

Spine Journal

Twenty-five years with the biopsychosocial model of low back pain - is it time to celebrate? A report from the Twelfth International Forum for Primary Care Research on Low Back Pain --Manuscript Draft--

Manuscript Number:	SPINE 130543R1
Full Title:	Twenty-five years with the biopsychosocial model of low back pain - is it time to celebrate? A report from the Twelfth International Forum for Primary Care Research on Low Back Pain
Article Type:	Health Services Research
Keywords:	Biopsychosocial model, back pain, pain related disability, return to work, clinical research, clinical practice, international conference
Corresponding Author:	Tamar Pincus, Ph.D Royal Holloway University of London Egham, Surrey UNITED KINGDOM
Corresponding Author Secondary Information:	
Corresponding Author's Institution:	Royal Holloway University of London
Corresponding Author's Secondary Institution:	
First Author:	Tamar Pincus, Ph.D
First Author Secondary Information:	
Order of Authors:	Tamar Pincus, Ph.D Peter Michael Kent, PhD Gert Bronfort, Ph.D Patrick Loisel, Ph.D Glenn Pransky, MD Jan Hartvigsen, PhD
Order of Authors Secondary Information:	
Additional Information:	
Question	Response
Please select the level of evidence for this manuscript.	

To the editor-in-Chief
Spine

Dear Sir,

We thank the reviewers for their comments and have amended the manuscript accordingly. Specifically:

1. Reviewer 1 requested that we comment on the comprehensiveness of the evidence described in the manuscript, considering the fact that it was based on key note presentations, albeit by world leaders, rather than systematic reviews. Reviewer 2 similarly requests that we acknowledge the possibility of a selection bias in presenting research findings. We agree with both reviewers, and have now added the following in the introduction:
The evidence reviewed constitutes a synthesis of key-note presentations and discussions. Although citations are provided to illustrate the arguments, and where possible, we rely on evidence from systematic reviews, we recognise that possible bias and lack of comprehensiveness may be inherent in this review.
2. Reference 3 was incorrect, as spotted by reviewer 1. We apologise for this and have corrected it in the list and in the text.
3. The second sentence in the “biological” section has been amended. It now reads:
These include the use of diagnostic imaging to quantify the degree of disk degeneration, vertebral marrow (Modic) changes, endplate lesions, and vertebral joint degeneration.
4. Reviewer 2 commented: “The biopsychosocial model such as introduced by Waddell in a Spine paper in 1987 is currently the prevailing paradigm in low back pain research. It hasn't been fully adopted in clinical practice so far according to the study authors despite the fact that many researchers nowadays take psychological and social factors into account when studying low back pain. This might be a problem of implementation but on the other hand we as researchers have to admit that we know very little. The explained variances of regression models in the field of low back pain are still low and treatment effect sizes small to modest, also in treatments addressing psychosocial factors. We still know very little about low back pain despite the biopsychosocial model and I miss that point when reading this paper.”

We have now inserted the following in the concluding paragraph:

In taking stock of the current state of knowledge, it seems evident that vast gaps remain in our understanding about the aetiology, prognosis and effective interventions in back pain, despite the biopsychosocial model.

5. Reviewer 2 also requested that we amend the structure to the traditional structure of introduction, method, results and discussion. In this instance we disagree with this opinion. We believe that synthesis of key notes and discussions from the Forum cannot be captured in a formal methods section, nor does it lend itself to replication. We note that narrative reviews traditionally are structured under similar sub-headings to our preferred structure, and that similar syntheses from previous Forums published in *Spine* have been written in our preferred structure (e.g. Pransky et al., 2011).
6. Reviewer 2 requested that the conclusion be summarised as a list of bullet points. We note that this is already done under Key points.
7. We thank reviewer 3 for their positive comments and endorsement of the manuscript.

Twenty-five years with the biopsychosocial model of low back pain – is it time to celebrate? A report from the Twelfth International Forum for Primary Care Research on Low Back Pain

Tamar Pincus, PhD a

Peter Kent, PhD b

Gert Bronfort, PhD c,d

Patrick Loisel, PhD e,f

Glenn Pransky, MD MOccH g,h,i

Jan Hartvigsen, PhD j,d

a Department of Psychology, Royal Holloway, University of London, England, UK.

b Research Department, Spine Centre of Southern Denmark, Institute of Regional Health Services Research, Hospital Lillebaelt, University of Southern Denmark, Middelfart, Denmark

c Musculoskeletal Research Program, Northwestern Health Sciences University, Minnesota, USA

d Nordic Institute of Chiropractic and Clinical Biomechanics, Odense, Denmark

e Dalla Lana School of Public Health, University of Toronto, Toronto, Canada

f Canadian Memorial Chiropractic College, Toronto, Canada

g Liberty Mutual Research Institute for Safety, Hopkinton, Massachusetts, USA

h Department of Family Medicine and Community Health, University of Massachusetts Medical School, Worcester Massachusetts, USA

i Harvard School of Public Health, Boston, USA

j Institute of Sports Science and Clinical Biomechanics, University of Southern Denmark, Odense, Denmark

Address for correspondence:

Prof Tamar Pincus

Department of Psychology, Clinical, Health and Social Psychology, Royal Holloway, University of London, England.

Phone: +44 1784 443523

Contact Email: t.pincus@rhul.ac.uk

No benefits in any form have been, or will be, received from a commercial party related directly or indirectly to the subject of this manuscript.

Study Design. An integrated review of current knowledge about the biopsychosocial model of back pain for understanding aetiology, prognosis and interventions, as presented at the plenary sessions of the

XII International Forum on LBP Research in Primary Care (Denmark 17-19 October 2012).

Objectives. To evaluate the utility of the model in reference to rising rates of back pain related disability, by identifying a) the most promising avenues for future research in biological, psychological and social approaches, b) promising combinations of all three approaches and c) obstacles to effective implementation of biopsychosocial based research and clinical practice.

Summary of Background Data. The biopsychosocial model of back pain has become a dominant model in the conceptualisation of the aetiology and prognosis of back pain, and has led to the development and testing of many interventions. Despite this back pain remains a leading source of disability worldwide.

Method. The review is a synthesis based on the plenary sessions and discussions at the XII International Forum on LBP Research in Primary Care. The presentations included evidence-based reviews of the current state of knowledge in each of the three areas (biological, psychological and social), identification of obstacles to effective implementation and missed opportunities, and identification of the most promising paths for future research.

Results. While there is good evidence for the role of biological, psychological and social factors in the aetiology and prognosis of back pain, synthesis of the three in research and clinical practice has been suboptimal.

Conclusion. The utility of the biopsychosocial framework cannot be fully assessed until we truly adopt and apply it in research and clinical practice.

- It is 25 years since Gordon Waddell's seminal paper on the biopsychosocial model in back pain was published by SPINE.
- Back pain remains an alarming worldwide health problem and is now the leading cause of disability.
- This may be a consequence of the mostly restrictive way the biopsychosocial model in back pain has been understood and applied rather than a failure of the model itself.
- The utility of the biopsychosocial framework cannot be fully assessed until we truly adopt and apply it in research and clinical practice.

