amount of long-term evidence of
28 effectiveness. However, due to the lack of head-to-head comparisons between the
29 antidepressant classes, the committee could not recommend duloxetine in
30 preference to the other antidepressants for which there was evidence. The decision

1 of which antidepressant to try should be based on a fully informed discussion with
2 the person with chronic primary pain, taking into account the risks and benefits.

3 Although none of the antidepressants have marketing authorisations for chronic
4 primary pain, there are no licensed alternatives for this indication and these
5 medications are already used in practice.

6 The committee agreed that the risk of withdrawal symptoms should be considered
7 when prescribing antidepressants and these should not be continued if they were not
8 effective. They recommended that the recommendations in the [NICE guideline on](#)
9 [depression in adults](#) should be followed if stopping or reducing antidepressants.

10 ***Cannabis-based medicinal products for chronic primary pain***

11 No evidence was identified on the effectiveness of cannabis-based products for
12 chronic primary pain, and some evidence suggested that the treatment could cause
13 adverse events in the short term. However, this was limited evidence from a small
14 study. Although the committee agreed that more research would be useful to inform
15 future practice, it was decided this was adequately covered within [the NICE guideline](#)
16 [on cannabis-based medicinal products](#).

17 ***Opioids for chronic primary pain***

18 No evidence was identified on the effectiveness of opioids for chronic primary pain.
19 Although there were limitations, evidence from non-randomised studies on the long-
20 term use (more than 6 months) of opioids for chronic pain suggested an increased
21 risk of dependence. Based on their experience, the committee agreed that even
22 short-term use of opioids could be harmful for a chronic condition. The lack of
23 evidence for effectiveness of opioids, along with evidence of long-term harm,
24 persuaded the committee to recommend against opioid use for people with chronic
25 primary pain.

26 ***Benzodiazepines and NSAIDs for chronic primary pain***

27 Limited evidence suggested a lack of benefit of benzodiazepines and non-steroidal
28 anti-inflammatory drugs (NSAIDs) for chronic primary pain. Evidence suggested that
29 psychological and physical functioning were poorer with benzodiazepines than with

1 placebo. Although there was no evidence for long-term use, the committee noted the
2 addictive properties of benzodiazepines and agreed to recommend against their use
3 for chronic primary pain.

4 Evidence suggested that short-term use of NSAIDs made no difference to people's
5 quality of life, pain or psychological distress. A small amount of evidence suggested
6 that NSAIDs reduced physical function, compared with placebo. In view of the risks
7 of harm with NSAIDs (gastrointestinal bleeding) and the lack of evidence of short-
8 term or long-term effectiveness, the committee decided to recommend against their
9 use for chronic primary pain.

10 ***Anti-epileptics for chronic primary pain***

11 Limited evidence suggested a lack of benefit of gabapentinoids for chronic primary
12 pain. No evidence was identified on the long-term safety of gabapentinoids, however
13 the committee were aware of reports of harm and risk of misuse and dependence
14 highlighted by the MHRA notification of the reclassification of gabapentinoids as a
15 class C substance controlled under the Misuse of Drugs Act 1971 and scheduled
16 under the Misuse of Drugs Regulations 2001 as schedule 3. There was no evidence
17 identified for any other anti-epileptics for chronic primary pain. Taking this into
18 account, alongside the lack of evidence of effectiveness compared with placebo, the
19 committee agreed to recommend against their use for chronic primary pain
20 generally. They were aware that gabapentinoids are currently recommended for
21 neuropathic pain and expert opinion within the committee suggested that complex
22 regional pain syndrome (CRPS) is sometimes understood as a neuropathic pain
23 disorder. Based on the expert opinion of some committee members they therefore
24 decided to make a research recommendation for the use of gabapentinoids for
25 CRPS to inform future practice.

26 ***Local anaesthetics for chronic primary pain***

27 Evidence for local anaesthetics was limited. A small amount of evidence for short-
28 term use suggested that there is either no benefit or that their use could result in
29 worse outcomes for pain than placebo. The committee therefore agreed to
30 recommend against the use of local anaesthetics for chronic primary pain. However,
31 based on the expert opinion of some members of the committee, it was noted that
32 local anaesthetics may be useful for people with CRPS who are under-represented

1 in randomised controlled trials. They therefore decided to make a research
2 recommendation for the use of local anaesthetics for CRPS to inform future practice.

