Relieving patients' pain with expectation interventions

Citation for published version (APA):

Peerdeman, K. J., van Laarhoven, A. I. M., Keij, S. M., Vase, L., Rovers, M. M., Peters, M. L., & Evers, A. W. M. (2016). Relieving patients' pain with expectation interventions: a meta-analysis. *Pain*, 157(6), 1179-1191. <https://doi.org/10.1097/j.pain.0000000000000540>

Document status and date:

Published: 01/06/2016

DOI:

[10.1097/j.pain.0000000000000540](https://doi.org/10.1097/j.pain.0000000000000540)

Document Version:

Publisher's PDF, also known as Version of record

Document license:

Taverne

Please check the document version of this publication:

- A submitted manuscript is the version of the article upon submission and before peer-review. There can be important differences between the submitted version and the official published version of record. People interested in the research are advised to contact the author for the final version of the publication, or visit the DOI to the publisher's website.
- The final author version and the galley proof are versions of the publication after peer review.
- The final published version features the final layout of the paper including the volume, issue and page numbers.

[Link to publication](#)

General rights

Copyright and moral rights for the publications made accessible in the public portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

- Users may download and print one copy of any publication from the public portal for the purpose of private study or research.
- You may not further distribute the material or use it for any profit-making activity or commercial gain
- You may freely distribute the URL identifying the publication in the public portal.

If the publication is distributed under the terms of Article 25fa of the Dutch Copyright Act, indicated by the "Taverne" license above, please follow below link for the End User Agreement:

www.umlib.nl/taverne-license

Take down policy

If you believe that this document breaches copyright please contact us at:

repository@maastrichtuniversity.nl

providing details and we will investigate your claim.

Relieving patients' pain with expectation interventions: a meta-analysis

Kaya J. Peerdeman^{a,b,*}, Antoinette I.M. van Laarhoven^{a,b}, Sascha M. Keij^a, Lene Vase^{c,d}, Maroeska M. Rovers^e, Madelon L. Peters^f, Andrea W.M. Evers^{a,b,g}

Abstract

Patients' expectations are important predictors of the outcome of analgesic treatments, as demonstrated predominantly in research on placebo effects. Three commonly investigated interventions that have been found to induce expectations (verbal suggestion, conditioning, and mental imagery) entail promising, brief, and easy-to-implement adjunctive procedures for optimizing the effectiveness of analgesic treatments. However, evidence for their efficacy stems mostly from research on experimentally evoked pain in healthy samples, and these findings might not be directly transferable to clinical populations. The current meta-analysis investigated the effects of these expectation inductions on patients' pain relief. Five bibliographic databases were systematically searched for studies that assessed the effects of brief verbal suggestion, conditioning, or imagery interventions on pain in clinical populations, with patients experiencing experimental, acute procedural, or chronic pain, compared with no treatment or control treatment. Of the 15,955 studies retrieved, 30 met the inclusion criteria, of which 27 provided sufficient data for quantitative analyses. Overall, a medium-sized effect of the interventions on patients' pain relief was observed (Hedges $g = 0.61$, $I^2 = 73\%$), with varying effects of verbal suggestion ($k = 18$, $g = 0.75$), conditioning (always paired with verbal suggestion, $k = 3$, $g = 0.65$), and imagery ($k = 6$, $g = 0.27$). Subset analyses indicated medium to large effects on experimental and acute procedural pain and small effects on chronic pain. In conclusion, patients' pain can be relieved with expectation interventions; particularly, verbal suggestion for acute procedural pain was found to be effective.

Keywords: Meta-Analysis, Systematic review, Expectation, Expectancy, Placebo effect, Verbal suggestion, Conditioning, Imagery, Pain, Analgesia, Patients

1. Introduction

Expectations are important predictors of the outcome of analgesic treatments.^{16,27,58,77} As posed in the expectancy theory,^{43,44} expectations of pain relief can directly elicit and/or enhance actual pain relief. The importance of expectations has particularly become clear in research on placebo effects, of which expectancy is believed to be a core mechanism.^{7,43,44,69} Placebos, such as sugar pills and saline injections, have

repeatedly been found to provide pain relief, with effects at both subjective^{80,81} and neurobiological levels.^{3,71} These and other findings suggest that interventions that induce expectations of pain relief, ie, analgesic expectation inductions, are promising for optimizing the effectiveness of standard analgesic treatments in clinical practice. However, evidence for the efficacy of expectation inductions stems mostly from laboratory research using experimental pain in samples of healthy participants, whereas research in clinical samples (eg, patients with chronic back pain or postoperative pain) is limited. Although experimentally evoked pain in healthy samples is generally considered a good model for clinical pain, these findings might not be directly transferable to clinical populations. On the one hand, patients with pain, especially chronic pain, have a more extensive and complex history of pain and, often unsuccessful, pain treatment. This might make them more resistant to expectation interventions.^{23,41} On the other hand, patients are likely to have a higher desire for pain relief, possibly making them more sensitive to expectation interventions.^{24,38,70,82}

Three common, brief, and easy-to-implement interventions that have been found to induce and/or enhance expectations are promising for implementation in clinical practice: verbal suggestion, conditioning, and imagery. Verbal suggestion entails instructions regarding treatment outcomes given by, for example, a health care provider. Verbal suggestions, such as saying that a placebo or active treatment is an effective analgesic, can induce expectations of pain relief and produce corresponding experiences of pain relief.^{4,74,76} Conditioning entails the pairing of a neutral stimulus with an unconditioned stimulus that triggers

Sponsorships or competing interests that may be relevant to content are disclosed at the end of this article.

^a Unit Health, Medical and Neuropsychology, Leiden University, Leiden, the Netherlands, ^b Leiden Institute for Brain and Cognition, Leiden University, Leiden, the Netherlands, ^c Department of Psychology and Behavioural Sciences, Aarhus University, Aarhus, Denmark, ^d Danish Pain Research Center, Aarhus University Hospital, Aarhus, Denmark, ^e Radboud Institute for Health Sciences, Radboud University Medical Center, Nijmegen, the Netherlands, ^f Department of Clinical Psychological Science, Maastricht University, Maastricht, the Netherlands, ^g Department of Medical Psychology, Radboud University Medical Center, Nijmegen, the Netherlands

*Corresponding author. Address: Unit Health, Medical and Neuropsychology, Leiden University, PO Box 9555, 2300 RB Leiden, the Netherlands. Tel.: +31-71-527-3622; fax: +31-71-527-3619. E-mail address: k.j.peerdeman@fsw.leidenuniv.nl (K.J. Peerdeman).

Supplemental digital content is available for this article. Direct URL citations appear in the printed text and are provided in the HTML and PDF versions of this article on the journal's Web site (www.painjournalonline.com).

PAIN 157 (2016) 1179–1191

© 2016 International Association for the Study of Pain
<http://dx.doi.org/10.1097/j.pain.0000000000000540>

a certain response. For example, pairing a placebo treatment with reduced pain stimulation can produce expected and experienced pain relief when merely receiving the placebo treatment,^{1,15,45,46,59,78} especially when conditioning is paired with a verbal suggestion.^{6,46,52} Mental imagery of a future event or desired outcome entails actively generating a multisensory cognitive representation of an event and often involves relatively implicit suggestions.^{28,35} For example, imagining an optimal future self or health can increase general positive expectations (ie, optimism)^{56,60,61} and correspondingly reduce pain and medical care utilization.^{29,42} Thus, inducing expectations of pain relief, through verbal suggestion, conditioning, and imagery, can reduce pain. However, the comparative effectiveness of these expectation inductions, particularly in clinical populations, is mostly unclear.

The primary aim of the current meta-analysis was to investigate the effects of brief and easy-to-implement expectation interventions for relieving patients' pain. Specifically, the effects of verbal suggestion, conditioning, and imagery on pain relief in clinical populations are investigated. Furthermore, we compared the effects on experimental vs clinical pain, and acute procedural (pain during or directly after a medical procedure, eg, post-operative pain) vs chronic (long-lasting pain associated with a medical condition, eg, chronic back pain or recurrent migraine) clinical pain. Additional outcome analyses explored the effects on expected pain, affective pain, and anxiety.

2. Methods

2.1. Protocol and registration

The systematic review and meta-analysis were performed in accordance with the PRISMA Statement⁵⁷ and the recommendations of the Cochrane Collaboration.³² The study protocol was registered in the international prospective register of systematic reviews Prospero (CRD42013006575).