25 years after Gordon Waddell's seminal paper on the biopsychosocial model, back pain remains a worldwide health challenge. Whether this is a result of problems in the model or its understanding and application was explored at the International Forum for Primary Care Research on Low Back Pain.

1 Twenty-five years with the biopsychosocial model of low back pain – is it time to
2 celebrate? A report from the Twelfth International Forum for Primary Care
3 Research on Low Back Pain
4
5
6

7 Tamar Pincus, PhD ^a

8
9 Peter Kent, PhD ^b

10
11 Gert Bronfort, PhD ^{c,d}

12
13 Patrick Loisel, PhD ^{e,f}

14
15 Glenn Pransky, MD MOcCH ^{g,h,i}

16
17 Jan Hartvigsen, PhD ^{j,d}
18
19

20 ^a Department of Psychology, Royal Holloway, University of London, England, UK.

21
22 ^b Research Department, Spine Centre of Southern Denmark, Institute of Regional
23 Health Services Research, Hospital Lillebaelt, University of Southern Denmark,
24 Middelfart, Denmark
25

26
27 ^c Musculoskeletal Research Program, Northwestern Health Sciences University,
28 Minnesota, USA
29

30
31 ^d Nordic Institute of Chiropractic and Clinical Biomechanics, Odense, Denmark

32
33 ^e Dalla Lana School of Public Health, University of Toronto, Toronto, Canada

34
35 ^f Canadian Memorial Chiropractic College, Toronto, Canada

36
37 ^g Liberty Mutual Research Institute for Safety, Hopkinton, Massachusetts, USA

38
39 ^h Department of Family Medicine and Community Health, University of
40 Massachusetts Medical School, Worcester Massachusetts, USA
41

42
43 ⁱ Harvard School of Public Health, Boston, USA

44
45 ^j Institute of Sports Science and Clinical Biomechanics, University of Southern
46 Denmark, Odense, Denmark
47

48
49
50 Address for correspondence:

51 Prof Tamar Pincus

52
53 Department of Psychology, Clinical, Health and Social Psychology, Royal
54 Holloway, University of London, England.
55

56
57 Phone: +44 1784 443523

58
59 Contact Email: t.pincus@rhul.ac.uk
60
61
62
63
64
65

1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65

No benefits in any form have been, or will be, received from a commercial party related directly or indirectly to the subject of this manuscript.

Structured abstract

1
2 **Study Design.** An integrated review of current knowledge about the
3
4 biopsychosocial model of back pain for understanding aetiology, prognosis and
5
6 interventions, as presented at the plenary sessions of the
7
8 XII International Forum on LBP Research in Primary Care (Denmark 17-19
9
10 October 2012).

11 **Objectives.** To evaluate the utility of the model in reference to rising rates of
12
13 back pain related disability, by identifying a) the most promising avenues for
14
15 future research in biological, psychological and social approaches, b) promising
16
17 combinations of all three approaches and c) obstacles to effective
18
19 implementation of biopsychosocial based research and clinical practice.

20 **Summary of Background Data.** The biopsychosocial model of back pain has
21
22 become a dominant model in the conceptualisation of the aetiology and
23
24 prognosis of back pain, and has led to the development and testing of many
25
26 interventions. Despite this back pain remains a leading source of disability
27
28 worldwide.

29 **Method.** The review is a synthesis based on the plenary sessions and discussions
30
31 at the XII International Forum on LBP Research in Primary Care. The
32
33 presentations included evidence-based reviews of the current state of knowledge
34
35 in each of the three areas (biological, psychological and social), identification of
36
37 obstacles to effective implementation and missed opportunities, and
38
39 identification of the most promising paths for future research.

40 **Results.** While there is good evidence for the role of biological, psychological and
41
42 social factors in the aetiology and prognosis of back pain, synthesis of the three
43
44 in research and clinical practice has been suboptimal.

45 **Conclusion.** The utility of the biopsychosocial framework cannot be fully
46
47 assessed until we truly adopt and apply it in research and clinical practice.
48
49
50
51
52
53
54
55

Key words

56
57 Biopsychosocial model, back pain, pain related disability, return to work, clinical
58
59 research, clinical practice, international conference
60
61

1
2 **Mini abstract**

3
4 25 years after Gordon Waddell's seminal paper on the biopsychosocial model,
5
6 back pain remains a worldwide health challenge. Whether this is a result of
7
8 problems in the model or its understanding and application was explored at the
9
10 International Forum for Primary Care Research on Low Back Pain.

11
12
13 **Key points**

- 14
15 • It is 25 years since Gordon Waddell's seminal paper on the biopsychosocial
16
17 model in back pain was published by SPINE.
18
19 • Back pain remains an alarming worldwide health problem and is now the
20
21 leading cause of disability.
22
23 • This may be a consequence of the mostly restrictive way the biopsychosocial
24
25 model in back pain has been understood and applied rather than a failure of
26
27 the model itself.
28
29 • The utility of the biopsychosocial framework cannot be fully assessed until
30
31 we truly adopt and apply it in research and clinical practice.
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65

Background

The state of the art

Gordon Waddell's seminal paper on the biopsychosocial model in back pain published by SPINE¹ marked a fundamental change in the conceptualization of back pain. The model suggests that back pain should be more broadly understood than is possible from a biomedical perspective alone, because for many individuals the main problem lies not with the common and frequently transient experience of pain, but rather in their own and society's perceptions and reactions to pain. Inappropriate reactions may include unnecessary avoidance of physical activity and social interactions, absenteeism from work, and high health care utilization.

The 25 year anniversary of Waddell's publication was a focus of the Forum for Research in Back Pain in Primary Care XII that was held in Odense, Denmark October 17-19 2012. The goal of the Forum is to share the latest concepts, methods, and results of research on low back pain diagnosis, evaluation, treatment, and disability prevention. The presentations described here addressed the three dimensions of the biopsychosocial model, how it has been applied, and promising areas for research to further develop this conceptual view of LBP. The evidence reviewed constitutes a synthesis of key-note presentations and discussions. Although citations are provided to illustrate the arguments, and where possible, we rely on evidence from systematic reviews, we recognize that possible bias and lack of comprehensiveness may be inherent in this review.