3 ***Paracetamol, ketamine, corticosteroids, anaesthetic/corticosteroid***
4 ***combinations and antipsychotics for chronic primary pain***

5 No evidence was identified for paracetamol, ketamine, corticosteroids,
6 anaesthetic/corticosteroid combinations, or antipsychotics. From their own
7 experience, and from the summaries of product characteristics, the committee
8 agreed that these medicines have possible harms. The committee agreed that not
9 commenting on these medicines could result in their continued use in practice, which
10 would be inappropriate given the lack of evidence and possible harms, so they
11 recommended against the use of these treatments.

12 ***Withdrawing medicines***

13 The committee agreed that when recommendations had been made against the use
14 of medicines, there should be guidance for people who are already taking these.
15 They therefore included a recommendation based on expert opinion to explain the
16 risks of continuing a medicine, to inform a decision about whether the risks
17 outweighed the benefits and whether the medicine should be reduced or stopped. A
18 recommendation was also made to highlight possible withdrawal symptoms after
19 stopping some medicines.

20 ***How the recommendations might affect practice***

21 There is currently variation in the drugs used to treat chronic primary pain. The
22 recommendations are likely to have a resource impact in the short term because
23 there may be increased resource use from helping people to stop treatments,
24 particularly opioids and gabapentinoids. SNRI antidepressants are also slightly more
25 expensive than other types of antidepressant such as tricyclics, but this does depend
26 on dose. In the longer term, the recommendations should reduce the use of drugs for
27 managing chronic primary pain, with a consequent reduction in harms and cost
28 savings. This is likely to have wider benefits both to an individual and to society by,
29 for example, enabling people to return to the workforce.

1 Full details of the evidence and the committee's discussion are in evidence review J:
2 Pharmacological management.

3 [Return to recommendations](#)

4 **Context**

5 Chronic pain is often difficult to treat. It can be associated with many different types
6 of tissue injuries and disease processes. Sometimes no underlying disease can be
7 found. Pain has a significant impact on individuals and their families and carers. It
8 affects mood, sleep, mobility, role within the family, ability to work and other aspects
9 of life. Current mood, anxiety about pain, previous experience of pain, and
10 unpleasant life events can influence how pain is perceived.

11 ***Key facts and figures***

12 The prevalence of chronic pain has been difficult to define: but estimates range from
13 8.7% to 64.4%, with a pooled mean of 31%. In the UK chronic pain may affect
14 between one-third and one-half of the population, but it is not known what proportion
15 of people meeting the criteria for chronic pain either need or wish for treatment.

16 Almost half of people with chronic pain have a diagnosis of depression and two-
17 thirds of people are unable to work outside the home. Studies of disability in relation
18 to a number of long-term health conditions show that pain contributes the most to
19 disability measures.

20 Attempts to treat chronic pain are costly to the healthcare system. In 2016,
21 £537 million was spent on prescribing analgesics, with at least an additional 50%
22 cost incurred from the prescription of other drug classes such as antidepressants
23 and anti-epileptics. Further healthcare costs include visits to primary care, referrals
24 to secondary care for opinions from specialists, and costs of investigations and
25 interventions, including surgery.

26 The economic impact of pain is high due to absenteeism, poor productivity and
27 people with pain leaving the workforce. Painful conditions such as arthritis and back
28 pain account for one-third of all claims for disability benefits in the UK. The annual

1 indirect (productivity) cost of back pain in the UK was estimated to be between
2 £5 billion and £10.7 billion.

3 ***Current practice***

4 There is no medical intervention, pharmacological or non-pharmacological, that is
5 helpful for more than a minority of people with chronic pain, and benefits of
6 treatments are modest in terms of effect size and duration. Additional morbidity
7 resulting from treatment for chronic pain is not unusual, so it is important to evaluate
8 the treatments we offer for chronic pain, to focus resources appropriately and to
9 minimise harm.

10 The complexity of chronic pain and the association with significant distress and
11 disability can influence clinical interactions. People often expect a clear diagnosis
12 and effective treatment, but these are rarely available. GPs and specialists in other
13 fields find chronic pain very challenging to manage and often have negative
14 perceptions of people with pain. This is despite the fact that in every specialty there
15 are some people with chronic pain. This can have important consequences for the
16 therapeutic relationship between healthcare professionals and patients.

17 A clear understanding of the evidence for the effectiveness of chronic pain
18 treatments:

- 19 • improves the confidence of healthcare professionals in their conversations about
20 pain, and
- 21 • helps healthcare professionals and patients to have realistic expectations about
22 outcomes of treatment.

23 **Finding more information and resources**

24 To find out what NICE has said on topics related to this guideline, see our web
25 pages on [neurological conditions](#) and [musculoskeletal conditions](#) .

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