2.2. Information sources and search strategy

The electronic bibliographic databases PubMed, PsycINFO, EMBASE, Cochrane CENTRAL, and the Cochrane Methodology Register were searched from inception until June 19, 2015, using search terms describing the 3 expectation inductions and pain (see Supplementary File 1 for the full search strategy, available online as Supplemental Digital Content at <http://links.lww.com/PAIN/A249>). The search was restricted to humans when possible in the databases. In addition, the reference lists of eligible studies and studies that cited the eligible studies were searched for relevant articles.

2.3. Eligibility criteria

Studies were included if they assessed the effect of 1 of the 3 expectation inductions (verbal suggestion, conditioning, or imagery) on pain relief in a clinical sample (ie, adult patients with a somatic condition and/or undergoing medical treatment). The review was restricted to studies that compared a brief intervention (verbal suggestion, conditioning, or imagery; maximum 1 day) that was believed to induce expectations of pain relief to a control condition consisting of no treatment/treatment as usual, or a control treatment that was believed to not induce expectations of pain relief. If the studied intervention consisted of multiple components (eg, both imagery and relaxation), the expectation induction had to be the main component of the intervention (ie, duration > 50% of intervention time). Studies in which uncertain

expectations of pain relief (eg, 50/50 chance of receiving active or inactive treatment like during blinded treatment administration) were induced in either condition were excluded. Experimental (ie, experimentally evoked pain), acute procedural (ie, pain during or directly after a medical procedure), or chronic (ie, long-lasting pain associated with a medical condition) pain had to be assessed with a self-report rating scale that provided numerical values of experienced pain intensity (eg, visual analogue scale). Only original research results that were presented in full-length English language empirical articles were included (ie, not abstracts, case studies, reviews, and reanalyses).

2.4. Study selection

Titles and abstracts of articles retrieved using the search strategy were screened by 1 of 2 review authors (K.J.P. or S.M.K.) to identify studies that potentially met the eligibility criteria outlined above. The full texts of these articles were retrieved (online, through Dutch academic libraries, or through study authors) and assessed for eligibility (K.J.P. or S.M.K.). Full texts that were considered to be eligible for inclusion or about which doubts existed were also assessed for eligibility by a second review author (K.J.P. or S.M.K.). Any remaining doubts were resolved through discussion with other review authors (A.W.M.E., A.I.M.v.L., and L.V.).

2.5. Data extraction

A standardized form was independently used by 2 review authors (K.J.P. and S.M.K.) to extract data regarding the following from the included studies: expectation induction, control condition, study design, study population, type of pain, and pain outcome measure. Statistical data for meta-analysis (ie, sample size, mean, and SD of all postintervention pain measurements and secondary outcomes, or alternative values) were extracted by 1 review author (K.J.P.), and accuracy was checked by a second review author (S.M.K.). If it was not possible to extract sufficient data for the calculation of postintervention effect sizes for the primary and secondary outcomes, the study authors were contacted. When sufficient data could not be acquired, alternative statistics (eg, SE, confidence interval [CI], *t* or *F* value, *P* value, or mean change scores) were inspected. When appropriate alternative statistics were available, effect sizes were calculated using these, otherwise the study was excluded from quantitative analysis (Table 1).

2.6. Risk of bias assessment

Risk of bias within each of the included studies was assessed independently by 2 review authors (K.J.P. and S.M.K.) with the Cochrane risk of bias tool, version 5.1.0.³² The following items were evaluated at study level: "Random sequence generation" (selection bias), "Allocation concealment" (selection bias), "Incomplete outcome data" (attrition bias), "Selective outcome reporting" (reporting bias), and "Other bias" (focused on differences in sample characteristics—sex, age, and baseline pain). A priori, it was decided not to judge the items "Blinding of participants and personnel" (performance bias) and "Blinding of outcome assessors" (detection bias), because it is not possible to blind participants to the expectation inductions or to blind outcome assessors for self-reported outcomes. Disagreements between the authors regarding judgment of the risk of bias were resolved by discussion, with involvement of a third review author (A.I.M.v.L.) where necessary.

Table 1**Study characteristics of all studies included in the quantitative and qualitative meta-analysis.**

Study Author	Intervention			Sample		Outcome	
	Intervention (route of administration)	Control condition	Comparison	N	Patient population	Pain type	Timing of pain measurement
Verbal suggestion Amanzio et al. (2001) ²	Verbal suggestion referring to active treatment* (injection)	Control treatment	Between subjects	73/69	Patients undergoing thoracic surgery	Acute procedural pain (postoperative pain)	15, 30, 45, and 60 min after intervention†
Benedetti et al. (1995) ⁸	Verbal suggestion referring to placebo treatment* (injection)	Control treatment	Between subjects	13/11	Patients undergoing thoracic surgery	Acute procedural pain (postoperative pain)	15 and 60 min after intervention‡
Benedetti et al. (2003) ¹⁰	Verbal suggestion referring to active treatment* (injection)	Control treatment	Between subjects	21/21	Patients undergoing thoracic surgery	Acute procedural pain (postoperative pain)	30 and 60 min after intervention†
Benedetti et al. (2006) ⁹	Verbal suggestion referring to active treatment* (cutaneous)	Control treatment	Within subjects	28	Patients with Alzheimer's disease	Acute procedural pain (pain after venipuncture)	15 min after intervention§
Bialosky et al. (2014) ¹¹	Verbal suggestion referring to placebo treatment (other)	No treatment	Between subjects	27/28	Patients with low back pain	Experimental pain (mechanical pain)	1 × postintervention
Charron et al. (2006) ¹³	Verbal suggestion referring to placebo treatment (injection)	Control treatment	Between subjects	8/8	Patients with chronic low back pain	Chronic pain (low back pain)	Every 2 min for 20 min after intervention‡¶
de Craen et al. (2001) ¹⁸	Verbal suggestion referring to placebo or active treatment (oral)	Control treatment	Between subjects	55/56	Patients with chronic pain	Chronic pain (chronic pain)	0.5, 1, 2, 4, 6, 8, and 24 h after intervention†
Gryll and Katahn (1978) ²⁶	Verbal suggestion referring to placebo treatment (oral)	No treatment	Between subjects	40/40	Patients undergoing dental surgery	Acute procedural pain (injection pain)	1 × postintervention
Hashish et al. (1988) ³⁰	Verbal suggestion referring to placebo treatment (other)	No treatment	Between subjects	25/25	Patients undergoing dental surgery	Acute procedural pain (postoperative pain)	24 h after intervention
Ho et al. (1988) ³⁴	Verbal suggestion referring to placebo treatment (other)	No treatment	Between subjects	16/16	Patients undergoing dental surgery	Acute procedural pain (postoperative pain)	1 × postintervention#
Kam-Hansen et al. (2014) ³⁹	Verbal suggestion referring to placebo or active treatment (oral)	Control treatment	Within subjects	120	Patients with migraine (episodic)	Chronic pain (migraine pain)	2 h after intervention
Levine and Gordon (1984) ⁵¹	Verbal suggestion referring to placebo treatment* (injection)	Control treatment	Between subjects	12/12	Patients undergoing dental surgery	Acute procedural pain (postoperative pain)	50 min after intervention‡
Liberman (1964) ⁵²	Verbal suggestion referring to placebo treatment (injection)	No treatment	Between subjects	51/30	Patients undergoing labor	Acute procedural pain (labor pain)	15 and 30 min during labor#
Petersen et al. (2012) ⁶⁴	Verbal suggestion referring to active treatment* (cutaneous)	Control treatment	Within subjects	19	Patients with neuropathic pain	Chronic pain (spontaneous neuropathic pain)	1 × postintervention
Petersen et al. (2014) ⁶³	Verbal suggestion referring to active treatment* (cutaneous)	Control treatment	Within subjects	18	Patients with neuropathic pain	Chronic pain (ongoing neuropathic pain)	1 × postintervention
Pollo et al. (2003) ⁶⁵	Verbal suggestion referring to placebo treatment (injection)	No treatment	Between subjects	20/17	Patients undergoing assessment of autonomic functions	Experimental pain (electrical pain stimulus)	1 × postintervention
Price et al. (2007) ⁶⁸	Verbal suggestion referring to placebo treatment (other)	No treatment	Within subjects	9	Patients with irritable bowel syndrome	Experimental pain (rectal distension pain)	Last 5 of 7 consecutive stimuli¶
Schmid et al. (2015) ⁷³	Verbal suggestion referring to placebo treatment (injection)	Control treatment	Within subjects	17	Patients with irritable bowel syndrome	Experimental pain (rectal distension pain)	8 distensions postintervention†
Vase et al. (2003) ⁸²	Verbal suggestion referring to placebo treatment (other)	No treatment	Within subjects	13	Patients with irritable bowel syndrome	Experimental pain (rectal distension pain)	5, 15, 20, 40, and 50 min after intervention**
Vase et al. (2005) ⁸³	Verbal suggestion referring to placebo treatment (other)	No treatment	Within subjects	16	Patients with irritable bowel syndrome	Experimental pain (rectal distension pain)	5, 10, 15, 20, 25, 30, 35, and 40 min after intervention† (each time mean pain of 2 distensions)