Back pain remains an alarming worldwide health problem and is now the leading cause of disability, with an estimated 632 million people affected.² When considering both death and disability, musculoskeletal conditions have the fourth greatest impact on the health of the world population and back pain accounts for nearly half of this. Disability due to musculoskeletal disorders is estimated to have increased by 45% from 1990 to 2010, and, with increasingly obese, sedentary and aging societies, is expected to increase even more in the

1 years to come.³ Against this backdrop, one can hardly say that the introduction of
2 the biopsychosocial model in research and practice has been a public health
3 success. In fact, alongside with the increasing rates of disability, and against
4 guideline advice, are increases in tests ⁴, and in the provision of biologic mono-
5 therapies that are costly and mostly ineffective.⁵ The question therefore is
6 whether it is the model itself that has failed to deliver or whether it is the
7 scientific and healthcare communities that have failed to adopt the model.
8
9
10
11
12
13

14 **Discussion**

15 *Explaining the current status*

16 Understanding the underlying principles of a condition is a prerequisite for
17 designing effective interventions, and while we are still struggling to identify the
18 precise biological basis for most back problems, there is good evidence to
19 suggest that psychological constructs such as pre-existing somatization,
20 depression, anxiety, fear avoidance beliefs, poor coping strategies and poor self-
21 efficacy are significant predictors of outcomes such as more severe pain, greater
22 functional disability and work loss. Similar constructs play a role in the transition
23 from acute to persistent pain and disability.⁶⁻⁸ Nevertheless results from trials
24 testing interventions aimed at changing psychological factors have been
25 disappointing⁹ and findings from systematic reviews of psychological
26 interventions for chronic pain groups show that effects are at best modest.¹⁰
27
28
29
30
31
32
33
34
35
36
37
38
39
40

41 Evidence also suggests that social and organizational factors influence the
42 consequences of back pain such as work absenteeism, but only a few trials have
43 evaluated the effect of social interventions.¹¹ Furthermore, regardless of whether
44 interventions are based on biological, psychological or social approaches, results
45 consistently show only small to moderate effects.¹²
46
47
48
49
50
51

52 One explanation may be that interventions in trials have rarely integrated all
53 three components of the biopsychosocial model. In addition, some interventions
54 that have attempted to integrate psychological methods into general practice and
55 physiotherapy care have been compromised by delivery at suboptimal levels of
56 dosage, content, fidelity and mode of delivery.⁹
57
58
59
60
61

1
2 Progress has also been compromised by lack of clarity about the selection of
3 appropriate outcomes. Thus, the experience of back pain per se and the
4 consequential disability and loss of social participation, such as work absence are
5 often confused in studies. These domains of health status are only weakly
6 associated and one should not be considered to be a proxy for the others.¹³ For
7 example, a certain level of back pain intensity may occur in one patient with
8 significant pain-related disability and work absence, while another patient with
9 an equal level of pain may continue to have an active life without loss of work
10 participation. Thus in a study that used work participation as the outcome, the
11 first patient would be classified as having a poor outcome and the second a good
12 outcome, whereas in a study focusing on pain intensity, both would be classified
13 as having a poor outcome. Such examples highlight the need for multi-domain
14 assessment and interpretation in clinical studies.^{1,14-16}

15
16
17
18
19
20
21
22
23
24
25
26
27
28 Finally within clinical practice, there is mostly little reward or opportunity for
29 primary care practitioners to utilize a comprehensive biopsychosocial approach
30 given current practice and payment structures. This may explain why
31 practitioners appear reluctant to attempt to influence the social aspects of the
32 pain experience, especially those related to work.¹⁷ Even in the occupational
33 health context of the USA, where there is sufficient payment and other incentives
34 based on outcome evaluations, providers retreat to the 'safe' biological arena
35 when faced with psychosocial problems.¹⁸ Finally, training for most of the
36 professions that treat back pain remains bio-medically focused and grounded in
37 profession-specific tradition rather than on contemporary evidence.¹⁹

38
39
40
41
42
43
44
45
46
47
48
49 *The biopsychosocial model: New and promising findings from the three*
50 *components*

51 Biological

52
53
54 The absence of established biomarkers of back pain has led to calls for increased
55 efforts to understand the biological components of back pain.²⁰⁻²² These include
56 the use of diagnostic imaging to quantify the degree of disk degeneration,
57 vertebral marrow (Modic) changes, endplate lesions, and vertebral joint

1 degeneration.^{23,24} These findings have shown positive associations with the
2 presence and severity of back pain symptoms on a population level, but currently
3 they are not a useful way of diagnostically classifying individual patients, nor of
4 informing treatment choice.^{25,26}
5
6
7

8
9 Spinal intersegmental motion assessment technology (e.g. quantitative video
10 fluoroscopy,²⁷ kinematic MRI,²⁸ and tissue elastography²⁹ has now reached a
11 level of sophistication that its application in research is likely to provide a
12 greater understanding of the association between spinal biomechanical
13 dysfunction and back pain. Using previous technology, it is possible to
14 distinguish low back pain patients from healthy controls by comprehensive
15 biomechanical analysis of trunk motion associated with standardized functional
16 tasks.³⁰ However, there is considerable variability both within and between the
17 populations with and without pain on these tasks and we have no knowledge
18 about the role of spinal functional performance as a treatment effect modifier or
19 prognostic factor. Currently, there is no evidence for a causal path between such
20 manifestations, disability and pain.
21
22
23
24
25
26
27
28
29
30
31

32
33 Central nervous system sensitization and abnormal central processing of pain is
34 emerging as an important biologic explanation for the persistence of pain.³¹⁻³⁵
35 There is even evidence that persistent back pain may alter brain morphology by
36 reducing the volume of grey matter in the prefrontal area and the thalamus³³ and
37 that such changes may be reversible once the pain is effectively treated.³⁶ Such
38 mechanisms may explain the small to moderate effects of numerous evidence-
39 based treatments, despite their being assumed to have very different
40 mechanisms of action.^{12,37} Early evidence suggests that it may be feasible to
41 normalize pain processing through real time functional MRI feedback training.³⁸
42
43
44
45
46
47
48
49
50

51 Another potentially important biological mechanism is epigenetics, which
52 through interactions with environmental factors, controls the expression of
53 genetic predispositions. Genetic factors have been shown to strongly influence
54 various spinal pain phenotypes³⁹ and epigenetic modulation has been shown to
55 be involved in the transition from acute to chronic pain,⁴⁰ in addition to the
56
57
58
59
60
61

1 degree of spinal disc degeneration.⁴¹ Thus the ability to influence epigenetic
2 expression in the future may lead to improvements in back pain treatment.⁴²
3
4

5 Psychological 6

7 Challenges being addressed by research into psychological aspects of back pain
8 can be divided into two broad goals: a) To better understand which
9 psychological risk factors impact on which outcomes and b) To elucidate
10 mechanisms related both to psychological dysfunction and to recovery.
11
12
13
14

15
16 In relation to both goals, an international consensus panel⁴³ recognized the need
17 to standardize the predictors included in prospective cohort research
18 investigating the transition from acute to persistent back pain. Identifying the
19 unique contribution of factors within specific subgroups will require extremely
20 large samples. In addition, the consortium recognized the potential impact of
21 social arrangement, health structures and local cultural beliefs, which have been
22 largely ignored in most previous research. Pooling of samples from international
23 regions is now possible, and provides a promising avenue to address limitations
24 in current knowledge.⁴⁴
25
26
27
28
29
30
31
32
33

34
35 In addition, recent emerging evidence about practitioners beliefs⁴⁵, behaviours⁴⁶,
36 and perceptions of their role, especially in reference to patient's work,⁴⁷ present
37 both a potential and a challenge for future research, because it implies that
38 practitioners may inadvertently play a role in maintaining patients' disability.
39
40
41
42
43

44 Finally, a promising direction is the inclusion of new psychological approaches
45 that aim to increase acceptance of inevitable pain states and increase
46 engagement with all aspects of life through changes in psychological flexibility,
47 perceived values and mindfulness informed therapy.^{48,49}
48
49
50
51
52

53 Social 54

55 Social factors including potential obstacles to recovery, in the form of legislation,
56 compensation systems and social and economic conventions and infrastructures
57 are perhaps the most neglected area of research in back pain. Furthermore,
58
59
60
61

1 when studied as outcomes, social factors have been typically measured as
2 secondary outcomes, and in many cases studies have been insufficiently
3 powered to draw reliable conclusions from their findings.
4
5
6

7 Measurement of social factors can be problematic, as they include factors
8 operating both at an individual and at a group level. Thus they include factors
9 relating to the individuals' status (such as employment), those relating to the
10 individuals' perception and reaction to their status (such as job satisfaction),
11 those relating to group level, including regional or national level (such as
12 incapacity legislation), and those relating to the process at group level (such as
13 the time and ease of obtaining incapacity benefit). While the former factors have
14 been studied, comparisons between systems necessitate large samples and
15 careful coding of complex systems to enable clarification of the role they might
16 play in maintaining disability. Not surprisingly, the impact of compensatory
17 systems on the rising rates of back pain-related disability remains unclear. Yet
18 this is one of the most promising areas for future research, and register-based
19 information collectable at the level of incapacity and welfare systems provide a
20 comprehensive picture of how social structures influence disability at the
21 societal level.
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36

37 There is emerging evidence that social factors contribute substantially to
38 disability beyond the factors operating at the level of individuals.^{11,50,51} For
39 instance, Anema et al⁵⁰ compared sustainable return to work rates between six
40 different countries and found that differences in applied work interventions, job
41 characteristics and social disability systems were more important than medical
42 interventions, patient and injury related factors in explaining the large between
43 country differences.⁵⁰ In addition, the findings indicated that longer delays
44 before assigning permanent incapacity benefits, and availability of financial
45 support for partial return to work were associated with more favorable
46 outcomes. Eliminating compensation for pain and suffering after a whiplash
47 injury in one Canadian province was associated with a decreased incidence of
48 those injuries as well as improved prognosis for patients.⁵² Research on workers
49 with chronic musculoskeletal pain showed that personal and work-related
50
51
52
53
54
55
56
57
58
59
60
61

1 factors were more important than pain as determinants of work ability and
2 staying at work.⁵³ Taken together, the evidence suggests that the less
3 engagement and investment patients have with disability compensation systems,
4 and the more they are supported in work resumption, the better their outcomes.
5
6

7
8
9 The positive impact of engaging the workplace in preventing work disability and
10 supporting return to work in LBP is a consistent finding.^{54,55} Key components
11 include early and supportive communication from the workplace, arrangements
12 to ensure a safe return to work within the physical capabilities of the worker,
13 and ongoing support from supervisors and co-workers. Some of these
14 interventions are most effective if primarily focused in the workplace, and thus
15 have the benefit of avoiding an overly medical /disease orientation in
16 management of a condition that does not benefit greatly from medical
17 interventions.^{56,57} In those with more chronic work disability, multi-faceted
18 interventions involving workers, employers, and health care providers, along
19 with a return to work coordinator, may be required to achieve positive
20 results.^{58,59} Many approaches found to be effective are not easily evaluated in a
21 RCT, and thus the evidence is sometimes interpreted as weak, despite
22 consistency of findings across studies, countries, and conditions.⁶⁰ The high cost
23 of work disability for workers, employers, and society has led to conclusions that
24 diffusion of these principles into general practice is a priority.⁶¹
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40

41 In addition, recent qualitative studies have indicated that employer perceptions
42 about when an employee should return to work after a period of sick leave
43 because of back pain may result in longer periods off work than necessary⁶²,
44 suggesting that there is scope to intervene also at the employer level. Of
45 importance, that study identified problems associated with processes within
46 workplaces, healthcare, vocational rehabilitation, and workers compensation,
47 which operate to extend absence from work in patients. Lack of communication
48 between the different systems is at the core of increased disability, an
49 observation reflected in Waddell's call⁶³ for all stake holders to get outside if
50 disability is to be meaningfully reduced.
51
52
53
54
55
56
57
58
59
60
61

Conclusions

A synthesis of new directions

The Forum concluded with a discussion on the opportunities for future research and applications of the biopsychosocial model. One new and promising direction is stratified care for back pain, where patients are screened for known biopsychosocial risk factors using reliable and valid tools, and then referred to interventions designed to target their specific problem and risk profile.⁶⁴ The challenge is to develop appropriately validated instruments that stratify patients into streams of care that optimize their chance of a good outcome. Such research is underway but needs further development, testing, and wider validation, especially with respect to measuring social determinants of work disability outcomes.⁶⁴⁻⁶⁶ This approach may also eventually allow us to target the particular needs of subgroups in the population, such as older people, for whom back pain can lead to social isolation and reduction in physical activity or younger people, for whom preventing long-term work-related disability may change their life trajectory.¹¹ Lifespan research is also needed to clarify the changing impact of psychological factors at different points in a person's life course, including childhood and adolescence.^{67,68} Forum participants stressed the importance of distinguishing between psychological and social domains in both research and clinical practice.

Another approach is influencing beliefs and behaviours at the population level where mass media campaigns may be useful if delivered efficiently.⁶⁹ Whether at the population or individual person level, meaningful reduction in the burden of back pain will require integrating strategies, for example: seeking input and active engagement from stakeholders such as employers to the design of interventions; increasing incentives for appropriate clinician responses to social factors; and shifting public perceptions of the role of active self-management.

Lastly, clarity about which predictors of outcome are prognostic factors and which are potential treatment effect modifiers⁷⁰ may help guide best practice

1 treatment and the prevention of disability. Some factors exert an influence on
2 outcome regardless of treatment while some only influence response to specific
3 treatments. Applying such information to identifiable subgroups of patients and
4 at the individual patient level will require focused research and methodology
5 development but may be well worth the effort. Interventions for some high-risk
6 groups may be complex and costly, but expensive care that is appropriately
7 targeted may still prove to be cost-effective.
8
9
10
11
12
13

14 In taking stock of the current state of knowledge, it seems evident that vast gaps
15 remain in our understanding about the aetiology, prognosis and effective
16 interventions in back pain, despite the biopsychosocial model. In our view, the
17 biopsychosocial model has not failed to explain back pain - what has failed is the
18 mostly restrictive way it has been understood and applied. Forum discussants
19 concluded that the utility of the biopsychosocial framework cannot be fully
20 assessed until we truly adopt and integrate it into research and clinical practice.
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65