(continued on next page)

Table 1 (continued)

Study	Intervention	Sample		Outcome			
Author	Intervention (route of administration)	Control condition	Comparison	N	Patient population	Pain type	Timing of pain measurement
Conditioning							
Hashmi et al. (2014) ³¹	Conditioning with verbal suggestion referring to placebo or active treatment (other)	Control treatment	Within subjects	42	Patients with knee osteoarthritis	Experimental pain (heat pain)	2 × 6 stimuli postintervention¶
Klinger et al. (2007) ⁴⁷	Conditioning with verbal suggestion referring to placebo treatment (cutaneous)	Control treatment	Between subjects	12/12	Patients with atopic dermatitis	Experimental pain (electrical pain stimuli)	5 consecutive stimuli postintervention†
Laska and Sunshine (1973) ⁴⁹	Conditioning referring to placebo treatment (after active treatment) (oral)	Control treatment	Between subjects	95/16	Patients with postoperative, fracture, or somatic pain	Acute procedural pain or chronic pain (postoperative, fracture, or somatic pain)	30, 60, 120, 180, 240, 300, and 360 min after intervention#
Lee et al. (2012) ⁵⁰	Conditioning with verbal suggestion referring to placebo treatment (injection)	Control treatment	Within subjects	17	Patients with irritable bowel syndrome	Experimental pain (rectal distension pain)	1 × postintervention
Imagery							
Danhauer et al. (2007) ¹⁷	Imagery of sending warm energy to painful areas and of a pleasant place, and relaxation instructions (audio recording)	No treatment††	Between subjects	56/58	Patients undergoing colposcopy	Acute procedural pain (pain during colposcopy)	1 × postintervention (retrospect, ie, pain during procedure)
Foji et al. (2015) ²²	Imagery contents not reported (audio recording)	No treatment	Between subjects	31/31	Patients undergoing coronary angiography	Acute procedural pain (postangiography pain)	1 × postintervention
Gonzales et al. (2010) ²⁵	Imagery and progressive relaxation, biorhythmic music with positive statements (audio recording)	No treatment	Between subjects	22/22	Patients undergoing head and neck surgery	Acute procedural pain (postoperative pain)	1 and 2 h after intervention†
Jacobson (2006) ³⁷	Imagery of a pleasant place and cooling gloves (audio recording)	No treatment††	Between subjects	41/40	Patients undergoing peripheral i.v. therapy	Acute procedural pain (i. v. insertion pain)	1 × postintervention
Kwekkeboom et al. (2008) ⁴⁸	Imagery using glove anesthesia technique, transferring feeling of numbness to painful areas (audio recording)	Control treatment	Within subjects	31	Patients with cancer pain	Chronic pain (cancer pain)	1 × postintervention
Wells et al. (1989) ⁸⁴	Imagery of transferring feeling of numbness to painful area (audio recording)	Control treatment	Between subjects	10/10	Patients undergoing abortion	Acute procedural pain (abortion pain)	1 × worst pain during abortion, and 1 × pain in recovery room†

* Used open/hidden design.

† Average of effect sizes across time points is calculated.

‡ Only change from baseline available.

§ Postintervention score(s) calculated.

|| Within-subjects comparison also possible.

¶ Only average across time points available.

Insufficient data for meta-analysis.

** Only effect size (Cohen *d*) available.

†† Described by study authors as treatment as usual.

i.v., intravenous; N, either total sample size or sample size per condition (intervention/control condition).

2.7. Considerations regarding data selection

The following choices were made regarding the selection of intervention and control conditions. When a study contained multiple relevant intervention or control conditions, data were selected from the intervention most directly aimed at pain reduction,⁸⁴ the comparison of the most active expectation induction (eg, strongest verbal suggestion) vs the most passive control condition (eg, no treatment),^{26,30,34,39,47,51,64,84} or the control condition conducted before rather than after the intervention.⁶⁸ In 2 studies, the control condition involving hidden

administration of active medication was chosen rather than a no-treatment control condition, to avoid confusion with the effect of the active medication.^{63,64} With regard to the study design, between-subjects comparisons were included in the quantitative analyses if possible,^{13,47} because the majority of studies used a between-subjects design. With regard to the outcome measures, in the 4 studies that included several pain measures,^{13,63,64,82} the data of the most clinically relevant type of pain were included (eg, evoked visceral pain rather than evoked heat pain in patients with irritable bowel syndrome).

See Supplementary File 2 for an overview of the additional conditions and pain measures used in each study (available online as Supplemental Digital Content at <http://links.lww.com/PAIN/A249>).

2.8. Data analysis

All analyses were conducted by the first reviewer (K.J.P) and checked by a second reviewer (S.M.K.), using Comprehensive Meta-Analysis software, version 3.3.070 (Biostat, Englewood, CO). The effect size (Hedges g) was calculated as the mean postintervention pain intensity score for the control condition minus the mean postintervention pain intensity score for the intervention condition, divided by the pooled SD, and weighted according to the number of subjects in each study.³² When pain was assessed at multiple postintervention time points, the average effect across these time points was calculated. Positive values for g indicate lower postintervention pain ratings (or secondary outcome values, eg, expected pain) in the intervention condition than in the control condition. A value around 0.2 to 0.3 was considered a small effect, a value around 0.5 a medium effect, and a value of 0.8 or larger a large effect.¹⁴ The pooled effects were analyzed using a random-effects model, given the variability in research characteristics (eg, different expectation inductions and types of pain). The presence and magnitude of heterogeneity were assessed with the I^2 statistic, as well as by visual inspection of the forest plot. I^2 values of 25%, 50%, and 75% can be considered to indicate low, moderate, and high degrees of heterogeneity, respectively.³³ For within-subjects comparisons, the intervention–control condition correlation coefficient could not be derived from the included studies; therefore, an r of 0.5 was imputed. For subset analyses, τ^2 was not pooled because we did not expect the between-study variance to be the same for all subsets. The effect sizes in the subsets were compared descriptively rather than with statistical tests, given the small number of studies in most subsets (ie, insufficient statistical power). Meta-analysis was conducted only when the data of at least 3 studies were available.

The pooled effects of all 3 expectation inductions (verbal suggestion, conditioning, and imagery) were analyzed together and separately. Planned subset analyses compared the effects on different types of pain (experimental vs clinical pain, and acute procedural vs chronic clinical pain), which also served as a proxy for differential effects depending on the patient type (patients with somatic condition vs those undergoing medical treatment). Post hoc subset analyses assessed the influence of the route of treatment administration (oral, injection, cutaneous, and other) and compared studies using active (eg, analgesic medication) vs placebo (eg, saline injection) treatments. Additional outcome analyses explored the effects of the expectation inductions on expected pain, affective pain, and anxiety. Sensitivity analyses assessed the stability of the overall effect size in relation to: (1) the risk of bias within studies (by removing studies for which at least 1 item was judged to involve a high risk of bias); (2) publication bias (inspection of the funnel plot and trim and fill method); (3) the comparison with a control condition with or without a control treatment, as well as the inclusion of control treatments that might have induced some expectations; (4) the inclusion of both between-subjects and within-subjects comparisons; (5) the imputed intervention–control condition correlation coefficient (imputed $r = 0.5$, vs $r = 0.1$ or $r = 0.9$); (6) the inclusion of postintervention rather than change scores.

3. Results

3.1. Study selection

See **Figure 1** for the flowchart of the selection process. Through the initial search in the databases, 15,952 records were retrieved, 3 additional relevant studies were identified through other sources. Of these, 3678 records were duplicates, 11,835 records were excluded on the basis of screening of the titles/abstracts, and the full text of 15 studies that were considered possibly relevant was not available. The full texts of 427 records were retrieved. Of the 62 full texts that were initially selected, 32 studies were excluded for various reasons (eg, induction of negative expectations or no control condition). In total, 30 studies were included in the qualitative synthesis. For three studies, a measure of variance (eg, SD) was missing for the primary outcome (pain intensity).^{34,49,52} Sufficient data of 27 studies were available for meta-analysis.