References

1. Waddell G. A new clinical model for the treatment of low-back pain. *Spine* 1987;12:632-44.
2. Lim SS, Vos T, Flaxman AD, et al. A comparative risk assessment of burden of disease and injury attributable to 67 risk factors and risk factor clusters in 21 regions, 1990—2010: a systematic analysis for the Global Burden of Disease Study. *Lancet* 2012;380.
3. Murray CJ, Vos T, Lozano R, et al. Disability-adjusted life years (DALYs) for 291 diseases and injuries in 21 regions, 1990-2010: a systematic analysis for the Global Burden of Disease Study 2010. *The Lancet* 2012;380:2197-223.
4. Ivanova JI, Birnbaum HG, Schiller M, et al. Real-world practice patterns, health-care utilization, and costs in patients with low back pain: the long road to guideline-concordant care. *Spine J* 2011;11:622–32.
5. Deyo RA. Managing patients with back pain: putting money where our mouths are not. *Spine J* 2011;11:633-5.
6. Pincus T, Burton A, Vogel S, et al. A systematic review of psychological factors as predictors of chronicity/disability in prospective cohorts of low back pain. *Spine* 2002;27:E109-E20.
7. Gatchel RJ, Peng YB, Peters ML, et al. The biopsychosocial approach to chronic pain: scientific advances and future directions. *Psychol Bull* 2007;133:581-624.
8. Nicholas MK, Linton SJ, Watson PJ, et al. Early identification and management of psychological risk factors ("yellow flags") in patients with low back pain: a reappraisal. *Phys Ther* 2011;91:737-53.
9. van der Windt D, Hay EM, Jellema P, et al. Psychosocial interventions for low back pain in primary care: lessons learned from recent trials. *Spine* 2008;33:81-9.
10. Williams AC, Eccleston C, Morley S. Psychological therapies for the management of chronic pain (excluding headache) in adults. *Cochrane Database of Systematic Reviews* 2012;11.
11. Loisel P, Buchbinder R, Hazard R, et al. Prevention of work disability due to musculoskeletal disorders: the challenge of implementing evidence. *J Occup Rehabil* 2005;15:507-24.
12. Artus M, van der Windt DA, Jordan KP, et al. Low back pain symptoms show a similar pattern of improvement following a wide range of primary care treatments: a systematic review of randomized clinical trials. *Rheumatology (Oxford)* 2010;49:2346-56.
13. Mannion A, Junge A, Taimela S, et al. Active therapy for chronic low back pain - Part 3. Factors influencing self-rated disability and its change following therapy. *Spine* 2001;26:920-9.
14. Loisel P, Durand M-J, Berthelette D, et al. Disability prevention: The new paradigm of management of occupational back pain. *Dis Manage Health Outcomes* 2001;9:351-60.
15. Loisel P, Côté P. The Work Disability Paradigm and its Public Health Implications. In Anema PLaH ed. *Handbook of Work Disability: Prevention and Management*. New York: Springer Science+Business Media, 2013.
16. Sullivan MJ, Thibault P, Andrikonyte J, et al. Psychological influences on repetition-induced summation of activity-related pain in patients with chronic low back pain. *Pain* 2009;141:70-8.

17. Bishop P, Wing P. Compliance with clinical practice guidelines in family physicians managing workers compensation board patients with acute lower back pain. *Spine J* 2003;3:442-50.
18. Shaw WS, Pransky G, Winters T, et al. Does the presence of psychosocial 'yellow flags' alter patient-provider communication for work-related, acute low back pain? *J Occup Environ Med* 2009;51:1032-40.
19. Foster NE, Hartvigsen J, Croft PR. Taking responsibility for the early assessment and treatment of patients with musculoskeletal pain: a review and critical analysis. *Arthritis Res Ther* 2012;14.
20. Hancock MJ, Maher CG, Laslett M, et al. Discussion paper: what happened to the 'bio' in the bio-psycho-social model of low back pain? *Eur Spine J* 2011;20:2105-10.
21. Langevin H, M., Sherman KJ. Pathophysiological model for chronic low back pain integrating connective tissue and nervous system mechanisms. *Med Hypotheses* 2007;68:74-80.
22. Marras WS. The complex spine: the multidimensional system of causal pathways for low-back disorders. *Hum Factors* 2012;54:881-9.
23. Suri P, Dharamsi AS, Gaviola G, et al. Are facet joint bone marrow lesions and other facet joint features associated with low back pain? A pilot study. *PM R* 2013;5:194-200.
24. Jensen RK, Leboeuf-Yde C, Wedderkopp N, et al. Is the development of Modic changes associated with clinical symptoms? A 14-month cohort study with MRI. *Eur Spine J* 2012;21:2271-9.
25. Deyo R. Diagnostic evaluation of LBP: reaching a specific diagnosis is often impossible. *Arch Intern Med* 2002;162:1444-7.
26. Deyo RA, Mirza SK, Turner JA, et al. Overtreating chronic back pain: time to back off? *J Am Board Fam Med* 2009;22:62-8.
27. Breen AC, Teyhen DS, Mellor FE, et al. Measurement of intervertebral motion using quantitative fluoroscopy: report of an international forum and proposal for use in the assessment of degenerative disc disease in the lumbar spine. *Adv Orthop* 2012;802350.
28. Tan Y, Aghdasi BG, Montgomery SR, et al. Kinetic magnetic resonance imaging analysis of lumbar segmental mobility in patients without significant spondylosis. *European Spine Journal* 2012;21:2673-9.
29. Chan ST, Fung PK, Ng NY, et al. Dynamic changes of elasticity, cross-sectional area, and fat infiltration of multifidus at different postures in men with chronic low back pain. *Spine J* 2012;12:381-8.
30. Marras W, Ferguson S, Gupta P, et al. The Quantification of Low Back Disorder Using Motion Measures: Methodology and Validation. *Spine* 1999;24:2091.
31. Wand BM, O'Connell NE. Chronic non-specific low back pain - sub-groups or a single mechanism? *BMC Musculoskelet Disord* 2008;9.
32. Wand BM, Parkitny L, O'Connell NE, et al. Cortical changes in chronic low back pain: current state of the art and implications for clinical practice. *Man Ther* 2011;16:15-20.
33. Apkarian A, V., Sosa Y, Sonty S, et al. Chronic back pain is associated with decreased prefrontal and thalamic gray matter density. *J Neurosci* 2004;24:10410-5.