3.2. Study characteristics

The characteristics of all included studies are reported in **Table 1**. The majority of the studies that could be included in the quantitative meta-analysis assessed the analgesic effects of verbal suggestions (67%, $k = 18$) such as “The agent you have just been given is known to powerfully reduce pain in some patients”^{63,64,68,83} and “This drug is a local anesthetic and we use it to reduce the pain of the next stimulus. It takes a couple of minutes to work. Rest assured, the next stimulus will be less painful.”⁶⁵ Three studies assessed the effects of a conditioning procedure on pain, which was always combined with verbal suggestion of analgesic effects. Six studies assessed the effects of imagery, with images of pain reduction in four studies (eg, by imagining numbness). The images used in the other 2 studies were not specified. Regarding the presence of multiple intervention components, we note that the intervention in 4 of the imagery studies incorporated relaxation instructions, to maximally engage participants in imagery. In no other studies there were indications of components of the interventions that could not be qualified as an expectation induction in themselves. Because verbal suggestions are inherently incorporated in almost all types of psychological interventions, suggestions were probably included in the studied imagery interventions. In total, 1256 patients participated in the selected groups of the studies. The samples consisted of patients with various pathologies, eg, patients with irritable bowel syndrome ($k = 5$) and patients experiencing long-lasting pain such as chronic back pain or recurrent migraine ($k = 8$). For most studies, measurements of clinical pain could be included: acute procedural pain (eg, postoperative pain) was assessed in 12 studies, and chronic pain (eg, chronic back pain, including cancer pain) in six studies. Measurements of experimentally evoked pain (eg, electrical pain stimuli) were included in 9 studies. In all studies, patients reported their pain on a single-item pain scale (see Supplementary File 2, available online as Supplemental Digital Content at <http://links.lww.com/PAIN/A249>).

3.3. Description risk of bias within studies

Figures 2 and 3 show the results of the risk of bias (RoB) assessment in all included studies. Regarding selection bias, 63% of the studies reported that treatment allocation was random, but only 27% described adequate random sequence generation (low RoB). Randomization was not mentioned in 17% of the studies (unclear RoB), and incomplete or not performed at all in 20% of the studies (high RoB). Allocation concealment was

Figure 1. PRISMA flow diagram showing study selection process, including reasons for exclusion. Selection was conducted by 1 reviewer, unless otherwise stated.

reported adequately in only 13% of the studies (low RoB); in 1 study, allocation concealment was described, but insufficiently (unclear RoB). None of the other studies mentioned allocation concealment, but a high RoB was inferred if randomization was incomplete or not performed at all (20%). In 40% of the studies,

there were no signs of attrition bias due to incomplete outcome reporting (low RoB). For 10% of the studies, dropout was unbalanced and/or related to the outcome measure (high RoB). The judgment of reporting bias was challenged for the majority of studies (93%) because no preregistered study protocol could be

Figure 2. Risk of bias graph: review authors' judgments about each risk of bias item presented as percentages across all included studies. For the item "Selective outcome reporting" (reporting bias), the absence of a preregistered study protocol did not affect the judgment because a protocol was absent for 93% of the studies.

retrieved. When disregarding the presence of a protocol in the assessment (**Figs. 2 and 3**), 63% of the studies could be judged as having a low RoB. For 1 study, there was discordance between some measures mentioned in the methods and results section, whereas in another study, analyses did not include all available measurements of the primary outcome (high RoB). In 30% of the studies, no imbalances in sample characteristics of sex, age, and baseline pain were observed (low risk of "other bias"). All other studies (70%) reported insufficient data regarding equality of one or more of these sample characteristics (unclear RoB). Last, the risk of "other bias" was judged to be high in 1 study because of different study procedures in the intervention and control condition and in another study because of the insufficient reporting of study details such as the characteristics of the pain-reporting scale.

3.4. Primary meta-analysis: effect of expectation inductions on pain

See **Figure 4** for the effect sizes per study and the pooled effects. Meta-analysis indicated a medium overall effect of the expectation inductions on pain in clinical samples ($k = 27$, $g = 0.61$, 95% CI 0.42-0.79). A high degree of heterogeneity was observed ($I^2 = 73\%$), with the study effect sizes ranging between $g = -0.58$ and $g = 1.85$. The effect sizes for the different expectation inductions varied from a medium to large pooled effect of verbal suggestion ($k = 18$, $g = 0.75$, 95% CI 0.50-1.00, $I^2 = 78\%$), to a medium pooled effect of conditioning (always paired with verbal suggestion; $k = 3$, $g = 0.65$, 95% CI 0.18-1.11, $I^2 = 56\%$), and to a small pooled effect of imagery ($k = 6$, $g = 0.27$, 95% CI 0.02-0.53, $I^2 = 42\%$).

The overall effect of the expectation inductions corresponded with an average pain reduction of 1.16 points on a scale of 0 to 10 (95% CI 0.77-1.54). Verbal suggestion reduced pain with 1.39 points (95% CI 0.85-1.93), conditioning with 1.03 points (95% CI 0.30-1.76), and imagery with 0.62 points (95% CI 0.10-1.15).

The results of the studies for which sufficient data for meta-analyses were not available were in line with the observed pooled effects. Ho et al.³⁴ found a mean difference of 18.3 on a scale of 0 to 100 between a verbal suggestion condition and a no-treatment condition. Liberman⁵² observed that patients reported significantly less labor, postpartum, and experimental pain in a verbal suggestion condition compared with a control condition ($P < 0.001$). Laska and Sunshine⁴⁹ found that participants reported less pain when a placebo followed an active analgesic

(ie, conditioning) rather than when it followed a placebo (ie, no conditioning; difference between 0.5 and 3.6 on a sum of pain intensity differences scale).

3.5. Subset analyses

3.5.1. Effects on different types of pain

A comparison of the effects of the expectation inductions on different types of pain (see **Table 1** for specifications) indicated a medium to large pooled effect on experimental pain ($k = 9$, $g = 0.72$, 95% CI 0.43-1.01, $I^2 = 52\%$) and a medium pooled effect on clinical pain ($k = 18$, $g = 0.55$, 95% CI 0.33-0.78, $I^2 = 77\%$). A further comparison of acute procedural vs chronic clinical pain indicated a medium pooled effect on acute procedural pain ($k = 12$, $g = 0.67$, 95% CI 0.36-0.97, $I^2 = 74\%$) compared with a small pooled effect on chronic pain ($k = 6$, $g = 0.33$, 95% CI 0.04-0.62, $I^2 = 70\%$). Comparing the effects on the different types of pain for the separate expectation inductions was possible only for verbal suggestion. The effects of verbal suggestion on experimental pain were comparable to the overall effect ($k = 6$, $g = 0.79$, 95% CI 0.37-1.21, $I^2 = 59\%$), but the difference between the effects on acute procedural and chronic pain was considerably larger ($k = 7$, $g = 1.03$, 95% CI 0.79-1.27, $I^2 = 24\%$ vs $k = 5$, $g = 0.25$, 95% CI -0.06 to 0.56, $I^2 = 66\%$, respectively).

3.5.2. Post hoc: route of treatment administration

Verbal suggestions or conditioning referring to treatments that were administered through injection (see **Table 1** for relevant studies) were associated with large pooled effects ($k = 8$, $g = 0.90$, 95% CI 0.58-1.21, $I^2 = 52\%$), whereas oral and cutaneous treatments were associated with a small to medium pooled effect ($k = 3$, $g = 0.42$, 95% CI -0.23 to 1.07, $I^2 = 91\%$ and $k = 4$, $g = 0.47$, 95% CI 0.00-0.94, $I^2 = 70\%$, respectively). When analyzing only the effects of verbal suggestion, comparable results were found ($k = 7$, $g = 0.87$, 95% CI 0.51-1.23, $I^2 = 56\%$ vs $k = 3$, $g = 0.42$, 95% CI -0.23 to 1.07, $I^2 = 91\%$ vs $k = 3$, $g = 0.56$, 95% CI 0.01-1.11, $I^2 = 77\%$, respectively).

3.5.3. Post hoc: active or placebo treatment

Studies that assessed the effects of verbal suggestion or conditioning that referred to an active treatment (see **Table 1** for relevant studies) found a medium to large pooled effect ($k = 5$, $g = 0.73$, 95% CI 0.35-1.10, $I^2 = 70\%$), compared with a large

	Random sequence generation (selection bias)	Allocation concealment (selection bias)	Incomplete outcome data (attrition bias)	Selective reporting (reporting bias)	Other bias
Amanzio et al. (2001)	?	?	+	+	+
Benedetti et al. (1995)	?	?	?	?	?
Benedetti et al. (2003)	?	?	+	+	?
Benedetti et al. (2006)	?	?	+	+	+
Bialosky et al. (2014)	+	+	?	+	+
Charron et al. (2006)	?	?	?	+	?
Danhauer et al. (2007)	+	?	?	?	+
de Craen et al. (2001)	?	+	?	+	+
Foji et al. (2015)	+	?	+	+	-
Gonzales et al. (2010)	+	?	+	+	+
Gryll & Katahn (1978)	?	?	?	?	?
Hashish et al. (1988)	+	?	?	+	?
Hashmi et al. (2014)	?	?	?	+	?
Ho et al. (1988)	?	?	?	?	?
Jacobson (2006)	+	?	?	?	?
Kam-Hansen et al. (2014)	+	+	-	+	+
Klinger et al. (2007)	?	?	+	+	?
Kwekkeboom et al. (2008)	-	-	-	+	+
Laska & Sunshine (1973)	?	+	-	?	?
Lee et al. (2012)	?	?	+	+	?
Levine & Gordon (1984)	?	?	+	+	?
Liberman (1964)	-	-	+	?	-
Petersen et al. (2012)	+	?	?	-	?
Petersen et al. (2014)	-	-	+	+	?
Pollo et al. (2003)	?	?	?	+	+
Price et al. (2007)	-	-	+	-	?
Schmid et al. (2015)	?	?	+	+	?
Vase et al. (2003)	-	-	?	?	?
Vase et al. (2005)	-	-	?	?	?
Wells (1989)	?	?	?	+	?

pooled effect in studies that used a placebo treatment ($k = 13$, $g = 0.90$, 95% CI 0.61–1.19, $I^2 = 58\%$). When analyzing only the effects of verbal suggestion, comparable results were found ($k = 5$, $g = 0.73$, 95% CI 0.35–1.10, $I^2 = 70\%$ vs $k = 11$, $g = 0.95$, 95% CI 0.63–1.26, $I^2 = 58\%$, respectively). No differential effects were indicated in studies in which both active and placebo treatments were used ($g = 0.25$, 95% CI -0.13 to 0.64 , $I^2 = 64\%$ and $g = 0.22$, 95% CI -0.15 to 0.59 , $I^2 = 62\%$, respectively).^{18,31,39}

3.6. Effect of expectation inductions on additional outcomes

See Figure 5 for the effect sizes per study and the pooled effects for each of the additional outcomes.

3.6.1. Expected pain

From 5 (of 7) studies, sufficient data were available to analyze the effects of expectation inductions ($k = 5$ verbal suggestion) on self-reported expectations of pain. A medium pooled effect was observed ($g = 0.66$, 95% CI 0.43–0.90, $I^2 = 0\%$).

3.6.2. Affective pain

From 7 (of 10) studies, sufficient data were available to analyze the effects of expectation inductions ($k = 4$ verbal suggestion, $k = 3$ imagery) on affective pain (ie, pain unpleasantness or pain distress). A medium pooled effect was observed ($g = 0.45$, 95% CI 0.21–0.70, $I^2 = 34\%$).

3.6.3. Anxiety

From 5 (of 6) studies, sufficient data were available to analyze the effects of expectation inductions ($k = 2$ verbal suggestion, $k = 3$ imagery) on anxiety (measured with the state version of the State-Trait anxiety Inventory or an anxiety visual analogue scale). A large pooled effect was observed ($g = 1.38$, 95% CI 0.11–2.66, $I^2 = 96\%$); however, when excluding an extreme outlier ($g = 7.93^{22}$), no effect was observed ($g = 0.03$, 95% CI -0.21 to 0.26 , $I^2 = 0\%$).^{30,73,83}

3.7. Sensitivity analyses for overall effect of expectation inductions on pain

3.7.1. Risk of bias within studies

Excluding studies that were judged to have a high risk of bias on one or more items ($k = 9$, Fig. 3) did not substantially affect the overall effect size ($g = 0.63$, 95% CI 0.38–0.87).

3.7.2. Publication bias

The funnel plot (Fig. 6) suggests publication bias. The trim and fill method indicated that 6 studies demonstrating below-average effects of an expectation induction on pain relief (the black dots in the figure) were estimated to be missing. Including these studies would lower the overall effect size to $g = 0.43$ (95% CI 0.24–0.62).

Figure 3. Risk of bias summary: review authors' judgments about each risk of bias item for each included study. For the item "Selective outcome reporting" (reporting bias), the absence of a preregistered study protocol did not affect the judgment, because a protocol was absent for 93% of the studies.

Figure 4. Forest plot of the random-effects meta-analysis indicating the effects of the expectation inductions (verbal suggestion, conditioning, and imagery) on pain relief in clinical samples. Positive values for g indicate lower postintervention pain ratings in the intervention condition than in the control condition.

3.7.3. Type of control condition

When expectation inductions were compared with a control condition with a control treatment, a pooled effect of $g = 0.58$ ($k = 16$, 95% CI 0.34-0.82) was observed, whereas for studies in which a no-treatment control condition was used, a pooled effect of $g = 0.65$ ($k = 11$, 95% CI 0.35-0.94) was found. Excluding 3 studies that involved a control condition in which some expectations of pain relief might have been induced^{18,48,84} resulted in an overall effect of $g = 0.67$ (95% CI 0.49-0.86).

3.7.4. Between- vs within-subjects comparisons

The pooled effect for studies for which between-subjects comparisons were reported was $g = 0.53$ ($k = 16$, 95% CI 0.26-0.80), compared with $g = 0.70$ for studies in which within-subjects comparison were used ($k = 11$, 95% CI 0.45-0.96). Inclusion of within- rather than between-subjects comparisons of 2 studies for which both comparisons could be made did not affect the overall effect size ($g = 0.60$, 95% CI 0.43-0.78).

3.7.5. Imputed correlation coefficients

Sensitivity analyses testing whether the imputed intervention-control correlation of $r = 0.5$ for within-subjects comparisons

affected the observed effects indicated a stable overall effect size (when $r = 0.1$, $g = 0.60$, 95% CI 0.41-0.79; when $r = 0.9$, $g = 0.61$, 95% CI 0.44-0.77).

3.7.6. Postintervention vs change scores

When excluding three studies for which only change scores were available,^{8,13,51} rather than the preferred postintervention scores, the overall effect size was $g = 0.55$ (95% CI 0.37-0.73). When selecting change scores rather than postintervention scores (available for 12 studies), the overall effect size was $g = 0.70$ (95% CI 0.49-0.90).

In summary, these sensitivity analyses indicate a relatively stable overall effect size, ranging from $g = 0.43$ to $g = 0.70$.

4. Discussion

The current meta-analysis assessed the pain-reducing effects of 3 expectation interventions, ie, verbal suggestion, conditioning, and imagery, in clinical samples. Meta-analysis of 27 studies showed an overall medium-sized (heterogeneous) effect of the interventions on patients' pain relief. The effects of verbal suggestion were most frequently studied and could be qualified as medium to large. Conditioning (always paired with verbal

Figure 5. Forest plot of the random-effects meta-analysis indicating the effects of the expectation inductions (verbal suggestion, conditioning, and imagery) on expected pain, affective pain, and anxiety in clinical samples. Positive values for *g* indicate lower postintervention values in the intervention condition than in the control condition.

suggestion) and imagery were studied much less frequently, and were associated with medium and small effects, respectively. The effect sizes varied depending on the type of pain that patients experienced, with medium to large effects in the case of experimental and acute procedural pain, but small effects on chronic pain. Thus, interventions that can induce analgesic

expectations, particularly verbal suggestions for acute procedural pain, were found to relieve patients' pain and can thus possibly be used to optimize the effectiveness of standard analgesic treatments in clinical practice.