- 1 34. Giesecke T, Gracely RH, Grant MA, et al. Evidence of augmented central pain
2 processing in idiopathic chronic low back pain. *Arthritis Rheum*
3 2004;50:613-23.
- 4 35. Baliki MN, Chialvo DR, Geha PY, et al. Chronic pain and the emotional brain:
5 specific brain activity associated with spontaneous fluctuations of intensity
6 of chronic back pain. *The Journal of Neuroscience* 2006;26:12165e73.
- 7 36. Baliki MN, Apkarian AV. Neurological effects of chronic pain. *J Pain Palliat*
8 *Care Pharmacother* 2007;21:59-61.
- 9 37. Kaptchuk TJ, Kelley JM, Conboy LA, et al. Components of placebo effect:
10 randomised controlled trial in patients with irritable bowel syndrome. *BMJ*
11 2008;336:999-1003.
- 12 38. deCharms RC, Maeda F, Glover GH, et al. Control over brain activation and
13 pain learned by using real-time functional MRI. *Proc Natl Acad Sci U S A*
14 2005;102:18626-31.
- 15 39. Hartvigsen J, Nielsen J, Kyvik KO, et al. Heritability of spinal pain and
16 consequences of spinal pain. A comprehensive genetic epidemiologic
17 analysis using a population-based sample of 35,315 Danish twins aged 20-71.
18 *Arthritis Rheum* 2009;61:1343-51.
- 19 40. Nachemson A. Adult scoliosis and back pain. *Spine* 1979;4:513-7.
- 20 41. Tajerian M, Alvarado S, Millicamps M, et al. DNA methylation of SPARC and
21 chronic low back pain. *Mol Pain* 2011;7.
- 22 42. Doehring A, Geisslinger G, Lötsch J. Epigenetics in pain and analgesia: an
23 imminent research field. *Eur J Pain* 2011;15:11-6.
- 24 43. Pincus T, Santos R, Breen A, et al. Multinational Musculoskeletal Inception
25 Cohort Study Collaboration. A review and proposal for a core set of factors
26 for prospective cohorts in low back pain: a consensus statement. *Arthritis*
27 *Rheum* 2008;59:14-24.
- 28 44. Spine Tango, an international spine registry, the Spine Society of Europe.
29 Available from: <http://www.eurospine.org/p31000381.html>. Accessed 28
30 March 2013.
- 31 45. Linton SJ, Vlaeyen J, Ostelo R. The back pain beliefs of health care providers:
32 are we fear-avoidant? . *J Occup Rehabil* 2002;12:223-32.
- 33 46. Morris J, Watson PJ. Investigating decisions to absent from work with low
34 back pain: a study combining patient and GP factors. *Eur J Pain* 2011;15:278-
35 85.
- 36 47. Pincus T, Greenwood L, McHarg E. Advising people with back pain to take
37 time off work: a survey examining the role of private musculoskeletal
38 practitioners in the UK. *Pain* 2011;152:2813-8.
- 39 48. Ussher M, Spatz A, Copland C, et al. Immediate effects of a brief mindfulness-
40 based body scan on patients with chronic pain. *J Behav Med* 2012;PubMed
41 PMID: 23129105.
- 42 49. McCracken L, M., Gutiérrez-Martínez O, Smyth C. "Decentering" Reflects
43 Psychological Flexibility in People With Chronic Pain and Correlates With
44 Their Quality of Functioning. *Health Psychol* 2012.
- 45 50. Anema JR, Schellart AJM, Cassidy JD, et al. Can Cross Country Differences in
46 Return-to-Work After Chronic Occupational Back Pain be Explained? An
47 Exploratory Analysis on Disability Policies in a Six Country Cohort Study. *J*
48 *Occup Rehabil* 2009;19:419-26.

- 1 51. Lippel K, Lötters F. Public Insurance Systems: A Comparison of Cause-Based
2 and Disability-Based Income Support Systems. In Anema PLaH ed. *Handbook*
3 *of Work Disability: Prevention and Management*. New York Springer
4 Science+Business Media, 2013 (in Press).
- 5 52. Cassidy JD, Carroll LJ, Côté P, et al. Effect of Eliminating Compensation for
6 Pain and Suffering on the Outcome of Insurance Claims for Whiplash Injury.
7 *N Engl J Med* 2000;342:1179-86.
- 8 53. de Vries HJ, Reneman MF, Groothoff JW, et al. Self-reported work ability and
9 work performance in workers with chronic nonspecific musculoskeletal pain.
10 *J Occup Rehabil* 2013;23:1-10.
- 11 54. Franche RL, Cullen K, Clarke J, et al. Workplace-based return-to-work
12 interventions: a systematic review of the quantitative literature. *J Occup*
13 *Rehabil* 2005;15:607-31.
- 14 55. Carroll C, Rick J, Pilgrim H, et al. Workplace involvement improves return to
15 work rates among employees with back pain on long-term sick leave: a
16 systematic review of the effectiveness and cost-effectiveness of interventions.
17 *J Disabil Rehabil* 2010;32:607-21.
- 18 56. Elders L, van der Beek A, Burdorf A. Return to work after sickness absence
19 due to back disorders--a systematic review on intervention strategies. *Int*
20 *Arch Occup Environ Health* 2000;73:339-48.
- 21 57. Werner EL, Laerum E, Wormgoor ME, et al. Peer support in an occupational
22 setting preventing LBP-related sick leave. *Occup Med (Lond)* 2007;57:590-5.
- 23 58. Shaw W, Hong QN, Pransky G, et al. A literature review describing the role of
24 return-to-work coordinators in trial programs and interventions designed to
25 prevent workplace disability. *J Occup Rehabil* 2008;18:2-15.
- 26 59. Higgins A, O'Halloran P, Porter S. Management of long term sickness
27 absence: a systematic realist review. *J Occup Rehabil* 2012;22:322-32.
- 28 60. van Oostrom SH, Driessen MT, de Vet HC, et al. Workplace interventions for
29 preventing work disability. *Cochrane Database Syst Rev* 2009;15:CD006955.
- 30 61. Squires H, Rick J, Carroll C, et al. Cost-effectiveness of interventions to return
31 employees to work following long-term sickness absence due to
32 musculoskeletal disorders. *J Public Health (Oxf)*. 2012;34:115-24.
- 33 62. MacEachen E, Kosny A, Ferrier S, et al. The "toxic dose" of system problems:
34 Why some injured workers don't return to work as expected. *J Occup Rehabil*
35 2010;20:349-66.
- 36 63. Waddell G. Preventing incapacity in people with musculoskeletal disorders.
37 *BMJ* 2006;77-78:55-69.
- 38 64. Hill JC, Whitehurst DG, Lewis M, et al. Comparison of stratified primary care
39 management for low back pain with current best practice (STarT Back): a
40 randomised controlled trial. *Lancet* 2011;378:1560-71.
- 41 65. Vibe Fersum K, O'Sullivan P, Skouen JS, et al. Efficacy of classification-based
42 cognitive functional therapy in patients with non-specific chronic low back
43 pain: A randomized controlled trial. *Eur J Pain* 2013; 17(6):916-28.
- 44 66. Shaw WS, Reme SE, Woiszwilllo MJ, et al. The PRICE (Pain Recovery Issues,
45 COncerns, and Expectations) questionnaire: a brief psychosocial screening
46 instrument to identify intervention needs among patients at elevated risk of
47 back disability. *J Occup and Environ Med* In press.
- 48 67. Dunn KM. Extending conceptual frameworks: life course epidemiology for
49 the study of back pain. *BMC Musculoskelet Disord* 2010;11:1-11.

- 1 68. Ramond A, Bouton C, Richard I, et al. Psychosocial risk factors for chronic
2 low back pain in primary care—a systematic review. *Fam Pract* 2011;28:12–
3 21.
- 4 69. Buchbinder R, Gross DP, Werner EL, et al. Understanding the characteristics
5 of effective mass media campaigns for back pain and methodological
6 challenges in evaluating their effects. *Spine* 2008;33:74-8.
- 7 70. Hancock M, Herbert R, Maher CG. A guide to interpretation of studies
8 investigating subgroups of responders to physical therapy interventions.
9 *Phys Ther* 2009;89:698-704.