The findings of this meta-analysis extend previous meta-analyses in which the pain-reducing effects of verbal suggestion

Figure 6. Funnel plot of SE by Hedges *g*.

and conditioning were studied in the context of placebo effects^{80,81} and a meta-analysis and systematic reviews in which the pain-reducing effects of imagery were studied,^{66,67,79} by directly comparing the effects of these expectation inductions, while focusing on brief interventions in clinical samples. The observed medium to large effects of verbal suggestion on experimentally evoked and acute procedural pain were generally in line with the findings of a previous meta-analysis⁸¹ and more recent studies in healthy participants,^{5,53,76} which supports the transferability of findings from healthy to clinical samples. In contrast, the effects of verbal suggestion on chronic pain were found to be small, possibly because of repeated negative treatment experiences in the past and consequently more negative expectations regarding pain treatment in general that cannot be easily molded by a brief verbal suggestion.²³ However, because within-study comparisons of experimental and chronic pain provided somewhat equivocal results,^{13,63,64} and given the heterogeneity of the studies, further research is required. Surprisingly, although conditioning procedures were always paired with corresponding verbal suggestions, their effects on pain in clinical samples were not larger than the effects of verbal suggestion alone. This finding is in contrast with previous research in healthy samples, where such procedures are generally observed to have more robust effects on pain than verbal suggestions alone.^{6,47,54} However, because the effects of conditioning in clinical samples could be analyzed only in 3 studies and were studied only on experimental pain, and because conditioning procedures were always paired with verbal suggestion, no firm conclusions can be drawn yet about the size of conditioning effects in clinical samples. Imagery was found to have a small effect on clinical pain in our meta-analysis. This is partially in contrast with previous reviews that indicated small to large effects of imagery on pain.^{66,67,79} Also, *a priori*, we considered that imagery might be more effective than verbal suggestion because visual thinking has been found to have a larger impact on emotions, and hence possibly also the subjective pain experience, than verbal thinking^{28,35} and because imagery entails more active involvement.²⁰ Several factors might explain these findings. First, the selected imagery interventions were brief, maximally 1 day (to increase comparability between the expectation inductions). Possibly more practice time is required to obtain substantial effects.⁷⁹ Second, imagery instructions were always delivered through audio recordings, whereas verbal suggestions were given by the experimenter. Personal communication might enhance the effects of expectation inductions.

Subsequent post hoc analyses demonstrated that the observed effects of verbal suggestion and conditioning varied depending on the route of administration of the medical treatment to which they referred, with larger effects for more invasive treatments (injections) than less invasive treatments (oral and cutaneous). This is in line with previous experimental placebo research and a meta-analysis of placebo arm data of clinical trials.^{19,40} In addition, the effects of verbal suggestion and conditioning were slightly larger when they referred to a placebo rather than an active treatment. However, direct comparisons within three studies indicated no differential effects.^{18,31,39} Also, research in healthy samples provides equivocal results regarding the relative effect sizes.^{4,72} Nonetheless, these findings underscore that expectation interventions are not only relevant in the context of placebo effects, but also that they can enhance the analgesic effects of active treatments in clinical samples.

The core working mechanism of verbal suggestion, conditioning, and imagery is believed to be expectancy, as already implied by the term “expectation inductions.” Our meta-analysis of the

subset of studies in which expectations were measured demonstrated that verbal suggestion indeed induced expectations of pain relief, and the study authors showed that these expectations predicted effects on actually experienced pain.^{13,63,73,82,83} Previous research in healthy samples confirmed that also conditioning and imagery induce expectations,^{12,29,45,46,56,61} but, because of a lack of research, this cannot yet be confirmed in clinical samples. Also, anxiety reduction has been considered as a possible psychological working mechanism.^{21,53,66,79} However, our meta-analysis could not demonstrate an effect of the expectation inductions on anxiety in clinical samples, with the exception of 1 study in which large effects of imagery on anxiety were observed. Preliminary evidence from another study⁸³ suggests possible effects on pain specific anxiety. Several other psychological processes (eg, general affect, attention, or sense of control) might be affected by the interventions, but this could not be assessed in the meta-analysis because necessary data were not available. We could not meta-analyze physiological and neuroimaging data because of the paucity and complexity of the data. Although several previous reviews illustrate the neurobiological mechanisms of placebo effects and imagery, it was predominantly in healthy samples.^{3,55,71} An inspection of the included studies in patient samples provides preliminary evidence that verbal suggestion might be able to reduce heart rate^{9,65} and c-reactive protein,³⁰ but not cortisol (possibly because of methodological issues).^{30,36,73} A study on imagery found no evidence for effects on physiological responses.²² At a neurobiological level, the effects of verbal suggestion and conditioning on pain have been found to be associated with pain-related brain activity and connectivity among different brain regions.^{9,31,50,68,73} Further research is required to allow more conclusive inferences of the effects of expectation interventions on physiological and neurological processes in clinical populations.^{9,30,65,73} (see further Refs. 3,71).

When evaluating the current results, certain methodological factors that could have affected the observed effect sizes should be considered. Despite considerable heterogeneity, sensitivity analyses indicated a relatively stable overall effect size in relation to the research design (type of control condition, within- vs between-subjects comparisons) and selected values for analyses (imputed correlation coefficients, postintervention rather than change scores). However, there were indications for publication bias, which might have inflated the overall effect size (although the adjusted effect size could still be qualified as medium). Bias in the individual studies could frequently not be judged decisively because of insufficiently detailed reporting and the absence of preregistered study protocols. Also, response bias due to the (partial) infeasibility of blinding cannot be excluded. Nevertheless, because excluding studies with a known high risk of bias barely influenced the observed overall effect size, the influence of study bias seems minor. Last, the observed pairing of conditioning with verbal suggestion and the frequent inherent inclusion of relaxation, and possibly also verbal suggestion, in imagery interventions, could have affected the observed effects and hampered judgments of the effectiveness of the separate intervention components.

Based on this meta-analysis, several directions for future research can be considered. Most importantly, given the current positive but heterogeneous and still limited findings, future research might focus particularly on further examining the elements that determine the effectiveness of the different expectation inductions and on maximizing therapeutic effects. Research on active intervention elements (eg, specifics of verbal suggestion and pure imagery), mediating factors (expectations, physiological and neurobiological responses, and, eg, anxiety and attention), moderating factors (eg, previous pain experiences, pain treatment

history, desire for pain relief, and personality characteristics), and outcome characteristics (eg, type of pain) could provide insight into what determines the effects of the expectation interventions, and for whom and when they are effective. Also, combining different expectation inductions might enhance the effects, and for patients with chronic pain, more extensive interventions (eg, also addressing general expectations regarding medical treatment and health) might be considered. Importantly, research should not only aim at inducing and/or enhancing positive expectations but should also address negative expectations regarding adverse effects.⁶² Furthermore, the current findings allow for conclusions regarding only the short-term effects of the expectation interventions; further research is warranted to determine whether the interventions have a long-lasting clinical impact. Last, more detailed methodological reporting of the research, including preregistration, would further advance the field and facilitate future meta-analyses.^{32,75}

In conclusion, the current meta-analysis indicated that brief expectation interventions, especially verbal suggestion, can relieve patients' acute procedural and, to a lesser extent, chronic pain. Most notably, the observed analgesic effects of verbal suggestions regarding placebo or active treatments underline the importance of the information a clinician provides when administering an analgesic treatment. Informing patients about, and emphasizing, the positive intended and expected outcomes of an analgesic intervention, without neglecting possible negative side effects, can optimize treatment effectiveness.

Conflict of interest statement

The authors have no conflicts of interest to declare.

This research is supported by an Innovation Scheme (Vidi) Grant of the Netherlands Organization for Scientific Research (granted to A.W.M.E., <http://www.nwo.nl/en>). The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript. The authors thank Alice Tillema of the Library of Medical Sciences at Radboud University for her help in developing the search strategy.

Appendix A. Supplemental Digital Content

Supplemental Digital Content associated with this article can be found online at <http://links.lww.com/PAIN/A249>.

Article history:

Received 16 November 2015

Received in revised form 27 January 2016

Accepted 22 February 2016

Available online 1 March 2016

References

- [1] Amanzio M, Benedetti F. Neuropharmacological dissection of placebo analgesia: expectation-activated opioid systems versus conditioning-activated specific subsystems. *J Neurosci* 1999;19:484–94.
- [2] Amanzio M, Pollo A, Maggi G, Benedetti F. Response variability to analgesics: a role for non-specific activation of endogenous opioids. *PAIN* 2001;90:205–15.
- [3] Atlas LY, Wager TD. A meta-analysis of brain mechanisms of placebo analgesia: consistent findings and unanswered questions. In: *Handbook of experimental pharmacology*, Vol. 225. Berlin: Springer, 2014. p. 37–69.
- [4] Atlas LY, Whittington RA, Lindquist MA, Wielgosz J, Sonty N, Wager TD. Dissociable influences of opiates and expectations on pain. *J Neurosci* 2012;32:8053–64.
- [5] Atlas LY, Wielgosz J, Whittington RA, Wager TD. Specifying the non-specific factors underlying opioid analgesia: expectancy, attention, and affect. *Psychopharmacology* 2014;231:813–23.
- [6] Bartels DJP, van Laarhoven AIM, Haverkamp EA, Wilder-Smith OH, Donders ART, van Middendorp H, van de Kerkhof PCM, Evers AWM. Role of conditioning and verbal suggestion in placebo and nocebo effects on itch. *PLoS One* 2014;9:e91727.
- [7] Benedetti F. *Placebo effects. Understanding the mechanisms in health and disease*. Oxford: Oxford University Press, 2008.
- [8] Benedetti F, Amanzio M, Maggi G. Potentiation of placebo analgesia by proglumide. *Lancet* 1995;346:1231.
- [9] Benedetti F, Arduino C, Costa S, Vighetti S, Tarenzi L, Rainero I, Asteggiano G. Loss of expectation-related mechanisms in alzheimer's disease makes analgesic therapies less effective. *PAIN* 2006;121:133–44.
- [10] Benedetti F, Maggi G, Lopiano L, Lanotte M, Rainero I, Vighetti S, Pollo A. Open versus hidden medical treatments: the patient's knowledge about a therapy affects the therapy outcome. *Prev Treat* 2003;6:1a.
- [11] Bialosky JE, George SZ, Horn ME, Price DD, Staud R, Robinson ME. Spinal manipulative therapy-specific changes in pain sensitivity in individuals with low back pain (nct01168999). *J Pain* 2014;15:136–48.
- [12] Boselie JJLM, Vancleef LMG, Smeets T, Peters ML. Increasing optimism abolishes pain-induced impairments in executive task performance. *PAIN* 2014;155:334–40.
- [13] Charron J, Rainville P, Marchand S. Direct comparison of placebo effects on clinical and experimental pain. *Clin J Pain* 2006;22:204–11.
- [14] Cohen J. *Statistical power analysis for the behavioral sciences*, Vol. 2. Hillsdale: Lawrence Erlbaum Associates, 1988.
- [15] Colloca L, Petrovic P, Wager TD, Ingvar M, Benedetti F. How the number of learning trials affects placebo and nocebo responses. *PAIN* 2010;151:430–9.
- [16] Cormier S, Lavigne GL, Choiniere M, Rainville P. Expectations predict chronic pain treatment outcomes. *PAIN* 2015;157:329–38.
- [17] Danhauer SC, Marler B, Rutherford CA, Lovato JF, Asbury DY, McQuellon RP, Miller BE. Music or guided imagery for women undergoing colonoscopy: a randomized controlled study of effects on anxiety, perceived pain, and patient satisfaction. *J Low Genit Tract Dis* 2007;11:39–45.
- [18] de Craen AJM, Lampe-Schoenmaeckers AJEM, Kraal JW, Tjissen JGP, Kleijnen J. Impact of experimentally-induced expectancy on the analgesic efficacy of tramadol in chronic pain patients: a 2 × 2 factorial, randomized, placebo-controlled, double-blind trial. *J Pain Symptom Manage* 2001;21:210–17.
- [19] de Craen AJM, Tjissen JGP, de Gans J, Kleijnen J. Placebo effect in the acute treatment of migraine: subcutaneous placebos are better than oral placebos. *J Neurol* 2000;247:183–8.
- [20] Fazio RH, Chen J, McDonel EC, Sherman SJ. Attitude accessibility, attitude behavior consistency, and the strength of the object evaluation association. *J Exp Soc Psychol* 1982;18:339–57.
- [21] Flaten MA, Aslaksen PM, Lyby PS, Bjorkedal E. The relation of emotions to placebo responses. *Philos Trans R Soc* 2011;366:1818–27.
- [22] Foji S, Tadayonfar MA, Mohsenpour M, Rakhshani MH. The study of the effect of guided imagery on pain, anxiety and some other hemodynamic factors in patients undergoing coronary angiography. *Complement Ther Clin Pract* 2015;21:119–23.
- [23] Geers AL, Fowler SL, Wellman JA, Helfer SG, Close S, France CR. Prior experience with a pain stimulus as a predictor of placebo analgesia. *J Behav Med* 2015;38:136–42.
- [24] Geers AL, Weiland PE, Kosbab K, Landry SJ, Helfer SG. Goal activation, expectations, and the placebo effect. *J Pers Soc Psychol* 2005;89:143–59.
- [25] Gonzales EA, Ledesma RJA, McAllister DJ, Perry SM, Dyer CA, Maye JP. Effects of guided imagery on postoperative outcomes in patients undergoing same-day surgical procedures: a randomized, single-blind study. *AANA J* 2010;78:181–8.
- [26] Gryll SL, Katahn M. Situational factors contributing to the placebo effect. *Psychopharmacology (Berl)* 1978;57:253–61.
- [27] Haanstra TM, Kamper SJ, Williams CM, Spiensma AS, Lin CWC, Maher CG, de Vet HCW, Ostelo RWJG. Does adherence to treatment mediate the relationship between patients' treatment outcome expectancies and the outcomes of pain intensity and recovery from acute low back pain? *PAIN* 2015;156:1530–6.
- [28] Hackmann A, Bennett-Levy J, Holmes EA. *Oxford guide to imagery in cognitive therapy*. Oxford University Press: Oxford, United Kingdom, 2011.
- [29] Hanssen MM, Peters ML, Vlaeyen JWS, Meevissen YMC, Vancleef LMG. Optimism lowers pain: evidence of the causal status and underlying mechanisms. *PAIN* 2013;154:53–8.
- [30] Hashish I, Hai HK, Harvey W, Feinmann C, Harris M. Reduction of postoperative pain and swelling by ultrasound treatment: a placebo effect. *PAIN* 1988;33:303–11.
- [31] Hashmi JA, Kong J, Spaeth R, Khan S, Kaptchuk TJ, Gollub RL. Functional network architecture predicts psychologically mediated