References

1. Waddell G. A new clinical model for the treatment of low-back pain. *Spine* 1987;12:632-44.
2. Lim SS, Vos T, Flaxman AD, et al. A comparative risk assessment of burden of disease and injury attributable to 67 risk factors and risk factor clusters in 21 regions, 1990—2010: a systematic analysis for the Global Burden of Disease Study. *Lancet* 2012;380.
3. Murray CJ, Vos T, Lozano R, et al. Disability-adjusted life years (DALYs) for 291 diseases and injuries in 21 regions, 1990-2010: a systematic analysis for the Global Burden of Disease Study 2010. *The Lancet* 2012;380:2197-223.
4. Ivanova JI, Birnbaum HG, Schiller M, et al. Real-world practice patterns, health-care utilization, and costs in patients with low back pain: the long road to guideline-concordant care. *Spine J* 2011;11:622-32.
5. Deyo RA. Managing patients with back pain: putting money where our mouths are not. *Spine J* 2011;11:633-5.
6. Pincus T, Burton A, Vogel S, et al. A systematic review of psychological factors as predictors of chronicity/disability in prospective cohorts of low back pain. *Spine* 2002;27:E109-E20.
7. Gatchel RJ, Peng YB, Peters ML, et al. The biopsychosocial approach to chronic pain: scientific advances and future directions. *Psychol Bull* 2007;133:581-624.
8. Nicholas MK, Linton SJ, Watson PJ, et al. Early identification and management of psychological risk factors ("yellow flags") in patients with low back pain: a reappraisal. *Phys Ther* 2011;91:737-53.
9. van der Windt D, Hay EM, Jellema P, et al. Psychosocial interventions for low back pain in primary care: lessons learned from recent trials. *Spine* 2008;33:81-9.
10. Williams AC, Eccleston C, Morley S. Psychological therapies for the management of chronic pain (excluding headache) in adults. *Cochrane Database of Systematic Reviews* 2012;11.
11. Loisel P, Buchbinder R, Hazard R, et al. Prevention of work disability due to musculoskeletal disorders: the challenge of implementing evidence. *J Occup Rehabil* 2005;15:507-24.
12. Artus M, van der Windt DA, Jordan KP, et al. Low back pain symptoms show a similar pattern of improvement following a wide range of primary care treatments: a systematic review of randomized clinical trials. *Rheumatology (Oxford)* 2010;49:2346-56.
13. Mannion A, Junge A, Taimela S, et al. Active therapy for chronic low back pain - Part 3. Factors influencing self-rated disability and its change following therapy. *Spine* 2001;26:920-9.
14. Loisel P, Durand M-J, Berthelette D, et al. Disability prevention: The new paradigm of management of occupational back pain. *Dis Manage Health Outcomes* 2001;9:351-60.
15. Loisel P, Côté P. The Work Disability Paradigm and its Public Health Implications. In Anema PLaH ed. *Handbook of Work Disability: Prevention and Management*. New York: Springer Science+Business Media, 2013.

16. Sullivan MJ, Thibault P, Andrikonyte J, et al. Psychological influences on repetition-induced summation of activity-related pain in patients with chronic low back pain. *Pain* 2009;141:70-8.
17. Bishop P, Wing P. Compliance with clinical practice guidelines in family physicians managing workers compensation board patients with acute lower back pain. *Spine J* 2003;3:442-50.
18. Shaw WS, Pransky G, Winters T, et al. Does the presence of psychosocial 'yellow flags' alter patient-provider communication for work-related, acute low back pain? *J Occup Environ Med* 2009;51:1032-40.
19. Foster NE, Hartvigsen J, Croft PR. Taking responsibility for the early assessment and treatment of patients with musculoskeletal pain: a review and critical analysis. *Arthritis Res Ther* 2012;14.
20. Hancock MJ, Maher CG, Laslett M, et al. Discussion paper: what happened to the 'bio' in the bio-psycho-social model of low back pain? *Eur Spine J* 2011;20:2105-10.
21. Langevin H, M., Sherman KJ. Pathophysiological model for chronic low back pain integrating connective tissue and nervous system mechanisms. *Med Hypotheses* 2007;68:74-80.
22. Marras WS. The complex spine: the multidimensional system of causal pathways for low-back disorders. *Hum Factors* 2012;54:881-9.
23. Suri P, Dharamsi AS, Gaviola G, et al. Are facet joint bone marrow lesions and other facet joint features associated with low back pain? A pilot study. *PM R* 2013;5:194-200.
24. Jensen RK, Leboeuf-Yde C, Wedderkopp N, et al. Is the development of Modic changes associated with clinical symptoms? A 14-month cohort study with MRI. *Eur Spine J* 2012;21:2271-9.
25. Deyo R. Diagnostic evaluation of LBP: reaching a specific diagnosis is often impossible. *Arch Intern Med* 2002;162:1444-7.
26. Deyo RA, Mirza SK, Turner JA, et al. Overtreating chronic back pain: time to back off? *J Am Board Fam Med* 2009;22:62-8.
27. Breen AC, Teyhen DS, Mellor FE, et al. Measurement of intervertebral motion using quantitative fluoroscopy: report of an international forum and proposal for use in the assessment of degenerative disc disease in the lumbar spine. *Adv Orthop* 2012;802350.
28. Tan Y, Aghdasi BG, Montgomery SR, et al. Kinetic magnetic resonance imaging analysis of lumbar segmental mobility in patients without significant spondylosis. *European Spine Journal* 2012;21:2673-9.
29. Chan ST, Fung PK, Ng NY, et al. Dynamic changes of elasticity, cross-sectional area, and fat infiltration of multifidus at different postures in men with chronic low back pain. *Spine J* 2012;12:381-8.
30. Marras W, Ferguson S, Gupta P, et al. The Quantification of Low Back Disorder Using Motion Measures: Methodology and Validation. *Spine* 1999;24:2091.
31. Wand BM, O'Connell NE. Chronic non-specific low back pain - sub-groups or a single mechanism? *BMC Musculoskelet Disord* 2008;9.