- analgesia related to treatment in chronic knee pain patients. *J Neurosci* 2014;34:3924–36.
- [32] Higgins JPT, Green S. (editors). *Cochrane handbook for systematic reviews of interventions*. Version 5.1.0 [updated March 2011]. The Cochrane Collaboration, 2011. Available from www.cochrane-handbook.org.
 - [33] Higgins JPT, Thompson SG, Deeks JJ, Altman DG. Measuring inconsistency in meta-analyses. *Br Med J* 2003;327:557–60.
 - [34] Ho KH, Hashish I, Salmon P, Freeman R, Harvey W. Reduction of post-operative swelling by a placebo effect. *J Psychosom Res* 1988;32:197–205.
 - [35] Holmes EA, Arntz A, Smucker MR. Imagery rescripting in cognitive behaviour therapy: images, treatment techniques and outcomes. *J Behav Ther Exp Psychiatry* 2007;38:297–305.
 - [36] Horan JJ, Laying FC, Pursell CH. Preliminary study of effects of “in vivo” emotive imagery on dental discomfort. *Perceptual Motor Skills* 1976;42:105–6.
 - [37] Jacobson AF. Cognitive-behavioral interventions for IV insertion pain. *AORN J* 2006;84:1031–48.
 - [38] Jensen MP, Karoly P. Motivation and expectancy factors in symptom perception—a laboratory study of the placebo-effect. *Psychosom Med* 1991;53:144–52.
 - [39] Kam-Hansen S, Jakubowski M, Kelley JM, Kirsch I, Hoaglin DC, Kaptchuk TJ, Burstein R. Altered placebo and drug labeling changes the outcome of episodic migraine attacks. *Sci Transl Med* 2014;6:218a5.
 - [40] Kaptchuk TJ, Stason WB, Davis RB, Legedza ATR, Schnyer RN, Kerr CE, Stone DA, Nam BH, Kirsch I, Goldman RH. Sham device v inert pill: randomised controlled trial of two placebo treatments. *Br Med J* 2006;332:391–7.
 - [41] Kessner S, Wiech K, Forkmann K, Ploner M, Bingel U. The effect of treatment history on therapeutic outcome: an experimental approach. *JAMA Intern Med* 2013;173:1468–9.
 - [42] King LA. The health benefits of writing about life goals. *Pers Soc Psychol Bull* 2001;27:798–807.
 - [43] Kirsch I. Response expectancy as a determinant of experience and behavior. *Am Psychol* 1985;40:1189–202.
 - [44] Kirsch I. Response expectancy theory and application: a decennial review. *Appl Prev Psychol* 1997;6:69–79.
 - [45] Kirsch I, Kong J, Sadler P, Spaeth R, Cook A, Kaptchuk T, Gollub R. Expectancy and conditioning in placebo analgesia: separate or connected processes? *JAMA Intern Med* 2014;151:51–9.
 - [46] Kirsch I, Lynn SJ, Vigorito M, Miller RR. The role of cognition in classical and operant conditioning. *J Clin Psychol* 2004;60:369–92.
 - [47] Klinger R, Soost S, Flor H, Worm M. Classical conditioning and expectancy in placebo hypoalgesia: a randomized controlled study in patients with atopic dermatitis and persons with healthy skin. *PAIN* 2007;128:31–9.
 - [48] Kwekkeboom KL, Wanta B, Bumpus M. Individual difference variables and the effects of progressive muscle relaxation and analgesic imagery interventions on cancer pain. *J Pain Symptom Manage* 2008;36:604–15.
 - [49] Laska E, Sunshine A. Anticipation of analgesia. A placebo effect. *Headache* 1973;13:1–11.
 - [50] Lee HF, Hsieh JC, Lu CL, Yeh TC, Tu CH, Cheng CM, Niddam DM, Lin HC, Lee FY, Chang FY. Enhanced affect/cognition-related brain responses during visceral placebo analgesia in irritable bowel syndrome patients. *PAIN* 2012;153:1301–10.
 - [51] Levine JD, Gordon NC. Influence of the method of drug administration on analgesic response. *Nature* 1984;312:755–6.
 - [52] Liberman R. An experimental study of the placebo response under three different situations of pain. *J Psychiat Res* 1964;2:233–46.
 - [53] Lyby PS, Aslaksen PM, Flaten MA. Is fear of pain related to placebo analgesia? *J Psychosom Res* 2010;68:369–77.
 - [54] Martin-Pichora AL, Mankovsky-Arnold TD, Katz J. Implicit versus explicit associative learning and experimentally induced placebo hypoalgesia. *J Pain Res* 2011;4:67–77.
 - [55] McNorgan C. A meta-analytic review of multisensory imagery identifies the neural correlates of modality-specific and modality-general imagery. *Front Hum Neurosci* 2012;6:285.
 - [56] Meevissen YMC, Peters ML, Alberts HJEM. Become more optimistic by imagining a best possible self: effects of a two week intervention. *J Behav Ther Exp Psychiat* 2011;42:371–8.
 - [57] Moher D, Liberati A, Tetzlaff J, Altman DG; Prisma Group. Preferred reporting items for systematic reviews and meta-analyses: the Prisma statement. *PLoS Med* 2009;6:e1000097.
 - [58] Mondloch MV, Cole DC, Frank JW. Does how you do depend on how you think you'll do? A systematic review of the evidence for a relation between patients' recovery expectations and health outcomes. *Can Med Assoc J* 2001;165:174–9.
 - [59] Montgomery GH, Kirsch I. Classical conditioning and the placebo effect. *PAIN* 1997;72:107–13.
 - [60] Peerdeman KJ, van Laarhoven AIM, Donders ART, Hopman MTE, Peters ML, Evers AWM. Inducing expectations for health: effects of verbal suggestion and imagery on pain, itch, and fatigue as indicators of physical sensitivity. *PLoS One* 2015;10:e0139563.
 - [61] Peters ML, Flink IK, Boersma K, Linton SJ. Manipulating optimism: can imagining a best possible self be used to increase positive future expectancies? *J Posit Psychol* 2010;5:204–11.
 - [62] Petersen GL, Finnerup NB, Colloca L, Amanzio M, Price DD, Jensen TS, Vase L. The magnitude of nocebo effects in pain: a meta-analysis. *PAIN* 2014;155:1426–34.
 - [63] Petersen GL, Finnerup NB, Grosen K, Pilegaard HK, Tracey I, Benedetti F, Price DD, Jensen TS, Vase L. Expectations and positive emotional feelings accompany reductions in ongoing and evoked neuropathic pain following placebo interventions. *PAIN* 2014;155:2687–98.
 - [64] Petersen GL, Finnerup NB, Norskov KN, Grosen K, Pilegaard HK, Benedetti F, Price DD, Jensen TS, Vase L. Placebo manipulations reduce hyperalgesia in neuropathic pain. *PAIN* 2012;153:1292–300.
 - [65] Pollo A, Vighetti S, Rainero I, Benedetti F. Placebo analgesia and the heart. *PAIN* 2003;102:125–33.
 - [66] Posadzki P, Ernst E. Guided imagery for musculoskeletal pain: a systematic review. *Clin J Pain* 2011;27:648–53.
 - [67] Posadzki P, Lewandowski W, Terry R, Ernst E, Stearns A. Guided imagery for non-musculoskeletal pain: a systematic review of randomized clinical trials. *J Pain Symptom Manage* 2012;44:95–104.
 - [68] Price DD, Craggs J, Verne GN, Perlstein WM, Robinson ME. Placebo analgesia is accompanied by large reductions in pain-related brain activity in irritable bowel syndrome patients. *PAIN* 2007;127:63–72.
 - [69] Price DD, Finniss DG, Benedetti F. A comprehensive review of the placebo effect: recent advances and current thought. *Annu Rev Psychol* 2008;59:565–90.
 - [70] Price DD, Milling LS, Kirsch I, Duff A, Montgomery GH, Nicholls SS. An analysis of factors that contribute to the magnitude of placebo analgesia in an experimental paradigm. *PAIN* 1999;83:147–56.
 - [71] Schedlowski M, Enck P, Rief W, Bingel U. Neuro-bio-behavioral mechanisms of placebo and nocebo responses: Implications for clinical trials and clinical practice. *Pharmacol Rev* 2015;67:697–730.
 - [72] Schenk LA, Sprenger C, Geuter S, Buchel C. Expectation requires treatment to boost pain relief: an fMRI study. *PAIN* 2014;155:150–7.
 - [73] Schmid J, Langhorst J, Gass F, Theysohn N, Benson S, Engler H, Gizewski ER, Forsting M, Elsenbruch S. Placebo analgesia in patients with functional and organic abdominal pain: a fMRI study in IBS, UC and healthy volunteers. *Gut* 2015;64:418–27.
 - [74] Schmid J, Theysohn N, Gass F, Benson S, Gramsch C, Forsting M, Gizewski ER, Elsenbruch S. Neural mechanisms mediating positive and negative treatment expectations in visceral pain: a functional magnetic resonance imaging study on placebo and nocebo effects in healthy volunteers. *PAIN* 2013;154:2372–80.
 - [75] Schulz KF, Altman DG, Moher D, Group C. Consort 2010 statement: updated guidelines for reporting parallel group randomised trials. *Br Med J* 2010;340:c332.
 - [76] Scott DJ, Stohler CS, Egnatuk CM, Wang H, Koeppe RA, Zubieta JK. Placebo and nocebo effects are defined by opposite opioid and dopaminergic responses. *Arch Gen Psychiat* 2008;65:220–31.
 - [77] Sohl SJ, Schnur JB, Montgomery GH. A meta-analysis of the relationship between response expectancies and cancer treatment-related side effects. *J Pain Symptom Manage* 2009;38:775–84.
 - [78] Stewart-Williams S, Podd J. The placebo effect: dissolving the expectancy versus conditioning debate. *Psychol Bull* 2004;130:324–40.
 - [79] Van Kuiken D. A meta-analysis of the effect of guided imagery practice on outcomes. *J Holist Nurs* 2004;22:164–79.
 - [80] Vase L, Petersen GL, Riley JL III, Price DD. Factors contributing to large analgesic effects in placebo mechanism studies conducted between 2002 and 2007. *PAIN* 2009;145:36–44.
 - [81] Vase L, Riley JL, Price DD. A comparison of placebo effects in clinical analgesic trials versus studies of placebo analgesia. *PAIN* 2002;99:443–52.
 - [82] Vase L, Robinson ME, Verne GN, Price DD. The contributions of suggestion, desire, and expectation to placebo effects in irritable bowel syndrome patients. An empirical investigation. *PAIN* 2003;105:17–25.
 - [83] Vase L, Robinson ME, Verne GN, Price DD. Increased placebo analgesia over time in irritable bowel syndrome (IBS) patients is associated with desire and expectation but not endogenous opioid mechanisms. *PAIN* 2005;115:338–47.
 - [84] Wells N. Management of pain during abortion. *J Adv Nurs* 1989;14:56–62.