32. Wand BM, Parkitny L, O'Connell NE, et al. Cortical changes in chronic low back pain: current state of the art and implications for clinical practice. *Man Ther* 2011;16:15-20.
33. Apkarian A, V. , Sosa Y, Sonty S, et al. Chronic back pain is associated with decreased prefrontal and thalamic gray matter density. *J Neurosci* 2004;24:10410-5.
34. Giesecke T, Gracely RH, Grant MA, et al. Evidence of augmented central pain processing in idiopathic chronic low back pain. *Arthritis Rheum* 2004;50:613-23.
35. Baliki MN, Chialvo DR, Geha PY, et al. Chronic pain and the emotional brain: specific brain activity associated with spontaneous fluctuations of intensity of chronic back pain. *The Journal of Neuroscience* 2006;26:12165e73.
36. Baliki MN, Apkarian AV. Neurological effects of chronic pain. *J Pain Palliat Care Pharmacother* 2007;21:59-61.
37. Kaptchuk TJ, Kelley JM, Conboy LA, et al. Components of placebo effect: randomised controlled trial in patients with irritable bowel syndrome. *BMJ* 2008;336:999-1003.
38. deCharms RC, Maeda F, Glover GH, et al. Control over brain activation and pain learned by using real-time functional MRI. *Proc Natl Acad Sci U S A* 2005;102:18626-31.
39. Hartvigsen J, Nielsen J, Kyvik KO, et al. Heritability of spinal pain and consequences of spinal pain. A comprehensive genetic epidemiologic analysis using a population-based sample of 35,315 Danish twins aged 20-71. *Arthritis Rheum* 2009;61:1343-51.
40. Nachemson A. Adult scoliosis and back pain. *Spine* 1979;4:513-7.
41. Tajerian M, Alvarado S, Millecamps M, et al. DNA methylation of SPARC and chronic low back pain. *Mol Pain* 2011;7.
42. Doehring A, Geisslinger G, Lötsch J. Epigenetics in pain and analgesia: an imminent research field. *Eur J Pain* 2011;15:11-6.
43. Pincus T, Santos R, Breen A, et al. Multinational Musculoskeletal Inception Cohort Study Collaboration. A review and proposal for a core set of factors for prospective cohorts in low back pain: a consensus statement. *Arthritis Rheum* 2008;59:14-24.
44. Spine Tango, an international spine registry, the Spine Society of Europe. Available from: <http://www.eurospine.org/p31000381.html>. Accessed 28 March 2013.
45. Linton SJ, Vlaeyen J, Ostelo R. The back pain beliefs of health care providers: are we fear-avoidant? *J Occup Rehabil* 2002;12:223-32.
46. Morris J, Watson PJ. Investigating decisions to absent from work with low back pain: a study combining patient and GP factors. *Eur J Pain* 2011;15:278-85.
47. Pincus T, Greenwood L, McHarg E. Advising people with back pain to take time off work: a survey examining the role of private musculoskeletal practitioners in the UK. *Pain* 2011;152:2813-8.
48. Ussher M, Spatz A, Copland C, et al. Immediate effects of a brief mindfulness-based body scan on patients with chronic pain. *J Behav Med* 2012;PubMed PMID: 23129105.

49. McCracken L, M., Gutiérrez-Martínez O, Smyth C. "Decentering" Reflects Psychological Flexibility in People With Chronic Pain and Correlates With Their Quality of Functioning. *Health Psychol* 2012.
50. Anema JR, Schellart AJM, Cassidy JD, et al. Can Cross Country Differences in Return-to-Work After Chronic Occupational Back Pain be Explained? An Exploratory Analysis on Disability Policies in a Six Country Cohort Study. *J Occup Rehabil* 2009;19:419-26.
51. Lippel K, Lötters F. Public Insurance Systems: A Comparison of Cause-Based and Disability-Based Income Support Systems. In Anema PLaH ed. *Handbook of Work Disability: Prevention and Management*. New York Springer Science+Business Media, 2013 (in Press).
52. Cassidy JD, Carroll LJ, Côté P, et al. Effect of Eliminating Compensation for Pain and Suffering on the Outcome of Insurance Claims for Whiplash Injury. *N Engl J Med* 2000;342:1179-86.
53. de Vries HJ, Reneman MF, Groothoff JW, et al. Self-reported work ability and work performance in workers with chronic nonspecific musculoskeletal pain. *J Occup Rehabil* 2013;23:1-10.
54. Franche RL, Cullen K, Clarke J, et al. Workplace-based return-to-work interventions: a systematic review of the quantitative literature. *J Occup Rehabil* 2005;15:607-31.
55. Carroll C, Rick J, Pilgrim H, et al. Workplace involvement improves return to work rates among employees with back pain on long-term sick leave: a systematic review of the effectiveness and cost-effectiveness of interventions. *J Disabil Rehabil* 2010;32:607-21.
56. Elders L, van der Beek A, Burdorf A. Return to work after sickness absence due to back disorders--a systematic review on intervention strategies. *Int Arch Occup Environ Health* 2000;73:339-48.
57. Werner EL, Laerum E, Wormgoor ME, et al. Peer support in an occupational setting preventing LBP-related sick leave. *Occup Med (Lond)* 2007;57:590-5.
58. Shaw W, Hong QN, Pransky G, et al. A literature review describing the role of return-to-work coordinators in trial programs and interventions designed to prevent workplace disability. *J Occup Rehabil* 2008;18:2-15.
59. Higgins A, O'Halloran P, Porter S. Management of long term sickness absence: a systematic realist review. *J Occup Rehabil* 2012;22:322-32.
60. van Oostrom SH, Driessen MT, de Vet HC, et al. Workplace interventions for preventing work disability. *Cochrane Database Syst Rev* 2009;15:CD006955.
61. Squires H, Rick J, Carroll C, et al. Cost-effectiveness of interventions to return employees to work following long-term sickness absence due to musculoskeletal disorders. *J Public Health (Oxf)*. 2012;34:115-24.
62. MacEachen E, Kosny A, Ferrier S, et al. The "toxic dose" of system problems: Why some injured workers don't return to work as expected. *J Occup Rehabil* 2010;20:349-66.
63. Waddell G. Preventing incapacity in people with musculoskeletal disorders. *BMJ* 2006;77-78:55-69.

64. Hill JC, Whitehurst DG, Lewis M, et al. Comparison of stratified primary care management for low back pain with current best practice (STarT Back): a randomised controlled trial. *Lancet* 2011;378:1560–71.
65. Vibe Fersum K, O’Sullivan P, Skouen JS, et al. Efficacy of classification-based cognitive functional therapy in patients with non-specific chronic low back pain: A randomized controlled trial. *Eur J Pain* 2013; 17(6):916-28.
66. Shaw WS, Reme SE, Woiszwilllo MJ, et al. The PRICE (Pain Recovery Issues, COncerns, and Expectations) questionnaire: a brief psychosocial screening instrument to identify intervention needs among patients at elevated risk of back disability. *J Occup and Environ Med* In press.
67. Dunn KM. Extending conceptual frameworks: life course epidemiology for the study of back pain. *BMC Musculoskelet Disord* 2010;11:1-11.
68. Ramond A, Bouton C, Richard I, et al. Psychosocial risk factors for chronic low back pain in primary care—a systematic review. *Fam Pract* 2011;28:12–21.
69. Buchbinder R, Gross DP, Werner EL, et al. Understanding the characteristics of effective mass media campaigns for back pain and methodological challenges in evaluating their effects. *Spine* 2008;33:74-8.
70. Hancock M, Herbert R, Maher CG. A guide to interpretation of studies investigating subgroups of responders to physical therapy interventions. *Phys Ther* 2009;89:698-704.

*LWW Copyright Transfer and Disclosure Form

[Click here to download LWW Copyright Transfer and Disclosure Form: copyrightTransfer.pdf](#)

*LWW Copyright Transfer and Disclosure Form

[Click here to download LWW Copyright Transfer and Disclosure Form: copyrightTransfer \(3\) pransky pincus gp.pdf](#)

*LWW Copyright Transfer and Disclosure Form

[Click here to download LWW Copyright Transfer and Disclosure Form: copyrightTransfer_PK.pdf](#)

*LWW Copyright Transfer and Disclosure Form

[Click here to download LWW Copyright Transfer and Disclosure Form: copyrightTransfer JH.pdf](#)

*LWW Copyright Transfer and Disclosure Form

[Click here to download LWW Copyright Transfer and Disclosure Form: copyrightTransfer PL.pdf](#)

*LWW Copyright Transfer and Disclosure Form

[Click here to download LWW Copyright Transfer and Disclosure Form: bronfort spineCOI.pdf](#)